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Forensic Population Genetics-Letter to Editor

Dear JM,

DNA samples were analysed from 519 healthy, unrelated and consenting individuals who reside in the United Arab Emirates (UAE) and were randomly chosen for this study. The UAE is one of the middle-eastern countries located on the Arabian Gulf. It shares a border with Iran, Saudi Arabia and Oman. The UAE was founded in 1971, and consists of seven Emirates: Abu Dhabi, Dubai, Sharjah, Ajman, Ra's Al-Khaymah, Al-Fujairah and Umm Al-Quwain [1]. According to the National Bureau of Statistics, (2012), the total UAE population was reported to be around 8.26 million in 2010 [2]. The statistics showed that some 11.5% of the total population comprised of native Arabs, with majority of the population being of Indian and Pakistani ethnicities. In the early part of the twentieth century, the different Arabic tribes migrated in different directions in search of suitable locations to colonize. Some moved into coastal regions, while others inhabited the desert. Despite the modernization throughout the union, the basic family structure and pattern of native UAE Arab population has remained unchanged. Culturally, the preference for consanguineous marriages remains embedded in the society [3]. However, as the awareness of the social and medical impact of consanguinity increases and with diversification, non-consanguineous marriages appear to be on the increase, which has possibly resulted in greater genetic diversity throughout the population [4, 5]. The increase in genetic diversity in the population is of interest to assess whether STR markers can be used for forensic and paternity purposes. This study expands on previous publications with regards to the analysis of UAE populations with the amplification of additional STR markers and a larger population sample size [6].

The DNA samples analysed in the current study were obtained from indigenous UAE nationals residing in Abu Dhabi, UAE in accordance with approval from the Ethics committee of the Ministry of Health of the United Arab Emirates (2011). Informed consent was received from every volunteer during this collection process and de-identified data is presented. This study was also approved by the Ethics committee of the University of Central Lancashire (2014) as it was carried out as part of Masters Project in DNA profiling.

The DNA samples provided for this study were collected and extracted using the Genotek's Oragene-DNA kit (Genotek, Ottawa, Canada) in accordance with manufacturer's guidelines. The quantities of extracted DNA samples were determined using a NanoDrop spectrophotometer (Thermo Scientific, Wilmington DE, USA).

Using half volume (7.5 µl) reactions, samples were amplified using the GlobalFiler® PCR amplification kit (Life Technologies, Foster City CA, USA) and alleles were called using the allelic ladder provided by the manufacturer. The PCR was performed in the GeneAmp® PCR System 9700 (Life Technologies). The GlobalFiler® PCR amplification kit (Life Technologies) amplifies 21 autosomal STR loci, a Y-STR locus DYS 391, a Y-indel marker and Amelogenin. The 21 autosomal STR loci within this amplification kit were of interest for the purposes of this study. The 21 autosomal loci amplified and focused on within this study were D3S1358, vWA, D16S539, CSF1PO, TPOX, D8S1179, D21S11, D18S51, D2S441, D19S433, TH01, FGA, D22S1045, D5S818, D13S317, D7S820, SE33, D10S1248, D1S1656, D12S391, D2S1338.

The PCR products were analysed using an 8 capillary ABI 3500 DNA Genetic Analyser with POP-4™ polymer (Life Technologies). GeneMapper® Software version 4.0 (Life Technologies) was then used for analysis. LIZ-600 was used as internal Standard (Life Technologies).

The alleles from all loci reported here were designated using the allelic ladder supplied by the manufacturer, according to the published nomenclatures and the guidelines of the International Society for Forensic Genetics (ISFG) for performing STR analyses [7].

The STR allele frequencies along with the parameters of population genetics: observed and expected heterozygosity (H_o and H_e , respectively), power of discrimination (PD), probability of exclusion (PE), and polymorphic information content (PIC) were estimated using PowerStats version 1.2 (Promega, Madison, USA) (Supplementary Table 1).

Version 3.11 of the Arlequin software was used to perform an exact test to investigate any departures from the Hardy-Weinberg equilibrium (HWE) [8]. The theoretical profile frequency range was estimated signifying the rarest and most common heterozygous genotypes. Furthermore, the number of possible genotypes was also calculated (Supplementary Table 2).

significant differences at fewer loci when compared with populations from Kuwait, Egypt and India ($P > 0.05$). This is also supported by low F_{ST} value for the Iranian and Saudi Arabian populations. These results support the development of population or location specific databases even when considering populations that are geographically close such as within the Middle East (Supplementary Table 3).

This current dataset establishes the characteristics of the 21 STR loci panel for the identification of individuals, in paternity testing and for crime scene analysis in the UAE.

This manuscript of population data follows the journal guidelines for publication of data described [11, 12 and 13].

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Appendix A. Supplementary data

[Publication\Supplementary Table 1.xlsx](#)

[Publication\Supplementary Table 2.xlsx](#)

[Publication\Supplementary Table 3.xlsx](#)

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