Abstract

The Y chromosome contains the largest non-recombining portion in the human genome. Y-Binary polymorphisms, also known as a single nucleotide polymorphisms (SNPs), are a series of biallelic polymorphisms occurring on the non-recombining region of the Y chromosome (NRY), which represent a precious tool for human evolutionary studies and, potentially, for forensic applications. Low mutation rate, paternal inheritance, and absence of recombination make Y-SNPs particularly suitable for the identification of stable paternal lineages and the reconstruction of an ancestral state from which to explore the evolution of humans. Also, these markers would allow inference of the paternal ancestry of unknown samples which could be useful in forensic applications. In the present study we analyzed 22 biallelic Y-SNPs in 159 males belonging to three ethnic groups (Arab n = 42, Berber n = 67 and Sahrawi n = 50) from Morocco. A total of 10 different haplogroups were identified in this sample representative of Moroccan population. The most common Y chromosome haplogroups is E1b1b1, E1b1b1b and J1 with frequencies of 56%, 49% and 10% respectively.

Keywords: Y-chromosomal SNP; Mini-sequencing; SNaPshot; Morocco

Introduction

Biallelic markers, such as single nucleotide polymorphisms (SNPs) and insertion/deletions (indels), represent an important class of markers on the Y-chromosome [1]. Y-chromosome SNPs (Y-SNPs) are mostly used in molecular anthropology for evolutionary research. Moreover, typing a sample’s Y chromosome haplogroup allows paternal ancestry inference. This may be useful, in forensic applications, when a conventional Short Tandem Repeat (STR) profile, generated from DNA collected at a crime scene, does not match any of the identified suspects and doesn’t “hit” any profile on the available databases.

The kingdom of Morocco is a country located in the northwestern corner of the African continent with coasts on the Atlantic Ocean and the Mediterranean Sea; it is bordered by Algeria to the east and Mauritania to the south. Modern-day Morocco is inhabited by three major ethnic groups (Arab, Berber and Sahrawi). Several dialects such as Arabic (Moroccan dialect or Darija), Berber (Tarifit, Tachelhit and Tamazight) and Sahrawi (El Hassania) are spoken in the country. The aim of the present study was to develop an assay to genotype a selected panel of Y-SNPs using single base extension assay (SBE), or minisequencing [2], in order to determine the most frequent Y-chromosomal haplogroup in Moroccan population. The analysis of single nucleotide polymorphisms (SNPs) is a promising application in forensic casework; since the forensic scientists is often faced with degraded and/or very low amounts of DNA.

Materials and Methods

Population

Buccal swabs were collected from 159 unrelated healthy adult men belonging to the three ethnic groups (Arab n = 42, Berber n = 67 and Sahrawi n = 50). Informed consent was obtained from all participants in this study, and information about the geographical origin of their grand-parents and about their first language was recorded.

DNA Isolation

Genomic DNA was extracted from buccal swab punches, using DNA IQ™ System (Promega; Madison, Wisconsin) on the Biomek® 2000 (Beckman Coulter, Brea, CA) robotic platform according to manufacturer’s instructions.

Y-SNP selection and multiplex design

Assay design and development together with sample testing were conducted in the Forensic Molecular Biology Laboratory of the Forensic Sciences Department of The George Washington University. A total of 22 Y chromosome biallelic markers were selected for this study following a hierarchical strategy based on the phylogenetic tree of Y chromosome recognized by the Y Chromosome Consortium (YCC) [3]. Loci Nomenclature is based on Karafet et al. [4]. The reference sequence for each Y-SNP was taken from the International Society of Genetic Genealogy, Y-DNA Haplogroup Tree 2010 (http://www.isog.org/tree). The 22 markers were genotyped in two heptaplex, one hexaplex, and one duplex PCR reactions. The Multiplex MY01 allows the detection of major clades (A-R), A-M91 (A), B-M60 (B), C-RPS4Y711 (C), D-M174 (D), J-M267 (J1), J-M172 (J2a), and R-M207 (R). The Multiplexes MY02 E-P147 (E1), E-M132 (E1a), E-P189 (E1b1a), E-M215 (E1b1b), E-M81 (E1b1b1b), E-M54 (E2b), J-M172 (J2), MY03 J-L24 (J2a4h), J-M221 (J2b), R-L63 (R1a), R-M343 (R1b), R-M269 (R1b1b2), R-M68 (R1b1b2).

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T-M70 (T) and MY04 E-M35 (E1b1b1), E-M78 (E1b1b1a) subdivides haplogroups E, J, R and T to define the most frequent haplogroups in NorthWest Africa. Multiplex minisequencing primers were designed for each multiplex PCR. PCR primers used in the primer extension assays are listed in Table 1. The extension primers designed in four multiplex sets labeled as MY01, MY02, MY03 and MY04 are listed in Table 2.

**PCR amplification**

Amplifications were performed in a final volume of 25 µL using a master mix containing 1X GeneAmp® PCR Gold buffer (Applied Biosystems, Foster City, CA), 400 nM of dNTPs, 3 mM of MgCl₂, 0.65 mg/µL bovine serum albumin (BSA), and 1.5 units of AmpliTaq Gold® DNA polymerase (Applied Biosystems). The thermal cycling program was carried out on a GeneAmp 9700 (Applied Biosystems) using the following conditions: 95°C for 10 minutes, 35 cycles of 94°C for 30 seconds, 55°C for 30 seconds and 72°C for 30 seconds, and a final extension at 72°C for 10 minutes.

Following PCR amplification, unincorporated primers and dNTPs were removed by adding 1 unit of Exonuclease I (USB Corporation) and 1 unit of Shrimp Alkaline Phosphatase (SAP) to 5 µL of PCR products and incubating at 37°C for 70 minutes followed by 20 minutes at 65°C.

**Primer extension assay using fluorescence detection**

Multiplex primer extension reactions were carried out in a total volume of 15 µL using 1.5 µL of ABI Prism® SNaPshot™ multiplex kit mix (Applied Biosystems), 2 µL of purified PCR product and 1.5 pmols of each extension primer. Extension reaction was as follows: 25 cycles of 96°C for 10 seconds, 50°C for 5 seconds, and 60°C for 30 seconds.

A 1.5 µL of non-purified primer extension product was diluted in 9.8 µL of Hi-Di™ formamide and 0.2 µL of GeneScan-450 LIZ internal lane standard (Applied Biosystems) and analyzed on ABI Prism 3730 Genetic Analyzer (Applied Biosystems). Separation was performed on a 36 cm array using POP™-7 (Applied Biosystems). Results were analyzed using GeneMapper v 4.0 (Applied Biosystems).

**Results and Discussion**

A total of 10 haplogroups were observed across the 159 male samples that were tested (Table 3). The most common Y chromosome haplogroups in the combined Moroccan population resulted E1b1b1, E1b1b1b and J1 (56%, 49% and 10% respectively).

The E-M215 derivative E1b1b1 is observed in significant frequencies in Moroccan samples. It’s defined by the E-M35 SNP which appears to have originated in East Africa then migrating to the Near East and then on to North Africa and Europe.

Previous studies have shown that E1b1b1b (E-M81), formerly E3b1b or E3b2, is the predominant haplogroup in northwestern Africa [5 - 8]. Bosh et al. [9] obtained a higher frequency (64%) of E-M81 (E1b1b1b) in Northwest African populations than the present study. This disparity could be explained by the difference of geographic area and circumstances of sample collection. However, according to our results and those of Bosh et al. [9], it seems that the E1b1b1b haplogroup is more characteristic of Sahrawi than Berbers. It is also seen, although at low frequency, in the Iberian Peninsula (4%) and Sicily (3%) due to recent gene flow from Northwest Africa through Gibraltar strait [9 - 10]. Nevertheless, this haplogroup (E1b1b1b) is not found in sub-Saharan Africa and its frequency sharply declines through the continent towards the east.

The J1 haplogroup, defined by the single nucleotide polymorphism (SNP) M267, is most frequent in the Arabian Peninsula especially in Yemen (76%) [11]. This could be attributed to the early medieval period during which the Semitic expansion spread J1 out of Arabia into North Africa [12].

The multiplex assays described herein were designed to explore the shallowest branches of Y chromosome haplogroups in Moroccan population. They could also be applied to human evolution and human genetists studies as well as to forensic casework for ancestry inferences.

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