

Male genealogy study of the Iraqi Population

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Received: 17.08.2024

Revised: 19.09.2024

Accepted: 21.10.2024

ABSTRACT

The Y-chromosome of Short Tandem Repeats (STRs) plays a vital role in the field of forensics, particularly in the identification of male DNA in instances of sexual assault. Moreover, it is important in facilitating the connection of families through the application of genetic genealogy. In this study, the genetic polymorphisms of 23 Y-STR loci were analysed by the Power Plex® Y23 system in 383 healthy, unrelated male individuals randomly chosen from Iraqi populations. The result revealed 356 different haplotypes, including 337 unique haplotypes and 19 duplicate haplotypes. Two hundred and twelve different genetic variants (alleles), were detected at which are distributed across 23 specific locations on the Y chromosome. The study examined the highest and lowest frequencies at each genetic locus and found that at the DYS392 locus, the microvariant allele "11" was the most common and widespread allele within the Iraqi population. The highest frequency of haplotype was 0.0098, which was found in samples H44, H70, H140, H206, and H255, while 337 haplotypes had 0.0033 frequencies, and haplotype diversity was 0.997 with a discrimination capacity of 0.946, respectively. The gene diversity values ranged from 0.395062 at DYS392 to 0.93828 at DYS385a/b. The Y-haplogroups of male Iraqi Arabs were determined using the NevGen Y-DNA Haplogroup online Predictor program. The most common haplogroups in this study were J1 (38%), J2 (20%), R1 (11%), E1 (10%), and G2 (5%). Although less common, additional predicted haplogroups were also discovered. The present study indicated that the 23 Y-STR loci were highly polymorphic in the Bagdad population and played crucial roles in forensic applications as well as population genetics. The study revealed the genetic diversity of male lineages in the Bagdad population using a high-resolution 23 Y-STR collection, and as a result, it's contributed to familial research.

Keywords: Arabs, haplogroups, haplogroups, chromosome

INTRODUCTION

The Y chromosome's non-recombining region contains repeating areas called "Y-STRs," or short tandem repeats. These regions are made up of 2–7 base pair-long repetitive units. Due to their ease of typing and high level of polymorphism, Y-STRs are frequently used as markers on the Y chromosome. They also show a male inheritance pattern. Y-STR analysis is possible even in situations where DNA quality is reduced because PCR is a dependable method that can tolerate deteriorated DNA samples. Because of this, Y-STRs have a variety of uses, such as forensic sexual assault investigations, paternity testing when the purported father is unavailable for testing, circumstances with numerous male DNA samples being used in gang rape, and genetic research [1, 2].

Iraq is a country located in western Asia. Due to its good climate, fertile soil, and the presence of the Tigris and Euphrates rivers, Iraq is known as the cradle of the ancient civilization of Mesopotamia, including the Sumerian and Babylonian cultures. It is distinguished by its geographical characteristics, which include a varied terrain made up of mountains, deserts, and a productive river valley. Iraq is bordered by a number of nations, including Iran to the east, Turkey to the north, Syria, Jordan, and Saudi Arabia to the west, and Kuwait and Saudi Arabia to the south. The estimated population of Iraq in July 2011 was 30,399,572. Participants for the aforementioned study were chosen from Baghdad, which serves as Iraq's capital and largest city [3]. As a key metropolitan hub and the site of the research study's sampling, Baghdad was chosen. Iraq was exposed to the administration of many empires and civilizations over the span of its past history include the Mongols, Turkey, England and Iran. Arab Iraqi population made up of a wide of ethnic groups this highlights the significance conducting a study on history of society [4, 5]. The short tandem repeat loci (STR loci) are repeating sequences in of the human genome. It's highly polymorphic, different between people, have variable repeat counts which makes them very useful genetic markers. They can give information concern the bases behind the genetic origin of some diseases,

human migration and demographical research [6]. DNA typing used to determine individual's identification, paternity and maternity testes, investigated a criminal issue [7]. Y-STR markers demonstrate an inheritance pattern of specific male-transmission. Y-STRs can be used by researchers in several genetic applications [8, 9]. PCR-dependent techniques help to have accurate findings, even when dealing with degraded DNA samples. This ability to adjust to damage DNA is helpful for the analysis of ancient DNA samples to confirm the presence of individuals who lack amelogenin (a particular genetic condition) and other forensic applications [10, 11]. This particular power-plex Y 23 genotyping kit made from Promega is used to improve the analysis, accuracy, and discrimination power of Y-STR [12, 13]. Previous studies focused on the inherited characteristics of individuals representing different ethnic backgrounds in Iraq and its bordering countries. [14, 15, 16]. This study's goal is to investigate the genetic makeup of the Iraqi Arab population, haplogroup distributions, identification of a novel haplotypes and the creation of a reference database.

MATERIALS AND METHODS

Iraqi populations: The analysis of Y-STR included 383 healthy, unrelated men who were not related to each other and were not members of the same family. These people's blood was drawn at the Ministry of Health, more precisely at the Paternity and Kinship Section of the Medical Legal Directorate. The DNA for the analysis was derived from the acquired blood samples.

DNA extraction

In recent years, direct amplification has become more common as a faster and simpler substitute for conventional procedures. In comparison to earlier forensic science tests, more recent amplification kits seek to provide better and speedier results. However, some commercially available kits still call for pre-treating body fluids with washing chemicals or buffers in order to speed up the amplification procedure. The capacity to process more samples in less time and a significant reduction in analysis time are two key benefits of direct amplification without DNA purification. Blood samples spotted on FTA cards, air-dried then carefully packaged in paper envelopes until DNA extraction carried out [17].

STR amplification

An approximately 1.2 mm punch of male blood FTA cards used for amplification of 23 Y STR loci by using a PowerPlex system from Promega according to the manufacturer's instructions for ensuring accurate and reproducible results with 1 ± 2 ng of target DNA amplified using (Verity) a thermo cycler from Applied Biosystems [18].

Genotyping

The products of PCR amplification diluted with formamide (Hi-Di™) about 1:15, then analyzed with 3130 XL genetic analyzer using 3.2 version of gene mapper software (Applied Biosystems, USA).

STATISTICAL ANALYSIS

In the present study, the estimation of allele frequencies of haplotypes was used by gene counting methods. With exception, the locus DYS385 a/b is found on the same chromosome examined by the formula $D = (n/n-1)(1-\Sigma p_i^2)$, where (n) refers to the total allele number and (pi) refers to the frequency of one allele for calculating the genetic diversity of each tested haplotype. Population diversity was performed to estimate the discrimination ability of haplogroups. The new gen Y-DNA haplogroup predictor was used as a tool to identify ancestral genetic markers of the Y chromosome region. A molecular variance analysis (AMOVA) was conducted to estimate genetic diversity between the selected samples and compare it to the stored data from other populations using the Y-STR haplotype reference database 3.0 [19]. The multi-dimensional scaling (MDS) plot was performed using the results from the AMOVA analysis [20]. All the statistics used reveal information about ancestry, genetic origins, and the frequencies of Y haplotypes in a population [21, 22].

RESULTS

Statistical assessments were conducted to assess the appropriateness of the allele frequency databases derived from a sample of 383 Iraqi Arab individuals. The study performed an analysis to assess the frequency of particular haplotypes (combinations of genetic markers), in our sample population (Table 1). The DYS385a/b locus, which has a total of 55 distinct alleles, was discovered to have the most genetic diversity of the loci under study. Also high level of genetic variety was seen at the DYS458 locus, where a total of 15 distinct alleles were found. The loci DYS570 and DYS481, each showed 10 distinct alleles, also showed high levels of variability in addition to DYS458. Contrarily the lowest levels of genetic variety polymorphic loci were DYS391 with 4 alleles then followed by DYS533, DYS438, DYS437 and DYS393 with 5 alleles each locus (Table 1).

Table 1: Allele and Genotype frequencies in 23Y-STR haplotype of Iraqi Arab males.

DYS576		DYS389I		DYS448		DYS389II		DYS19		DYS385/a, b	
Allele	Ferq	Allele	Ferq	Allele	Ferq	Allele	Ferq	Allele	Ferq	Allele	Ferq
14	0.003	9	0.003	17	0.003	25	0.003	10	0.005	9,13	0.003
15	0.047	10	0.005	18	0.029	26	0.008	12	0.008	10,13	0.003
16	0.123	11	0.003	19	0.209	27	0.005	13	0.094	10,14	0.005
17	0.274	12	0.131	20	0.634	28	0.086	14	0.624	11,11	0.003
18	0.355	13	0.695	21	0.094	29	0.269	15	0.191	11,13	0.003
19	0.162	14	0.157	22	0.031	30	0.460	15,16	0.003	11,14	0.065
20	0.026	15	0.005			31	0.144	16	0.057	11,15	0.013
21	0.010	17	0.003			32	0.026	17	0.018	11,16	0.005
										11,17	0.005
										12,12	0.003
										12,14	0.021
										12,14.2	0.003
										12,15	0.029
										12,16	0.026
										12,17	0.005
										12,18	0.018
										12,19	0.003
										12,20	0.003
										12,22	0.005
										13,13	0.021
										13,14	0.018
										13,14.2	0.003
										13,15	0.047
										13,16	0.042
										13,17	0.037
										13.,17.2	0.003
										13,18	0.178
										13,19	0.112
										13,20	0.021
										13,21	0.003
										14,14	0.013
										14,15	0.016
										14,15.2	0.005
										14,16	0.050

DYS391		DYS481		DYS549		DYS533		DYS438	
Allele	Ferq	Allele	Ferq	Allele	Ferq	Allele	Ferq	Allele	Ferq
9	0.044	20	0.016	10	0.005	9	0.029	7	0.003
10	0.603	21	0.052	11	0.099	10	0.097	9	0.245
11	0.308	22	0.196	12	0.499	11	0.535	10	0.593
12	0.044	23	0.209	13	0.332	12	0.300	11	0.115
		24	0.154	14	0.055	13	0.039	12	0.044
		25	0.149	15	0.010				
		26	0.178						
		27	0.039						
		28	0.005						
		29	0.003						

DYS437		DYS570		DYS635		DYS390		DYS439	
Allele	Ferq	Allele	Ferq	Allele	Ferq	Allele	Ferq	Allele	freq
11	0.003	13	0.013	16	0.003	18	0.003	9	0.003
13	0.029	14	0.013	19	0.003	21	0.016	10	0.055
14	0.692	15	0.050	20	0.081	22	0.086	11	0.564
15	0.196	16	0.110	21	0.514	23	0.572	11,13	0.003
16	0.081	17	0.235	22	0.178	24	0.235	12	0.319
		18	0.326	23	0.162	25	0.073	13	0.052
		19	0.167	24	0.039	26	0.016	14	0.005
		20	0.063	25	0.013				
		21	0.016	26	0.005				
		22	0.008	28	0.003				

DYS392		DYS635		DYS393		DYS456		YGATAH ₄			
Allele	Ferq	Allele	Ferq	Allele	Ferq	Allele	Ferq	Allele	Ferq		
9	0.008	8	0.023	11	0.018	12	0.003	9	0.003	14,17	0.018
10	0.070	9	0.423	12	0.569	13	0.044	10	0.065	14,18	0.031
11	0.770	10	0.274	13	0.316	14	0.321	11	0.627	14,19	0.010
12	0.026	11	0.104	14	0.078	15	0.418	12	0.238	14,21	0.003
13	0.073	12	0.115	15	0.018	16	0.146	13	0.063	14,22	0.003
14	0.039	13	0.052			17	0.063	14	0.005	15,15	0.005
15	0.010	14	0.008			18	0.003			15,16	0.029
16	0.003					22	0.003			15,17	0.010
										15,21	0.003
										15,2,17	0.003
										16,16	0.008
										16,17	0.016
										16,18	0.018
										16,19	0.005
										16,20	0.005
										17,17	0.003
										17,18	0.010
										17,19	0.005
										18,18	0.003
										18,19	0.021
										19,19	0.005

DYS458	
Allele	Ferq
12	0.005
13	0.003
14	0.005
15	0.138
16	0.222
16.2	0.005
17	0.128
17.2	0.031
18	0.094
18.2	0.183
19	0.052
19.2	0.102
20	0.021
20.2	0.008
21.2	0.003

Frequency (Ferq): The bold refers to the most common allele's frequencies in each locus in the sample population. Except for **DYS385a/b**, the study calculated the genotype frequencies based on the combination of two alleles.

The study, as shown in Table 2, presents information about the occurrence of the highest and lowest allele frequencies at each genetic locus. Specifically, it points out that at the **DYS392** locus, the microvariant allele "11" was the most common and prevalent one in the Iraqi population. This information is important in genetic research as it helps understand the genetic makeup and diversity of a population.

Table 2: The occurrence of the highest and lowest allele frequencies at each genetic locus

	Freq		Freq		Freq		Freq
DYS576 High allele 18 Low allele 14	0.355 0.003	DYS391 High allele 13 Low allele 9,17	0.695 0.003	DYS448 High allele 17 Low allele 20	0.634 0.003	DYS389II High allele 25 Low allele 30	0.460 0.003
DYS391 High allele 14 Low allele 15,16	0.624 0.003	DYS385/a,b High allele 13,18 Low allele 9,13, 10,13, 11,11, 11,13 , 12,12, 12,14.2, 12,19, 12,20, 13,14.2,13.,17.2, 13,21	0.178 0.003	DYS481 High allele 23 Low allele 29	0.209 0.003	DYS549 High allele 12 Low allele 10	0.499 0.005
DYS391 High allele 10 Low allele 9,12	0.603 0.044	DYS533 High allele 11 Low allele 9	0.535 0.029	DYS438 High allele 10 Low allele 7	0.593 0.003	DYS437 High allele 14 Low allele 11	0.692 0.003
DYS570 High allele 18 Low allele 13,14	0.326 0.013	DYS635 High allele 21 Low allele 16,19	0.514 0.003	DYS390 High allele 23 Low allele 18	0.572 0.003	DYS439 High allele 11 Low allele 9,11,13	0.564 0.003
DYS392 High allele 11 Low allele 16	0.770 0.003	DYS635 High allele 9 Low allele 14	0.423 0.008	DYS393 High allele 12 Low allele 11,15	0.569 0.018	DYS456 High allele 15 Low allele 12,18,22	0.418 0.003
DYS458 High allele 16 Low allele 21.2	0.222 0.003	YGATAH4 High allele 11 Low allele 9	0.627 0.003				

There were a total of 383 different Y-STR haplotypes studied, out of these, 356 were unique, meaning they were not repeated in the dataset. This is seen as being quite very satisfying and demonstrates the potency of the PowerPlex Y 23 kit, a tool for examining genetic information. Several distinct haplotypes were discovered in the dataset more than once. For instance, there were three instances of each haplotype in samples H44, H70, H140, H206, and H255. These haplotypes were included in the sample population more frequently than others, compared to the other haplotypes, occurring at a frequency of 0.0098. In contrast, there were 337 different haplotypes that occurred with a lower frequency of 0.0033. Table (3) provides a genetic profile for each individual, showing the alleles they possess at various genetic markers.

Table 3: Y-STR haplotypes and haplotype frequencies in Iraq Arab populations

ID	DYS576	DYS389I	DYS448	DYS389II	DYS19	DYS391	DYS481	DYS549	DYS533	DYS438	DYS437	DYS570	DYS635	DYS390	DYS439	DYS392	DYS643	DYS393	DYS458	DYS385	DYS456	YGATAH4	FERQ
H1	18	13	20	30	17	10	25	12	12	11	14	20	23	25	10	11	10	13	16	10,13	16	12	0.0033
H2	17	13	20	31	15	10	24	13	12	11	14	19	23	26	11	12	10	13	15	10,14	15	12	0.0033
H3	19	13	20	30	16	11	23	11	12	11	14	20	23	25	10	11	10	13	15	10,14	16	12	0.0033
H4	17	13	20	30	15	11	23	12	12	10	14	19	23	25	11	11	10	13	15	11,11	15	11	0.0033
H5	16	14	19	30	14	10	20	11	11	10	14	20	22	23	11	12	11	14	16	11,13	14	12	0.0033
H6	17	13	19	30	14	10	22	12	12	12	15	17	21	24	12	12	10	12	16	11,14	15	12	0.0033
H7	17	13	19	30	14	10	22	13	12	12	15	17	23	24	12	13	9	12	16	11,14	15	12	0.0033
H8	17	13	19	30	14	10	22	13	12	12	15	17	23	24	12	13	10	12	16	11,14	15	12	0.0065
H9	17	13	19	30	14	10	22	13	12	12	15	17	23	25	12	13	10	12	16	11,14	16	12	0.0033
H10	17	13	19	29	14	10	22	14	12	12	15	17	23	24	13	13	10	12	16	11,14	15	13	0.0033
H25	18	13	20	30	17	10	23	12	12	11	14	18	23	23	11	11	10	13	15	11,16	15	13	0.0065
H37	18	13	20	30	14	10	23	12	11	9	14	16	21	23	11	11	10	12	17	12,15	16	11	0.0065
H44	19	12	21	28	15	10	22	12	9	10	16	19	21	23	12	11	11	15	16	12,16	15	12	0.0098
H58	18	14	19	30	14	10	24	11	10	11	15	17	24	23	11	11	10	15	15	12,22	15	12	0.0065
H70	16	13	20	29	14	10	24	13	11	10	15	18	24	23	10	11	11	12	16	13,16	15	11	0.0098
H93	18	13	20	30	14	10	25	12	11	10	14	18	20	23	11	10	9	13	17.2	13,16	15	11	0.0065
H140	18	13	20	30	14	11	26	13	11	10	14	18	21	23	11	11	9	12	18.2	13,18	14	11	0.0098
H150	19	13	20	30	14	11	26	12	10	9	14	18	21	23	11	11	9	12	19.2	13,18	13	11	0.0065
H175	18	13	20	30	14	11	25	12	11	10	14	18	21	23	11	11	9	12	18.2	13,19	14	11	0.0065
H185	18	13	20	31	17	10	25	13	11	10	14	18	21	23	11	11	9	13	19.2	13,19	16	11	0.0065
H195	17	13	20	30	14	11	26	13	11	10	14	18	21	23	11	11	9	12	18.2	13,20	14	11	0.0065
H204	16	13	20	29	14	10	22	13	11	9	14	16	21	22	12	11	8	12	15	14,15	16	11	0.0065
H206	17	13	21	30	15	10	20	12	12	9	16	18	21	26	11	11	9	13	16	14,16	15	11	0.0098
H217	19	13	20	30	15	10	22	11	11	9	15	16	22	23	11	11	12	12	18	14,16	15	12	0.0065
H228	17	13	20	31	14	10	24	14	12	10	14	18	20	23	11	11	9	12	18.2	14,18	15	11	0.0065
H255	16	13	19	30	13	10	25	12	12	10	14	20	23	24	12	11	13	13	16	15,16	17	11	0.0098
H248	17	12	20	29	13	9	26	13	11	10	14	18	20	23	12	11	11	14	17	15,17	15	12	0.0065

The haplotype diversity was calculated to be 0.997. This score shows that the Y-STR haplotypes under study have a very high level of variety. In other words, the population contains a large number of distinct haplotypes. Discrimination Capacity" (DC) now mean the degree to which a set of genetic markers can discriminate between individuals is measured by the DC. The ability to recognize unique haplotypes—those haplotypes that are encountered just once in the population is specifically at issue in this instance. The DC is determined by dividing the total number of haplotypes in the dataset by the number of unique haplotypes (Table 4).

Table 4: Genetic information of Y-STR haplotypes.

23Y-STR haplotype	
Sample size	383
Number of haplotypes	356
Unique haplotype	337
Repeating haplotype	19
Haplotype Diversity	0.997
Discrimination capacity	0.946

Average gene diversity (GD) is a measure used in genetic studies to assess the level of genetic variation within a population. In this context, the study population has an average gene diversity of 0.63981 (Table 5). This value indicates how genetically diverse or similar the individuals within this population are. The range of GD values within this population spans from 0.395062 to 0.93828. This range suggests that there is some variability in the genetic makeup of the individuals in the population, with some loci showing higher diversity and others showing lower diversity. The highest gene diversity was observed at the DYS385a/b loci, with a GD value of 0.93828. This means that these particular genetic markers (DYS385a/b) have a high degree of variation within the population; and the lowest gene diversity was found at the DYS392 locus, with a GD value of 0.395062. The NevGen Y-DNA Haplogroup online Predictor tool was used to identify the Y-haplogroups of male Iraqi Arabs. In this study, the most prevalent haplogroups were J1 (38%), J2 (20%), R1 (11%), E1 (10%), and G2 (5%). Additional expected haplogroups were also found, even though they were less common, as shown in Table 6.

Table 5: Gene Diversity in 23Y-STR haplotype of Iraqi Arab males

Locus	GD	Locus	GD
DYS576	0.756456	DYS570	0.793351
DYS3891	0.477233	DYS635	0.671155
DYS448	0.544598	DYS390	0.606141
DYS389II	0.689486	DYS439	0.576203
DYS19	0.563176	DYS392	0.395062
DYS391	0.538775	DYS635	0.720367
DYS481	0.83827	DYS393	0.570899
DYS549	0.630008	DYS458	0.860566
DYS533	0.613263	DYS456	0.696868
DYS438	0.574809	R-Y-GATA-H4	0.544079
DYS437	0.476782	DYS385a/b	0.93828

GD: Gene Diversity. The highest and lowest Gene Diversity in bold

Table 6: Haplogroups prediction for Iraqi Population.

Haplogroup	Number	Freq
J1	137	0.385
J2	72	0.202
R1	40	0.112
E1	37	0.104
G2	19	0.053
T	17	0.048
L1	6	0.017
Q	5	0.014
I2	4	0.011
R2	4	0.011
G1	4	0.011
Unsupported sub-clade	3	0.008
H1	2	0.006
N1	1	0.003
O2	1	0.003
I1	1	0.003
C2	1	0.003
O1	1	0.003
E2	1	0.003
	356	1.000

DISCUSSION

The Y-STR haplotypes, which are distinctive sets of genetic markers found on the Y chromosome, were examined in the context of this study. The analysis of these Y-STR haplotypes' diversity in the Iraqi Arab population was the main goal of the study. Y-STR haplotype databases are helpful for examining differences within a given population as well as between other demographic groupings and establishing databases, which is

essential for the forensic importance and reliability of Y-STR evidence [21]. The PowerPlex Y-23 kit contains more Y-STR loci than other Y-STR kits, which was done to improve the capacity to identify between individuals. It is a common option for use in forensic investigations and demographic research due to its strengthened discriminative power [22]. The result revealed 212 genetic variations (alleles) spread across 23 distinct locations on the Y chromosome known as Y-STR loci [Table 1]. A significant genetic variation within the population is indicated by polymorphism in these loci, which is important for different forensic and genetic research. The DYS385a/b locus had the most genetic diversity, with a total of 55 different alleles and highest gene diversity for this locus was 0.93828, indicating that different individuals within the population have distinct genetic profiles at these loci. These results are consistent with previous study published on the Arab and Kurdish populations in Iraq [23]. Conversely, the lowest gene diversity was found at the DYS392 locus, with a GD value of 0.395062. This suggests that the genetic variation at the DYS392 locus is relatively low within the population with 4 alleles, meaning that most individuals in the population have very similar genetic profiles at this specific genetic marker [24]. The observation that the GD values align with the polymorphism findings in the study indicates consistency in the genetic diversity patterns. In this study, it appears that the GD values correspond to the polymorphism levels observed at these genetic markers in the population. The fact that the GD values align with the study's polymorphism findings suggests that the genetic diversity patterns are consistent. In this situation, it appears that the GD values match the polymorphism levels seen at these genetic markers in the population. Polymorphism is the existence of several genetic variations (alleles) at a single genetic locus [25]. The micro-variant alleles "11" found in the DYS392 locus exhibited the highest frequency (0.770) and were widely prevalent throughout the Iraqi population. The aforementioned data holds significant relevance within the field of genetic research as it facilitates comprehension of the genetic composition and variability exhibited by a given population; this information plays a crucial role in population genetics and forensic investigations. They enable researchers to gain insights into genetic variety and facilitate the identification of individuals through the analysis of their DNA profiles [26]. A total of 356 haplotypes were identified as unique within the dataset, indicating their absence of repetition. This observation is regarded as highly satisfying and serves as evidence of the efficacy of the PowerPlex Y 23 kit, a genetic analysis tool. Additionally, multiple distinct haplotypes were found to occur more than once in the dataset. The correlation between a decrease in genetic diversity at specific genetic loci and an increased incidence of inbreeding has been established. Inbreeding observed in numerous regions of Iraq results in a gradual reduction in genetic variety over an extended period [27, 26, 29, 30]. The study revealed that the prevailing genetic groups, referred to as haplogroups, were J1 and J2, constituting 38% and 20% of the sample, respectively. Whit Athey's Haplogroup Predictor was used to identify Y-haplogroups in this research. According to the results, J1 was shown to be the most common haplogroup among Iraqi Arabs (38%). This fits in with what has been seen before. Many people from different parts of West Asia, North Africa, the Horn of Africa, Southern Europe, Central Asia, and South Asia share DNA with the haplogroup J1. People who speak other Semitic languages, most notably Arabic, are more likely to be familiar with it [27, 28, 31].

CONCLUSION

The current study successfully constructed a comprehensive database consisting of 23 Y-STR loci that were specifically designed and optimized for the Iraqi Arab community. The utilization of haplotype patterns and haplogroup predictions has demonstrated significant efficacy in understanding the geographic origins of individuals. The DYS385a/b loci showed the highest level of genetic diversity, followed by DYS458. In contrast, DYS391 exhibited the most limited extent of genetic diversity. The microvariant allele "11" has been observed to be the predominant allele at the DYS392 locus throughout the Iraqi population. The prevalence of the J1 haplogroup was noted to be highest among Iraqi Arabs. The result has significance for the development of a comprehensive population database, providing crucial insights for the fields of population genetics and forensic research. In addition, these tools enhance comprehension of genetic variability and allow for the differentiation of individuals through analysis of their DNA profiles.

Funding

There was no outside funding used to carry out the investigation.

Ethical approval

The study obtained ethical clearance by the institutional Ethics Committee of the Forensic DNA Centre for Research and Training at Al-Nahrain University, situated in Jadriya, Baghdad, Iraq. The approval was officially obtained on the: 10/9 /2022, and is linked to a distinct reference number (43).

Consent for publication

The consent to publish had been taken from each participant in this work.

Abbreviations

Y-STR: Y-chromosome of Short Tandem Repeats, PCR: polymerase chain reaction, AMOVA; Analysis of Molecular Variance, YHRD: STR Haplotype Reference Database, MDS: multidimensional scaling, DC: Discrimination Capacity, GD: Gene Diversity.

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