Multiplexed Microsphere Suspension-Array Assay for Urine Mitochondrial DNA Typing by C-Stretch Length in Hypervariable Regions

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Abstract

Background: The standard method for personal identification and verification of urine samples in doping control is short tandem repeat (STR) analysis using nuclear DNA (nDNA). The DNA concentration of urine is very low and decreases under most conditions used for sample storage; therefore, the amount of DNA from cryopreserved urine samples may be insufficient for STR analysis. We aimed to establish a multiplexed assay for urine mitochondrial DNA typing containing only trace amounts of DNA, particularly for Japanese populations.

Methods: A multiplexed suspension-array assay using oligo-tagged microspheres (Luminex MagPlex-TAG) was developed to measure C-stretch length in hypervariable region 1 (HV1) and 2 (HV2), five single nucleotide polymorphisms (SNPs), and one polymorphic indel. Based on these SNPs and the indel, the Japanese population can be classified into five major haplogroups (D4, B, M7a, A, D5). The assay was applied to DNA samples from urine cryopreserved for 1 - 1.5 years (n = 63) and fresh blood (n = 150).

Results: The assay with blood DNA enabled Japanese subjects to be categorized into 62 types, exhibiting a discriminatory power of 0.960. The detection limit for cryopreserved urine was 0.005 ng of nDNA. Profiling of blood and urine pairs revealed that 5 of 63 pairs showed different C-stretch patterns in HV1 or HV2.

Conclusions: The assay described here yields valuable information in terms of the verification of urine sources employing only trace amounts of recovered DNA. However, blood cannot be used as a reference sample.

Keywords: Suspension-array assay; MagPlex-TAG microspheres; Mitochondrial DNA; C-stretch length; Hypervariable region; Urine sample

Introduction

Urine samples collected via noninvasive methods are widely used in regulation, control, and monitoring of doping and are cryopreserved for a maximum of 10 years for re-examination. When substitution or manipulation of a urine sample is suspected, the standard method for personal identification and verification of urine samples in doping control is short tandem repeat (STR) analysis using nuclear DNA (nDNA) [1-5]. Urine nDNA concentration is very low and inconsistent [6-11] and decreases under most conditions used for sample preservation [7-13]. The success rate of STR analysis using 10 mL of unprocessed urine samples cryopreserved for 1 - 1.5 years has been reported to be < 100% [14]. Therefore, a simple assay exhibiting enhanced analytical sensitivity with trace amounts of nDNA has been concluded to be required in doping control.

Mitochondrial DNA (mtDNA) represents an advantageous target for personal identification and verification through its significantly elevated cellular copy number (100 - 1,000-fold compared with a single diploid representation of nDNA, which enables measurements even after prolonged storage). However, one disadvantage of mtDNA analysis is the greatly diminished inter-individual discriminatory capacity due to the maternal mode of inheritance.

The major haplogroups of the mtDNA characteristics of Japanese subjects were reported to be D4, B, M7a, A, and D5 [15], which can be classified by the assessment of five single nucleotide polymorphisms (SNPs; 3010G>A, 4386T>C, 5178C>A, 8794C>T, and 10397A>G) and one deletion (8272C>del) [16]. C-stretch length mutation was observed with a high frequency in the hypervariable regions, 16184-16193 (HV1C) and 303-315 (HV2C), of the mtDNA control region [17, 18]. Different HV2C profiles were observed, as a result of not following maternal inheritance, between mother and child [19-21], among siblings [22, 23], and among various tissues, including hair shafts from the same individual [24-27]. HV1C and HV2C mutants in cervical cells have been reported to be stable for 10 - 20 years in a longitudinal retrospective study [28].

Suspension-array technology (Luminex xMAP) [29] has been applied to a simultaneous detection method of various...
SNPs and indels in mtDNA [30-32]. Recently, the construction of a suspension-array assay was simplified using MagPlex-TAG microspheres (MagPlex-TAG), which are magnetic, differently-fluorescent beads covalently coupled to unique 24-mer “anti-TAG” oligonucleotide sequences [33, 34]. The assay with MagPlex-TAG consists of three steps: polymerase chain reaction (PCR), allele-specific primer extension (ASPE), and then hybridization between the ASPE product and MagPlex-TAG.

In this study, we report the development of a multiplexed suspension-array assay using MagPlex-TAG for mtDNA typing by measurement of C-stretch sequences in HV1C and HV2C as well as analysis of five SNPs and one deletion. The discrimination power and detection limit of the assay were evaluated by its application to blood and urine samples, respectively. In addition, mtDNA profiles in urine and blood pairs were compared to determine whether blood could be used as an appropriate reference sample.

### Materials and Methods

Nucleotide position numbers used are consistent with the revised Cambridge Reference Sequence [35].

### Materials

Streptavidin-R-PE, dGTP, dATP, dTTP, Biotin-dCTP, and Fast SYBR® Green Master Mix were purchased from Life Technologies Japan (Tokyo, Japan); ExoSAP-IT® from Affymetrix (Cleveland, Ohio, USA); Puregene® Blood Core kit and HotStar Taq® Master Mix kit from QIAGEN (Hilden, Germany); NucleoSpin® gDNA Clean-up and NucleoSpin® gDNA Clean-up XS kits from Macherey-Nagel (Duren, Germany); Novagen® human gDNA from Merck (Darmstadt, Germany); TaKaRa Taq™ Hot Start version from Takara Bio (Shiga, Japan); and MagPlex®-TAG microspheres from Luminex Japan (Tokyo, Japan). D17Z1 primers were synthesized by Life Technologies Japan, and mtDNA-specific PCR primers and ASPE primers by Integrated DNA Technologies (Austin, Texas, USA). All other chemicals and solvents were of the highest quality commercially available.

### Table 1. PCR Primer Sequences

| Target | Forward | Reverse | Amplicon size (bp) |
|--------|---------|---------|-------------------|
| SNP and deletion [30] | | | |
| 3010G>A | AGAGCGGGCGATAACACAGCA | GATCACGTTAGGACTTTAATCGTTGA | 369 |
| 4386T>C | TCGAACCCATCCCTGAGAATCC | GCTAATTTTGTCTCAGGGTTTGTTTAT | 234 |
| 5178C>A | CAGCATCGGACTATTTAGTATAT | GTGGCAGTTAGGAATGAGTAG | 269 |
| 8272C>del | GGGGTATACTACGGTCAATGCTC | TCATTTTGTCTCAGGGTTTGTTTAT | 378 |
| 8794C>T | CAAATATCTCATCAAACACCGACTA | CAGGGCTATTGGTTGAATGAGTAG | 367 |
| 10397A>G | ACCCCTACCATGAGCCTTACAA | GTGAGATGGTAAATGCTAGTATAAATAT | 239 |
| C-stretch [36] | | | |
| HV1C | CAAATGGGCCTGTCCTTGTA | TGTGGGGAATTTGATTTC | 519 |
| HV2C | GGTCTATACCCCTATTAACCAC | CTGTTAACAGTGTCATACCGCCA | 380 |
| Others | | | |
| ND1 [38] | TTCTAATCGCAATGGCCATTCCT | AAGGGTTGTGTAGCCCGTGAT | 109 |
| D17Z1 [37] | TTTTGCAGGATCTACAAGTGGA | AAGAGGTCTACATGTCCTCCTTG | 207 |

### Samples and DNA extraction

This study was reviewed and approved by the Review Board of the Japan Chemical Analysis Center, Anti-doping Research Laboratory. Permission was granted to use materials collected solely for research study. Japanese males aged 20 - 35 years, who were recruited from outside our laboratory, provided informed consent.

Urine and blood samples were collected at the Yana-gibashi-Clinical Trial Center (Tokyo, Japan) in January 2011, following confirmation that these subjects were negative for HIV and hepatitis B and C. DNA (nDNA and mtDNA) was extracted as previously described [14]. Briefly, DNA was extracted both from urine cryopreserved for 1 - 1.5 years (hereafter referred to as cryopreserved urine) and fresh blood using the Puregene Blood Core kit. Extracted urine DNA was purified with a NucleoSpin gDNA Clean-up kit or NucleoSpin gDNA Clean-up XS kit. DNA samples from cryopreserved urine (n = 63) and fresh blood (n = 150) were stored at -20 °C and 4 °C, respectively, until use. Due to the difficulty of quantifying mtDNA, DNA quantity was determined using the nDNA quantitative value rather than mtDNA.

### Multiplexed suspension-array assay with MagPlex-TAG

The assay was established following the manufacturer’s protocols [33] and modified to achieve appropriate sensitivity. The
Applied Biosystems Veriti Thermal Cycler (Life Technologies, Japan) was used with 8-tube PCR strips (0.2 mL) for PCR, ExoSAP-IT treatment, and ASPE, and with 96-well PCR plates (0.1 mL) for hybridization.

**Multiplexed PCR**

Multiplexed PCR was performed in a total volume of 20 µL with HotStar Taq Master Mix (10 µL, 1 unit HotStar Taq DNA Polymerase in PCR buffer with 3 mM MgCl₂, and 400 µM of each deoxynucleotide (dNTP)), 200 nM each of 8-primer pair (Table 1) [30, 36], and a DNA sample solution. Thermal cycling conditions were 95°C for 10 min; 35 cycles of 94°C for 30 s, 55°C for 30 s, and 72°C for 1 min; and 72°C for 10 min. The sizes of amplified products were confirmed by electrophoresis on 2% agarose gels in preliminary experiments.

Unincorporated PCR primers and dNTPs were removed from the reaction mixture after PCR by treatment with 3.5 µL of ExoSAP-IT at 37°C for 30 min or 90 min, followed by incubation at 80°C for 15 min.

**Multiplexed ASPE**

ASPE primers were designed to flank the targeted base at the 3’ ends and a TAG sequence (24-mer) at the 5’ termini. Sequences with melting temperatures (Tm) between 51°C - 56°C were selected from 18 - 24-mer sequences with the targeted base at the 3’ ends [33]. Tm of the sequences was calculated using the Oligonucleotide Properties Calculator (http://www.basic.northwestern.edu/biotools/oligocalc.html). A TAG sequence suitable for the selected sequence and the corresponding MagPlex-TAG were selected on advice provided by Luminex.

Two ASPE primers targeting each wild-type and mutant base were prepared for five SNPs and one deletion (Table 2). C-stretch sequences [18] were classified into nine groups for HV1C and six groups for HV2C according to the number of C before T or the absence of T in these regions. An ASPE primer was prepared for each group (Tables 3, 4). A total of 27 ASPE primers were used, of which 10 are for five SNPs, two for the deletion, nine for HV1C, and six for HV2C.

Multiplexed ASPE was conducted in 30 µL of PCR buffer (10 mM Tris-HCl, pH 8.3, 50 mM KCl, and 1.5 mM MgCl₂) containing 5 µM each of dATP, dTTP, dGTP, and biotin-dCTP, 1.125 U of TaKaRa Taq Hot Start DNA polymerase, 25 nM of each 27 ASPE primers and 7.5 µL of ExoSAP-IT-treated PCR products. Thermal cycling conditions were 32 cycles of 94°C for 30 s, 55°C for 30 s, and 72°C for 1 min.

**Hybridization**

Overall, 10 or 25 µL of ASPE products was hybridized with 2,000 particles of each of the 27 different MagPlex-TAGs in 50 µL of hybridization buffer (HB, 0.1 M Tris-HCl, pH 8.0, containing 0.2 M NaCl and 0.08% Triton X-100). The mixtures were incubated at 96°C for 90 s followed by 37°C for 30 min. After washing twice using a 96-well magnet plate, the hybridized MagPlex-TAGs were incubated with 75 µL of HB containing streptavidin-R-PE (2 µg/mL) at 37°C for 15 min. PE fluorescence of 50 µL of the mixture was measured as units of Median Fluorescence Intensity (MFI) at 37°C for a maximum of 200 s using a Luminex 200 analyzer (Austin, TX, USA) with the associated software (xPONENT®). Haplogroups were defined as A for 8794T, B for 8272del, D4 for 3010A and 5178A, D5 for 5178A and 10397G, and M7a for 4386C. A C-stretch pattern composed of more than two groups was expressed in descending order of MFI of the group.

**Discrimination power of the assay**

Blood DNA samples containing approximately 5 ng of nDNA were assayed under the same conditions as for ExoSAP-IT, with a treatment time of 30 min, and an ASPE product volume

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**Table 2. ASPE Primer Sequences for Detection of Five SNPs and One Deletion in MtDNA**

| Sequence (5’ to 3’) | Primer Sequence |
|---------------------|-----------------|
| 3010G>A            | (CTATCATTATATCTCTTTCTCTAATT) ATGTGGATCAGGACATCCCG |
| 4386T>C            | (CTCAAAACATTTACATTCTCTTT) CCTTACTTTAGGTTGGGGTGTG |
| 5178C>A            | (ATAAAGACACTCTTTACEATTT) ATATCTGCACCTGAAAAGGC |
| 8272C>del          | (AATTTCTCTCTCTTTAACAAT) CCCTATAGACCCCTCTC |
| 8794C>T            | (TTAAATAAATTCTCTCTTTCTCTCTTA) TGGGTGGTGGTTGTAATAAGT |
| 10397A>G           | (ATCTCAATTAAATTACACACAAA) TGACTACAAAAAGGATTAGACT |

Sequences in parentheses indicate 24-mer TAG sequences.
used for hybridization of 10 µL (n = 150). A positive result was defined as a value greater than four times the MFI obtained from the corresponding negative control. Discrimination power was calculated using the Discriminatory Power Calculator (http://insilico.ehu.es/mini_tools/discriminatory_power/index.php).

Detection limit of the assay for cryopreserved urine

Cryopreserved urine DNA samples containing < 0.15 ng of nDNA were assayed under the same conditions as for ExoSAP-IT, with a treatment time of 90 min, and an ASPE product volume used for hybridization of 25 µL. A positive result was defined as a value greater than two times the MFI obtained from the corresponding negative control.

Urine DNA samples containing 0.1 or 0.03 ng of nDNA were diluted to 0.005, 0.0025, and 0.00125 ng (n = 10). The lowest DNA quantity displaying the same mtDNA type as that for 0.1 or 0.03 ng was judged to be the detection limit.

Quantification of nDNA

Real-time PCR was performed using the Applied Biosystems ViiA 7 system (Life Technologies Japan) with Fast SYBR Green Master Mix containing 200 nM each of nDNA-specific D17Z1 primers (Table 1) [37] and 1 µL of standard human gDNA (0.0012 - 12 ng/µL) or DNA solution in a final volume of 15 µL, as previously described [14].

MtDNA quantity of cryopreserved urine relative to blood

DNA samples, diluted serially four times (1 µL), were amplified with 200 nM each of mtDNA-specific NADH-ubiquinone oxidoreductase chain 1 (ND1) primers (Table 1) [38] under the same conditions as for nDNA quantification to obtain Ct values. Separately, the nDNA quantities of the serially diluted DNA samples were measured as described above. The Y-axis intercept of an approximately straight line from the scatter plot of the Ct values of ND1 and the logarithm of nDNA quantities, Ct value of ND1/nDNA (1 ng), was obtained using Microsoft Excel 2010 tool. The Ct values of ND1/nDNA (1 ng) of urine and blood are referred to as Ct
erine and Ct
blood, respectively. The mtDNA quantity of cryopreserved urine (n = 5) relative to blood (n = 5) was calculated from the equation 2^{ΔCt} (ΔCt = Ct
blood - Ct
erine). Results were expressed as mean ± SD. Statistical analysis was performed using the Mann-Whitney U test, with P < 0.05 regarded as significant.

Results

Discrimination power of the assay

Japanese subjects (n = 150) were classified into 62 types con-

Table 4. ASPE Primer Sequences for Detection of HV2C Groups in MtDNA

| Group | Sequence (5’ to 3’) |
|-------|----------------------|
| 6CT   | (TCTCTTTAAACCATCTTCAAAACAAATAA)AAAAAAATTTTCACAAACCCCCCCCT |
| 7CT   | (CTCTCTTTAAACCATCTTCAAAACAAATAA)AAAAAAATTTTCACAAACCCCCCCCT |
| 8CT   | (CACTTCTCTTCTACATCTTCAAAACAAATAA)AAAAAAATTTTCACAAACCCCCCCCT |
| 9CT   | (CATACATCTCTTCTACATCTTCAAAACAAATAA)AAAAAAATTTTCACAAACCCCCCCCT |
| 11C   | (TACATCTCTTCTACATCTTCAAAACAAATAA)AAAAAAATTTTCACAAACCCCCCCCT |

Sequences in parentheses indicate 24-mer TAG sequences. Position at 316 is G.
sisting of a combination of C-stretch patterns of HV1C and HV2C and the major haplogroups of Japanese populations by blood DNA analysis (Table 5). Type 5CT>10C (HV1C)/7CT (HV2C)/D4 was identified as representing the highest frequency (20/150, 13.3%). The discrimination power of the assay was determined to be 0.960.

Haplogroup frequencies were 37.3% for D4, 10.7% for B, 9.3% for M7a, 7.3% for A, 3.3% for D5, and 32.0% for “others” (Table 5). There were 15 patterns, including 2.0% of “others” identified for HV1C (Table 6). The frequency of patterns consisting of one, two, or three groups was 10.7%, 57.3%, or 30.0%, respectively. Pattern 5CT>10C was the most frequent pattern observed (48.7%), followed by 10C>11C>9C (22.0%). A particular pattern was related to a particular haplogroup. The frequencies of 5CT>10C in D4 and M7a were 76.8% (43/56) and 71.4% (10/14), respectively. Furthermore, all patterns in B haplogroup were composed of 9C, 10C, and/or 11C. The discriminatory power of the assay that incorporated HV1C typing and haplotyping was 0.884.

There were 12 patterns, including 8.7% of “others” identified for HV2C (Table 7). The frequency of patterns consisting of one or two groups was 74.7% or 16.7%, respectively. Patterns 7CT and 8CT displayed similar high frequencies of 36.7% and 37.3%, respectively. As a result, 89.3% of Japanese subjects showed 7CT or/and 8CT. There was no particular pattern related to a particular haplogroup with > 70% frequency. The discrimination power of the assay that incorporated HV2C typing and haplotyping was 0.917.

Comparison of mtDNA type between urine and blood

MtDNA types of cryopreserved urine samples were compared with those of the corresponding blood samples (n = 63 pairs). Haplogroups of urine and blood were completely matched in all 63 pairs. However, C-stretch patterns were different in 18 pairs. The difference in 13 pairs resulted from the presence or absence of the group with the lowest MFI, a value near the defined positive and negative boundaries (Table 8). The remaining 5 pairs showed a difference in the MFI order of the same composing groups (Table 9). All patterns showing differences in HV1C consisted of three poly C-sequences (9C, 10C, and 11C). In particular, all urine samples corresponding to blood samples with 11C>10C>9C showed patterns different from blood. One pair showed a difference in HV2C pattern consisting of 6CT and 7CT. 10C>9C>11C (HV1C) and 7CT>6CT (HV2C) in urine were patterns not found in any of 150 blood samples (Tables 5-7).

Detection limit of the assay with MagPlex-TAG for cryopreserved urine

$C_{\text{urine}}$ and $C_{\text{blood}}$ values were 17.7 ± 0.5 and 21.0 ± 0.4, respectively (P < 0.05). MtDNA quantity of cryopreserved urine relative to blood calculated from $2^{\Delta C_{t}}$, in which $\Delta C_{t} = C_{\text{blood}} - C_{\text{urine}} = 3.3$, was 9.85. The result indicated that the mtDNA quantity was approximately 10 times more in cryopreserved urine than in blood, when both nDNA quantities were the same.

Analysis of DNA from cryopreserved urine samples revealed that the lowest nDNA quantity displaying the same mtDNA type as that derived from nDNA (0.1 or 0.03 ng) was 0.0025 or 0.005 ng (Table 10). We concluded that the detection limit of the assay was 0.005 ng when cryopreserved urine samples were used.

Discussion

We established a multiplexed suspension-array assay employing MagPlex-TAG for Japanese populations in which C-stretch length mutations in HV1C and HV2C as well as five SNPs and one deletion in mtDNA could be identified. Using a combination of two C-stretch typings and major haplotyping, Japanese subjects (n = 150) were classified into 62 types by blood DNA analysis. The discriminatory power of the assay was 0.960. The detection limit of the assay was 0.005 ng when DNA from cryopreserved urine samples was used. Our assay represents a potential method for the personal identification or verification of urine sample contributors containing only trace amounts of DNA following long-term cryopreservation in doping control.

According to the established assay, Japanese subjects were classified into 62 types combining six haplogroups, 15 HV1C patterns, and 12 HV2C patterns. Our haplogroup frequencies were consistent with results from 1,312 Japanese individuals: D4 (32.6%), B (13.3%), M7a (7.5%), A (6.9%), D5 (4.8%), and “others” (34.9%) [15]. Currently, there are no reports of C-stretch patterns consisting of multiple groups in Japanese populations. Therefore, the frequencies of patterns with 5CT (HV1C) or 8CT (HV2C) as the main group were summed to compare with previously published findings provided by direct sequencing [18, 39]. The calculated frequencies, 56.7% (5CT) and 48.7% (8CT), were close to the results of previous reports of 58.6% and 45.0% [18] and 62.6% and 40.8% [39]. Although we have not confirmed C-stretch patterns by other methods, these results demonstrated that our assay seems to exhibit high reliability in C-stretch typing.

HV2C typing contributed more to improving the discriminatory power of this assay involving haplotyping than HV1C characterization. The basis of this observation is likely to be the strong relation between an HV1C pattern and D4, M7a or B haplogroups. Our results that all HV1C patterns of B haplogroup consist only of groups with 16189C (9C, 10C, 11C) were consistent with the coexistence of 16189C with B haplogroup in Asian, African, Nicobarese, and Indonesian populations [40, 41]. The existence of a strong relation between HV1C pattern and haplogroup impeded the improvement of the discriminatory power of the assay.

There were two types of differences in C-stretch pattern observed between urine and the corresponding blood in some subjects. The first was different due to absence of the group with the lowest MFI in the pattern consisting of groups showing the same MFI order (Table 8). Except for one case showing 8CT↔other, the lowest MFI value was near the defined positive and negative boundaries, even when the group was
Table 5. Frequency of Type Consisting of HV1C Pattern, HV2C Pattern, and Major Haplogroups for Japanese Populations

| HV1C | HV2C | A | B | D4 | D5 | M7a | Others | Frequency N (%) |
|------|------|---|---|----|----|-----|--------|-----------------|
| 0CT  | 8CT  | 1 |   |    |    |     |        | 1 (0.7)         |
| 3CT  | 7CT  | 2 | 1 |    |    |     |        | 3 (2.0)         |
|      | 8CT  | 4 |   |    |    |     |        | 4 (2.7)         |
|      | 8CT>9CT | 1 |   |    |    |     |        | 1 (0.7)         |
| 5CT  | 7CT  | 1 |   |    |    |     |        | 1 (0.7)         |
|      | 8CT  | 2 |   |    |    |     |        | 2 (1.3)         |
|      | Others | 1 |   |    |    |     |        | 1 (0.7)         |
| 10C  | 7CT  | 1 |   |    |    |     |        | 1 (0.7)         |
| 11C  | 8CT>7CT | 1 |   |    |    |     |        | 1 (0.7)         |
|      | 13C>11C | 1 |   |    |    |     |        | 1 (0.7)         |
| 3CT>4CT | 9CT>8CT | 1 |   |    |    |     |        | 1 (0.7)         |
| 5CT>8CT | 7CT  | 1 |   |    |    |     |        | 1 (0.7)         |
| 5CT>10C | 7CT  | 20| 4 | 6  |    |     |        | 30 (20.0)       |
|      | 8CT  | 2 | 15| 2  | 10 |     |        | 29 (19.3)       |
|      | 7CT>8CT | 1 |   |    |    |     |        | 1 (0.7)         |
|      | 8CT>7CT | 6 | 1 |    |    |     |        | 7 (4.7)         |
|      | 8CT>9CT | 2 |   |    |    |     |        | 2 (1.3)         |
|      | 9CT>8CT | 1 | 1 |    |    |     |        | 2 (1.3)         |
|      | Others | 1 |   |    |    |     |        | 1 (0.7)         |
| 10C>5CT | 8CT  | 2 |   |    |    |     |        | 2 (1.3)         |
| 8CT>7CT | 1 |   |    |    |    |     |        | 1 (0.7)         |
| 10C>11C | 6CT  | 1 |   |    |    |     |        | 1 (0.7)         |
|      | 8CT  | 1 | 1 | 1  |    |     |        | 3 (2.0)         |
|      | Others | 2 |   |    |    |     |        | 4 (2.7)         |
| 5CT>10C>9C | 7CT | 1 | 1 | 1  |    |     |        | 3 (2.0)         |
| 8CT>7CT | 1 | 2 |   |    |    |     |        | 3 (2.0)         |
| 8CT>9CT | 1 |   |    |    |    |     |        | 1 (0.7)         |
| 10C>3CT>9C | 8CT | 1 |   |    |    |     |        | 1 (0.7)         |
| 10C>11C>9C | 7CT | 1 | 2 | 1  | 9  |    |        | 13 (8.7)        |
|      | 8CT  | 5 | 2 | 1  | 4  |    |        | 12 (8.0)        |
|      | 6CT>7CT | 1 |   |    |    |     |        | 1 (0.7)         |
|      | 7CT>13C | 1 |   |    |    |     |        | 1 (0.7)         |
|      | 8CT>7CT | 1 |   |    |    |     |        | 1 (0.7)         |
|      | Others | 3 |   |    |    |     |        | 5 (3.3)         |
| 11C>10C>9C | 7CT | 1 |   |    |    |     |        | 1 (0.7)         |
|      | 8CT  | 1 |   |    |    |     |        | 1 (0.7)         |
|      | 13C>9CT | 1 |   |    |    |     |        | 1 (0.7)         |
|      | Others | 1 |   |    |    |     |        | 1 (0.7)         |
| Others | 7CT  | 2 |   |    |    |     |        | 2 (1.3)         |
|      | 8CT  | 1 |   |    |    |     |        | 1 (0.7)         |
| N    | 11   | 16 | 56 | 5  | 14 | 48 |        | 150             |
| %    | 7.3  | 10.7| 37.3| 3.3| 9.3| 32.0|       |                 |
Table 6. Frequency of HV1C Pattern in the Major Haplogroups for Japanese Populations

| HV1C | A | B | D4 | D5 | M7a | Others | Frequency N (%) |
|------|---|---|----|----|-----|--------|-----------------|
| 0CT  |   |   | 1  |    |     | 1      | 1 (0.7)         |
| 3CT  | 7 | 1 |    |    |     | 8      | 8 (5.3)         |
| 5CT  | 3 | 1 | 1  |    |     | 4      | 4 (2.7)         |
| 10C  |   | 1 |    |    | 1   | 1      | 1 (0.7)         |
| 11C  | 1 | 1 | 2  |    | 1   | 2      | 2 (1.3)         |
| 3CT>4CT | |   |    |    |     | 1      | 1 (0.7)         |
| 5CT>8CT | |   | 1  |    | 1   | 1      | 1 (0.7)         |
| 5CT>10C | 3 | 43| 10 | 17 | 17  | 73     | 73 (48.7)       |
| 10C>5CT | 1 | 2 | 2  |    | 2   | 3      | 3 (2.0)         |
| 10C>11C | 3 | 2 | 3  |    | 3   | 8      | 8 (5.3)         |
| 5CT>10C>9C | | 3 | 3 | 1 | 1 | 7      | 7 (4.7)         |
| 10C>3CT>9C | | 1 |    |    |     | 1      | 1 (0.7)         |
| 10C>11C>9C | | 10| 4 | 2 | 1  | 16     | 16 (22.0)       |
| 11C>10C>9C | | 2 | 2 | 2 | 2  | 4      | 4 (2.7)         |
| Others | 1 | 2 | 2  |    | 2   | 3      | 3 (2.0)         |

Table 7. Frequency of HV2C Pattern in the Major Haplogroups for Japanese Populations

| HV2C | A | B | D4 | D5 | M7a | Others | Frequency N (%) |
|------|---|---|----|----|-----|--------|-----------------|
| 6CT  | 1 | 7 | 7  | 3  | 2   | 18     | 18 (37.3)       |
| 7CT  | 7 | 7 | 3  | 6  | 1   | 20     | 20 (36.7)       |
| 8CT  | 2 | 2 | 25 | 6  | 14  | 55     | 55 (36.7)       |
| 6CT>7CT | 1 | 8 | 3  | 2  | 2   | 13     | 13 (8.7)        |
| 7CT>8CT | 1 | 1 | 1  | 2  | 2   | 4      | 4 (2.7)         |
| 8CT>9CT | 1 | 2 | 2  |    | 2   | 3      | 3 (2.0)         |
| 9CT>8CT | 1 | 1 | 1  | 2  | 2   | 4      | 4 (2.7)         |
| 13CT>9CT | 1 | 1 | 1  |    | 1   | 1      | 1 (0.7)         |
| 13CT>11C | 1 | 1 | 1  | 1  | 1   | 1      | 1 (0.7)         |
| Others | 5 | 2 | 1  | 1  | 5   | 13     | 13 (8.7)        |

Table 8. Differences in C-stretch Patterns Between 13 Pairs of Urine and Blood Samples

| N  | Urine  | Blood  | Urine  | Blood  |
|----|--------|--------|--------|--------|
|    | HV1C   | HV2C   |        |        |
| 1  | 0CT>5CT| 0CT    |        |        |
| 2  | 5CT>10C| 5CT    |        |        |
| 1  | 5CT>10C| 5CT>10C>9C| 8CT   | 8CT>7CT|
| 1  | 10C>11C| 10C>11C>9C| 6CT>7CT| 6CT    |
| 1  | 10C>11C>9C| 10C>11C| 7CT    | 7CT>13C|
| 1  | 11C    | Others | 8CT    | 8CT>7CT|
| 1  | 8CT    | Others |        |        |
determined as negative, it could be distinguish from the other negative groups. The second was due to a different MFI order in the same composing groups. This type of difference was only observed in HV1C pattern consisting of 9C, 10C, and 11C as well as HV2C pattern consisting of 6CT and 7CT, in which there were patterns present only in urine but not in blood. These results indicated that blood cannot be used as a reference sample in our assay for the personal identity validation of urine samples.

The detection limit of 0.005 ng for cryopreserved urine samples was considerably less than the quantity of nDNA necessary for successful STR analysis (0.6 ng) [14]. An approximately 10 times higher ratio of mtDNA/nDNA in cryopreserved urine than in blood was believed to arise from the relatively diminished stability of nDNA under the preservation conditions of urine samples [7-13]. This apparent robustness of mtDNA in cryopreserved urine samples allowed the measurement of two C-stretch sequences and major haplogroups for Japanese populations at lower nDNA concentrations.

In conclusion, this study demonstrated that C-stretch typing, in addition to haplotyping, of samples containing only trace amounts of DNA was possible by a multiplexed suspension-array assay using MagPlex-TAG. Our assay can provide valuable information to ensure the authenticity of urine samples. However, confirmation of the long-term stability of the C-stretch sequence is required for our assay to be adopted in doping control. An advantage of our assay is the ease of introduction of new polymorphic targets of mtDNA, because a maximum of 80 types of MagPlex-TAG are commercially available. Therefore, we intend to develop a more comprehensive C-stretch sequence assay to obtain useful knowledge on the lifetime stability of C-stretch sequences in various samples.

Acknowledgments
The authors acknowledge the valuable support of Ms. Fujiko Sekine from Luminex, Japan.

Conflict of Interest
None.

Grant Support
This study was supported by a grant from the Japan Sport Council of 2014.

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Table 9. Differences in C-stretch Patterns Between 5 Pairs of Urine And Blood Samples

| N | Urine | HV1C | Blood | HV2C |
|---|---|---|---|---|
| 2 | 10C>11C>9C | 11C>10C>9C | |
| 1 | 10C>9C>11C | 11C>10C>9C | |
| 1 | 10C>9C>11C | 10C>11C>9C | |
| 1 | 7CT>6CT | 6CT>7CT | |

Table 10. Detection Limits of the Assay for DNA From Urine Samples Cryopreserved for 1 - 1.5 Years

| Sample No. | Haplogroup | MtDNA type | Detection limit ng as nDNA |
|---|---|---|---|
| 1 | A | 5CT>10C | 0.005 |
| 2 | A | 3CT | 0.0025 |
| 3 | B | 10C>11C>9C | 0.005 |
| 4 | B | 10C>11C>9C | 0.0025 |
| 5 | D4 | 5CT | Others | 0.005 |
| 6 | D4 | 5CT>10C | 7CT | 0.0025 |
| 7 | D5 | 10C>11C>9C | 7CT>6CT | 0.005 |
| 8 | D5 | 10C>11C>9C | 6CT>7CT | 0.0025 |
| 9 | M7a | 10C>11C>9C | 7CT | 0.005 |
| 10 | Others | 10C>9C>11C | 7CT | 0.0025 |
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