

A deafness-associated tRNA^{Asp} mutation alters the m¹G37 modification, aminoacylation and stability of tRNA^{Asp} and mitochondrial function

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ABSTRACT

In this report, we investigated the pathogenic mechanism underlying the deafness-associated mitochondrial(mt) tRNA^{Asp} 7551A > G mutation. The m.7551A > G mutation is localized at a highly conserved nucleotide(A37), adjacent (3') to the anticodon, which is important for the fidelity of codon recognition and stabilization in functional tRNAs. It was anticipated that the m.7551A > G mutation altered the structure and function of mt-tRNA^{Asp}. The primer extension assay demonstrated that the m.7551A > G mutation created the m¹G37 modification of mt-tRNA^{Asp}. Using cybrid cell lines generated by transferring mitochondria from lymphoblastoid cell lines derived from a Chinese family into mitochondrial DNA(mtDNA)-less (ρ^0) cells, we demonstrated the significant decreases in the efficiency of aminoacylation and steady-state level of mt-tRNA^{Asp} in mutant cybrids, compared with control cybrids. A failure in metabolism of mt-tRNA^{Asp} caused the variable reductions in mtDNA-encoded polypeptides in mutant cybrids. Impaired mitochondrial translation led to the respiratory phenotype in mutant cybrids. The respiratory deficiency lowered mitochondrial adenosine triphosphate production and increased the production of oxidative reactive species in mutant cybrids. Our data demonstrated that mitochondrial dysfunc-

tions caused by the m.7551A > G mutation are associated with deafness. Our findings may provide new insights into the pathophysiology of maternally transmitted deafness that was manifested by altered nucleotide modification of mitochondrial tRNA.

INTRODUCTION

The defects in mitochondrial protein synthesis have been associated with both syndromic deafness (hearing loss with other medical problems such as diabetes) and nonsyndromic deafness (hearing loss is the only obvious medical problem) (1–5). The mitochondrial translation machinery composed of 2 rRNAs and 22 tRNAs, encoded by mitochondrial DNA (mtDNA), and more than 150 proteins (ribosomal proteins, ribosomal assembly proteins, aminoacyl-tRNA synthetases, tRNA-modifying enzymes, and several initiation, elongation and termination factors), encoded by nuclear genes (6–8). Mutations in the *LARS2* and *NARS2* encoding mitochondrial leucyl-tRNA synthetase and asparaginyl-tRNA synthetase have been associated with deafness, respectively (9–10). The mtDNA mutations have been shown to be the important causes of both syndromic and nonsyndromic deafness (3–5). Of these, the m.1555A > G and m.1494C > T mutations in the 12S rRNA gene have been associated with both aminoglycoside-induced and nonsyndromic deafness in many families worldwide (3,4,11,12). The most prevalent mtDNA mutations associated with syndromic deafness are the MELAS-associated m.3243A > G mutation in the mt-

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tRNA^{Leu(UUR)} gene (13) and MERRF-associated m.8344A > G mutation in the mt-tRNA^{Lys} gene (14), while the non-syndromic deafness-associated mtDNA mutations included the mt-tRNA^{Ser(UCN)} 7445A > G, 7472insC, 7505T > C and 7511T > C, mt-tRNA^{His} 12201T > C, mt-tRNA^{Gly} 10003T > C and mt-tRNA^{Ile} 4295A > G mutations (15–21). These mt-tRNA mutations altered their structures and functions, including the processing of the mt-tRNA from the primary transcripts, stability of the folded secondary structure, the charging of the mt-tRNA, or the codon-anticodon interaction in the process of translation (5,22,23). The m.7445A > G mutation altered the processing of the 3' end mt-tRNA^{Ser(UCN)} precursor (24), the m.7511T > C mutations affected the stability of mt-tRNA^{Ser(UCN)} (25) and m.12201T > C mutation altered the aminoacylation of mt-tRNA^{His} (20). However, the pathophysiology of these tRNA mutations remains poorly understood.

As the part of a genetic screening program for deafness in a cohort of 2651 Han Chinese affected subjects, we identified the novel m.7551A > G mutation in the mt-tRNA^{Asp} gene in one Han Chinese pedigrees with maternal transmission of nonsyndromic deafness (19,26). As shown in Figure 1, the m.7551A > G mutation is localized at a highly conserved nucleotide (A37), adjacent (3') to the anticodon of mt-tRNA^{Asp} (22,23). There were no modifications of i⁶A37 or t⁶A37 in the human mitochondrial tRNA^{Asp} (27), although the nucleotides at position 37 (A or G) of tRNAs are often modified by methylthiolation (28–29). The modifications at position 37 were shown to contribute to the high fidelity of codon recognition and to the structural formation and stabilization of functional tRNAs (30–33). Thus, the substitution of A with G at position 37 of the mt-tRNA^{Asp} may introduce the m¹G37 modification of this tRNA, thereby altering the structure and function of mt-tRNA^{Asp}. In particular, the mutation may affect the aminoacylation capacity and stability of this mt-tRNA and then impair mitochondrial translation. It was also proposed that an impairment of mitochondrial translation caused by the mt-tRNA mutation alters the respiration, production of adenosine triphosphate (ATP) and reactive oxygen species (ROS). To further investigate the pathogenic mechanism of the m.7551A > G mutation, cybrid cell lines were constructed by transferring mitochondria from lymphoblastoid cell lines derived from an affected matrilineal relative in a Chinese family carrying the mtDNA mutation and from a control individual lacking the mtDNA mutation, into human mtDNA-less (ρ^0) cells (34–35). First, we examined if the m.7551A > G mutation created the m¹G37 modification of mt-tRNA^{Asp} by using primer extension. These resultant cybrid cell lines were then assessed for the effects of the mtDNA mutation on the aminoacylation capacity and stability of this mt-tRNA, mitochondrial translation, respiration and the production of ATP and ROS as well as mitochondrial membrane potential.

MATERIALS AND METHODS

Cell lines and culture conditions

Immortalized lymphoblastoid cell lines were generated from one affected matrilineal relative (III-6) of the Chinese

family carrying the m.7551A > G mutation (26) and one genetically unrelated Chinese control individual (A4) belonging to the same mtDNA haplogroup but lacking the mutation (Supplementary Table S1) (36). These cell lines were grown in RPMI 1640 medium with 10% fetal bovine serum. The bromodeoxyuridine (BrdU) resistant 143B.TK⁻ cell line was grown in Dulbecco's Modified Eagle Medium (Life Technologies) (containing 4.5 mg of glucose and 0.11 mg pyruvate/ml), supplemented with 100 μ g of BrdU/ml and 5% fetal bovine serum. The mtDNA-less ρ^0 206 cell line, derived from 143B.TK⁻ (34,35) was grown under the same conditions as the parental line, except for the addition of 50 μ g of uridine/ml. Transformation by cytoplasts of mtDNA-less ρ^0 206 cells using one affected subject (III-6) and one control individual (A4) was performed as described elsewhere (34,35,37). All cybrid cell lines constructed with enucleated lymphoblastoid cell lines were maintained in the same medium as the 143B.TK⁻ cell line. An analysis for the presence and level of the m.7551A > G mutation was carried out as described previously (26). The quantification of mtDNA copy number from different cybrids was performed as detailed elsewhere (38).

Detection of m¹G37 modification in mt-tRNA^{Asp} using primer extension

To determine the nucleotide positions of m¹G modification of mt-tRNA^{Asp}, a primer extension experiment was performed by a modified procedure, as detailed elsewhere (39,40). Mitochondria were isolated from three mutant and three control cybrids ($\sim 2.0 \times 10^8$ cells), as described previously (41). Total mitochondrial RNAs were isolated using TOTALLY RNATM kit (Ambion), as described previously. A DNA primer (5'-TGGUAAAGATATATAGGATTTAGCCTATAAT-3') complementary to the 3' end of the mt-tRNA^{Asp} was 5' end labeled with non-radioactive digoxin (DIG). Prime-script II 1st Strand cDNA Synthesis Kit (TAKARA) was used for reverse transcription with DIG-labeled oligodeoxynucleotide probe specific for the mt-tRNA^{Asp} by using total mt-RNA as templates. Extension reactions were carried out as detailed previously (39,40). Finally, samples were applied onto 15% polyacrylamide-7 M urea gel electrophoresis and electroblotted onto a positively charged nylon membrane. Quantification of density in each band was performed as detailed previously (40).

Mitochondrial tRNA Northern analysis

Two micrograms of total mitochondrial RNA were electrophoresed through a 10% polyacrylamide/7 M urea gel in Tris-borate-EDTA buffer (after heating the sample at 65°C for 10 min). The gels were then electroblotted onto a positively charged nylon membrane (Roche) for the hybridization analysis with DIG-labeled oligodeoxynucleotide probes for mt-tRNA^{Asp}, mt-tRNA^{Thr}, mt-tRNA^{Leu(CUN)}, mt-tRNA^{Lys} and mt-tRNA^{Ser(AGY)} were as detailed previously (20,24,25,42,43). DIG-labeled oligodeoxynucleotides were generated by using DIG oligonucleotide Tailing kit (Roche). The hybridization and quantification of density in each band were performed as detailed previously (20,24,25).

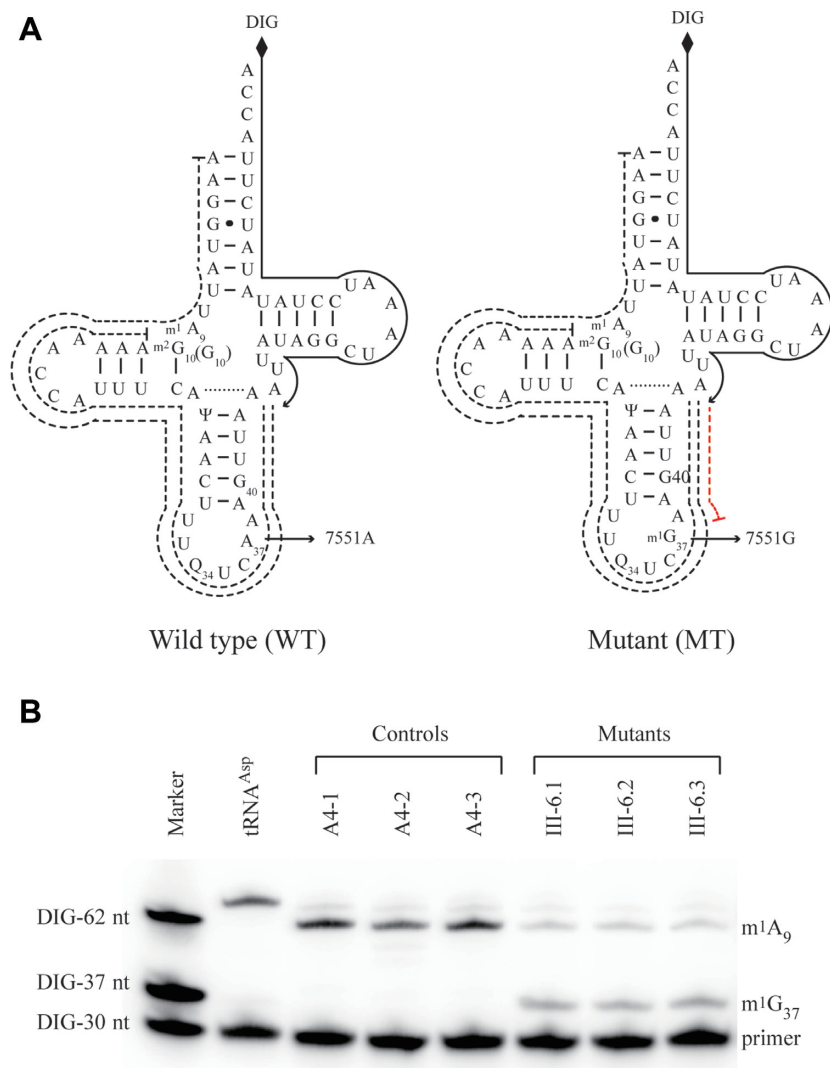


Figure 1. The m.7551A > G mutation introduced the methylation of G37 in mt-tRNA^{Asp}. **(A)** Schematic of methylation shown in the cloverleaf structures of human mitochondrial tRNA^{Asp}. An arrow denotes the location of the m.7551A > G mutation. Solid lines represent the DIG-labeled oligonucleotide probe specific for mt-tRNA^{Asp}. Broken lines represent the potential stops of primer extension caused by m¹A or m¹G modification. **(B)** Primer extension demonstrated the creation of m¹G37 in the mt-tRNA^{Asp} carrying the m.7511A > G mutation. One microgram of mitochondrial RNA from three control cybrids and three mutant cybrids were used for this investigation. DIG-labeled oligonucleotide specific for mt-tRNA^{Asp} was then annealed and the primer extensions were performed. The primer extension termination products caused by m¹A or m¹G modification are shown.

For the aminoacylation assays, total mitochondrial RNAs were isolated under acid conditions, and 2 µg of total mitochondrial RNAs was electrophoresed at 4°C through an acid (pH 3.0) 10% polyacrylamide–7 M urea gel to separate the charged and uncharged tRNA as detailed elsewhere (43,44). To further distinguish non-aminoacylated tRNA from aminoacylated tRNA, samples of tRNAs were deacylated by being heated for 10 min at 60°C at pH 8.3 and then run in parallel (35,44). The gels were then electroblotted onto a positively charged nylon membrane (Roche) for the hybridization analysis with oligodeoxynucleotide probes as described above. Quantification of density in each band was performed as detailed previously (20).

Western blot analysis

Western blotting analysis was performed as detailed previously (20,43). The antibodies used for this investigation were from Abcam [anti GAPDH (ab72655), p.MT-ND1 (ab74257), p.MT-ND5 (ab92624) and p.MT-ATP6 (ab101908), p.MT-CO2 (ab110258)], Santa Cruz Biotechnology [p.MT-ND4 (sc-20499-R) and p.MT-ND6 (sc-20667)] and Proteintech [p.MT-CYTB (55090-1-AP)]. Peroxidase Affini Pure goat anti-mouse IgG and goat anti-rabbit IgG (Jackson) were used as a secondary antibody and protein signals were detected using the ECL system (CW-BIO). Quantification of density in each band was performed as detailed previously (20,43).

Assays of activities of respiratory complexes

The enzymatic activities of complex I, II, III and IV were assayed as detailed elsewhere (43,45,46).

Measurements of oxygen consumption

The rates of oxygen consumption (OCR) in cybrid cell lines were measured with a Seahorse Bioscience XF-96 extracellular flux analyzer (Seahorse Bioscience), as detailed previously (20,43,47).

ATP measurements

The Cell Titer-Glo[®] Luminescent Cell Viability Assay kit (Promega) was used for the measurement of cellular and mitochondrial ATP levels, according to the modified manufacturer's instructions (20,43).

Assessment of mitochondrial membrane potential

Mitochondrial membrane potential was assessed with JC-10 Assay Kit-Microplate (Abcam) following general manufacturer's recommendations with some modifications, as detailed elsewhere (20,35).

Measurement of ROS production

ROS measurements were performed following the procedures detailed previously (20,43,48,49).

Computer analysis

Statistical analysis was carried out using the unpaired, two-tailed Student's *t*-test contained in the Microsoft-Excel program or Macintosh (version 2007). Differences were considered significant at a $P < 0.05$.

RESULTS

The description of one hearing-impaired Chinese pedigree and derived cell lines

The pedigree of the Chinese family with maternally inherited deafness was previously described (19,26). Six of 12 matrilineal relatives suffered from the variable degree of hearing impairment (two with mild hearing loss, two with moderate hearing loss and two with profound hearing loss). The age-at-onset of hearing loss ranged from 5 to 45 years, with the average of 26 years. These matrilineal relatives exhibited no other clinical abnormalities, including cardiac failure, muscular diseases, visual failure and neurological disorders. Immortalized lymphoblastoid cell lines were derived from one affected matrilineal relative carrying the m.7551A > G mutation, with profound hearing loss (III-6; male, 21 years) and from one genetically unrelated hearing normal individual lacking the m.7551A > G mutation belong to the same mtDNA haplogroup (A4, male, 20 years). The lymphoblastoid cells were enucleated, and subsequently fused to a large excess of mtDNA-less human $\rho^{\circ}206$ cells, derived from the 143B.TK- cell line (34,35). These cybrid clones were isolated by growing the fusion mixtures in the selective Dulbecco's

Modified Eagle Medium, containing BrdU and lacking uridine. Between 25 and 45 days after fusion, 10–15 presumptive mitochondrial cybrids derived from each donor cell line were isolated, and subsequently analyzed for the presence and level of the m.7551A > G mutation (26). The results confirmed the absence of the mtDNA mutation in the control clones and its presence in homoplasmy in all cybrids derived from the mutant cell line (Supplementary Figure S1). Three cybrids derived from each donor cell line with similar mtDNA copy numbers were used for the biochemical characterization described below.

The m.7551A > G mutation created the m¹G37 modification of mt-tRNA^{Asp}

There were no modifications of i⁶A37 or t⁶A37 detected in the human mitochondrial tRNA^{Asp} (27). To investigate if the m.7551A>G mutation produced the m¹G37 modification of mt-tRNA^{Asp}, we subjected mitochondrial RNAs from mutant and control cybrid cell lines to the reverse transcription with DIG-labeled oligonucleotide probe specific for mt-tRNA^{Asp} (Figure 1A). This results in a stop band one residue 3' to the methylation on 15% polyacrylamide gel. As shown in Figure 1B, the m¹G37 modification was present in mt-tRNA^{Asp} derived from three mutant cell lines, while the m¹G37 modification was not detected in the mt-tRNA^{Asp} derived from three control cell lines. Furthermore, the m¹A9 modifications were detected in mt-tRNA^{Asp} obtained from both control and mutant cell lines.

Reduction in the steady-state level of mt-tRNA^{Asp}

To examine if the m.7551A > G mutation affects the stability of mt-tRNA^{Asp}, we subjected mitochondrial RNAs from cybrids cell lines to Northern blots and hybridized them with DIG-labeled oligodeoxynucleotide probes for mt-tRNA^{Asp}, mt-tRNA^{Thr}, mt-tRNA^{Leu(CUN)}, mt-tRNA^{Lys} and mt-tRNA^{Ser(AGY)}, respectively. As shown in Figure 2A, the amount of mt-tRNA^{Asp} in three mutant cell lines were markedly decreased, compared with those in three control cybrid cell lines. For comparison, the average level of each mt-tRNA^{Asp} in control or mutant cell lines was normalized to the average levels in the same cell line for reference mt-tRNA^{Thr}, mt-tRNA^{Leu(CUN)}, mt-tRNA^{Lys} and mt-tRNA^{Ser(AGY)}, respectively. As shown in Figure 2B, the average steady-state level of mt-tRNA^{Asp} was significantly decreased in the mutant cells, compared with the wild-type cells. In particular, the average levels of mt-tRNA^{Asp} in the mutant cybrid cell lines were among ~58%, 52%, 64% and 57% of average values of three control cybrids after normalization to mt-tRNA^{Thr}, mt-tRNA^{Leu(CUN)}, mt-tRNA^{Lys} and mt-tRNA^{Ser(AGY)} ($P < 0.0018$ to 0.0005), respectively.

Deficient aminoacylation of mt-tRNA^{Asp}

We evaluated whether the m.7551A > G mutation alters the aminoacylation of mitochondrial tRNAs. For this purpose, the aminoacylation capacities of mt-tRNA^{Asp}, mt-tRNA^{Thr}, mt-tRNA^{Leu(CUN)}, mt-tRNA^{Lys} and mt-tRNA^{Ser(AGY)} in control and mutant cell lines were examined by the use of electrophoresis in an acidic

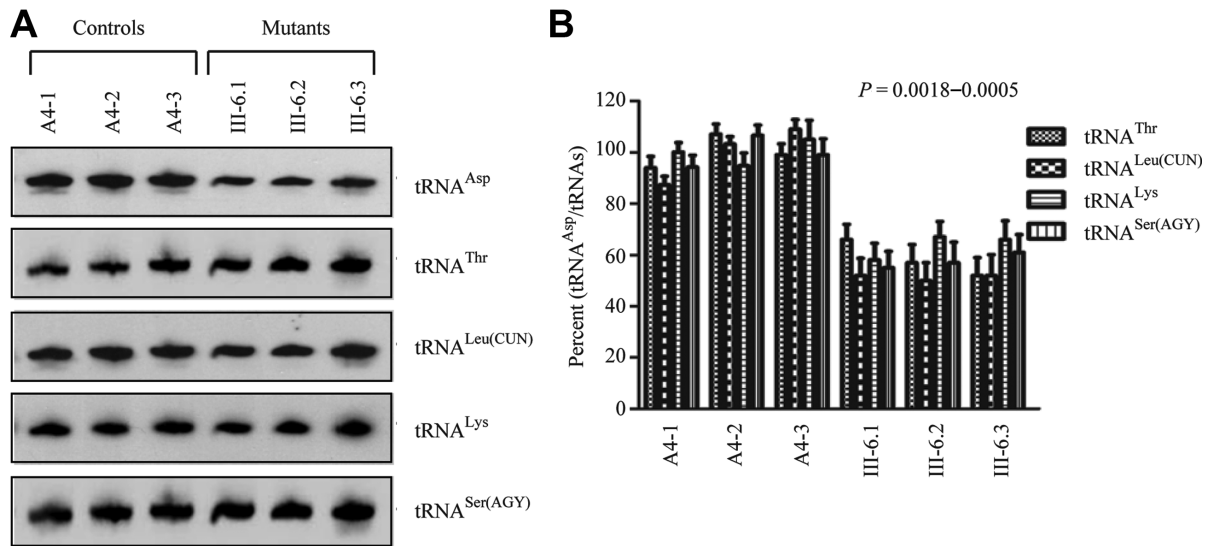


Figure 2. Northern blot analysis of mitochondrial tRNA. (A) Equal amounts of total mitochondrial RNA from various cell lines were electrophoresed through a denaturing polyacrylamide gel, electroblotted and hybridized with DIG-labeled oligonucleotide probes specific for the mt-tRNA^{Asp}, mt-tRNA^{Thr}, mt-tRNA^{Leu(CUN)}, mt-tRNA^{Lys} and mt-tRNA^{Ser(AGY)}, respectively. (B) Quantification of tRNA levels. Average relative mt-tRNA^{Asp} content per cell, was normalized to the average content per cell of mt-tRNA^{Thr}, mt-tRNA^{Leu(CUN)}, mt-tRNA^{Lys} and mt-tRNA^{Ser(AGY)} in three cybrid cell lines derived from one affected subject (III-6) carrying the m.7551A > G mutation and three cybrid cell lines derived from one Chinese control subject (A4). The values for the latter are expressed as percentages of the average values for the control cell lines. The calculations were based on three independent determinations of each tRNA content in each cell line and three determinations of the content of reference tRNA marker in each cell line. The error bars indicate two standard errors of the means. *P* indicates the significance, according to the *t*-test, of the differences between mutant and control cell lines.

polyacrylamide/urea gel system to separate uncharged tRNA species from the corresponding charged tRNA, electroblotting and hybridizing with the tRNA probes described above. As shown in Figure 3A, the upper band represents the charged tRNA, and the lower band represents uncharged tRNA. There were no obvious differences in electrophoretic mobility between the control and mutant cell lines. To further distinguish nonaminoacylated tRNA from aminoacylated tRNA, samples of tRNAs were deacylated by being heated for 10 min at 60°C at pH 8.3 and then run in parallel. As shown in Figure 3B, only one band (uncharged tRNA) was present in both mutant and control cell lines after deacylating. However, the efficiencies of aminoacylated mt-tRNA^{Asp} in these mutant cell lines carrying the m.7551A > G mutation varied from 74 to 84%, with average of 80%, relative to the average values of control cell lines (*P* = 0.0072) (Figure 3C). However, the levels of aminoacylation in mt-tRNA^{Thr}, mt-tRNA^{Leu(CUN)}, mt-tRNA^{Lys} and mt-tRNA^{Ser(AGY)} in the mutant cell lines were comparable with those in the control cell lines (Supplementary Figure S2).

Decreases in the level of mitochondrial proteins

To assess whether a failure in mt-tRNA^{Asp} metabolism caused by the m.7551A > G mutation impaired mitochondrial translation, a Western blot analysis was carried out to examine the steady state levels of seven subunits of respiratory complex in mutant and control cells with GAPDH as a loading control. As shown in Figure 4A, the levels of p.MT-CO1 and p.MT-CO2, subunits I and II of cytochrome *c* oxidase; p.MT-ND1, p.MT-ND4 and p.MT-ND5, subunits 1, 4 and 5 of NADH dehydrogenase; p.MT-ATP6, subunit 6 of

the H[±]ATPase; p.MT-CYTB, apocytochrome *b* exhibited the variable reductions in three mutant cell lines, as compared with those of three control cell lines. As shown in Figure 4B, the overall levels of seven mitochondrial translation products in the mutant cell lines ranged from ~58–76%, with an average of 65% (*P* = 0.0106), relative to the mean value measured in the control cell lines. Notably, the average levels of p.MT-ND1, p.MT-ND4, p.MT-ND5, p.MT-ND6, p.MT-CO2, p.MT-ATP6 and p.MT-CYTB in the mutant cells were 39%, 64%, 76%, 39%, 70%, 45% and 58% of the average values of control cells, respectively. However, the levels of polypeptide synthesis in mutant cells, relative to those in control cells, showed no significant correlation with either the number of codons or the proportion of aspartic acid residues (Supplementary Table S2).

Respiration defects in mutant cells

To evaluate if the m.7551A > G mutation alters cellular bioenergetics, we examined the OCR of three mutant cell lines carrying the m.7551A > G mutation and three control cell lines. As shown in Figure 5, the basal OCR in the mutant cell lines was ~59% (*P* = 0.047) relative to the mean value measured in the control cell lines. To investigate which of the enzyme complexes of the respiratory chain was affected in the mutant cell lines, OCR was measured after the sequential addition of oligomycin (to inhibit the ATP synthase), carbonyl cyanide *p*-(trifluoromethoxy) phenylhydrazone (to uncouple the mitochondrial inner membrane and allow for maximum electron flux through the ETC), rotenone (to inhibit complex I) and antimycin A (to inhibit complex III). The difference between the basal OCR and the drug-insensitive OCR yields the amount of ATP-linked

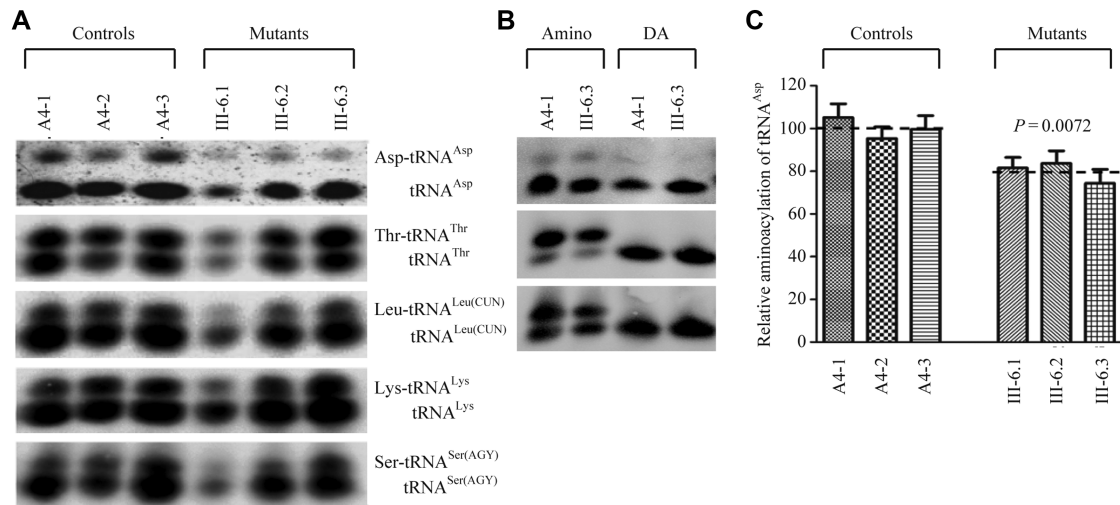


Figure 3. *In vivo* aminoacylation assays. (A) Two micrograms of total mitochondrial RNA purified from six cell lines under acid conditions were electrophoresed at 4°C through an acid (pH 3.0) 10% polyacrylamide-7 M urea gel, electroblotted and hybridized with a DIG-labeled oligonucleotide probe specific for the mt-tRNA^{Asp}. The blots were then stripped and rehybridized with mt-tRNA^{Thr}, mt-tRNA^{Leu(CUN)}, mt-tRNA^{Lys} and mt-tRNA^{Ser(AGY)}, respectively. (B) The samples from one control (A4.1) and mutant (III-6.3) cell lines were deacylated (DA) by heating for 10 min at 60°C at pH.8.3 and electrophoresed as above. Aminoacylation assays for mt-tRNA^{Asp} were carried out in parallel for aminoacylated and deacylated samples. (C) *In vivo* aminoacylated proportions of mt-tRNA^{Asp} in the mutant and controls. The calculations were based on three independent determinations. Graph details and symbols are explained in the legend to the Figure 2.

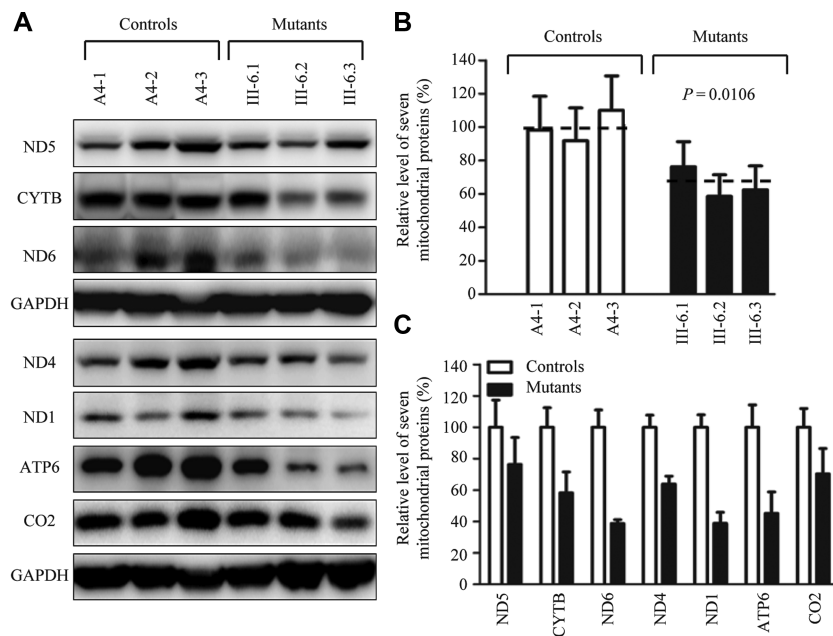


Figure 4. Western blot analysis of mitochondrial proteins. (A) Twenty micrograms of total cellular proteins from various cell lines were electrophoresed through a denaturing polyacrylamide gel, electroblotted and hybridized with 7 respiratory complex subunits in mutant and control cells with GAPDH as a loading control. p.MT-COI and p.MT-COII, indicate subunits I and II of cytochrome c oxidase; p.MT-ND1, p.MT-ND4 and p.MT-ND5, subunits 1, 4 and 5 of the reduced nicotinamide-adenine dinucleotide dehydrogenase; p.MT-ATP6, subunit 6 of the H±ATPase; and p.MT-CYTB, apocytochrome b. (B) Quantification of mitochondrial protein levels. Average relative p.MT-COI, p.MT-COII, p.MT-ND1, p.MT-ND4, p.MT-ND5, p.MT-ATP6 and p.MT-CYTB content per cell, normalized to the average content per cell of GAPDH in three mutant cell lines carrying the m.7551A > G mutation and three control cell lines lacking the mutation. The values for the latter are expressed as percentages of the average values for the control cell line. The calculations were based on three independent determinations. Graph details and symbols are explained in the legend to Figure 2.

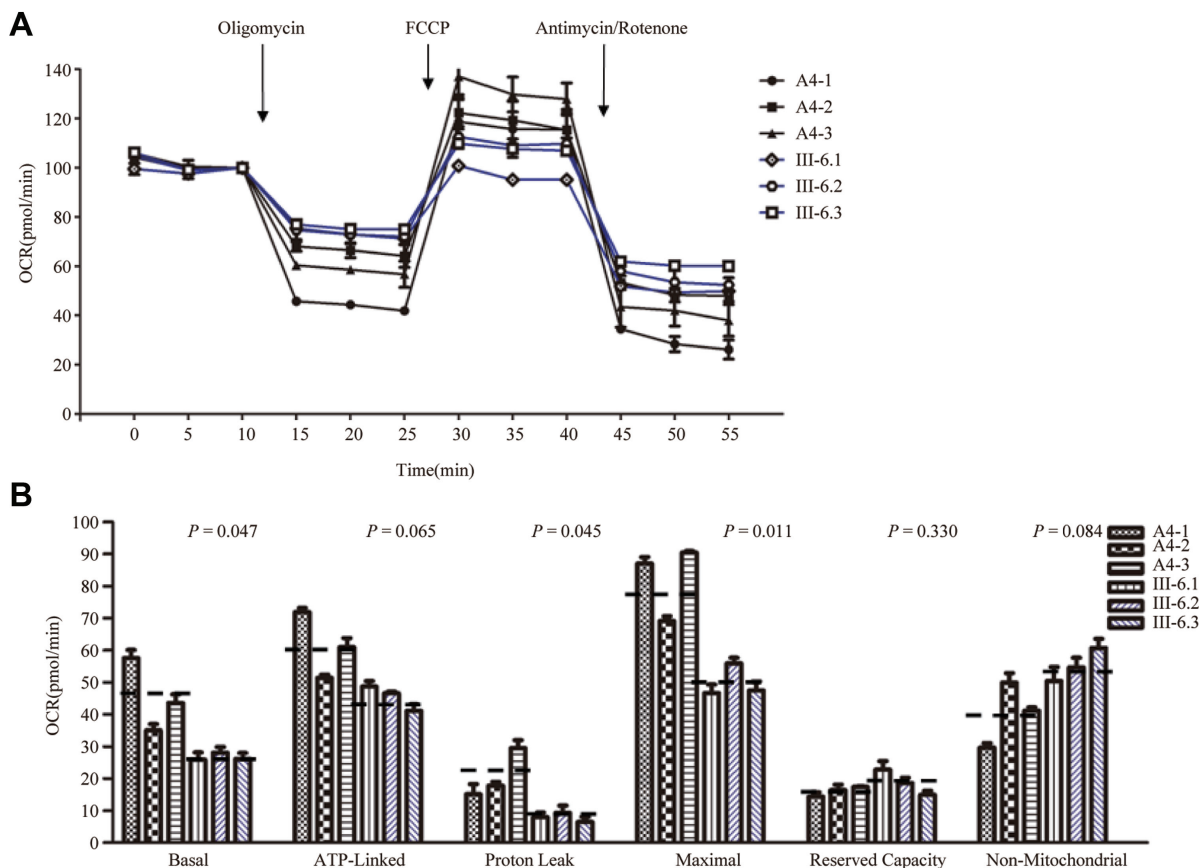


Figure 5. Respiration assays. (A) An analysis of O₂ consumption in the various cell lines using different inhibitors. The rates of O₂ (OCR) were first measured on 2×10^4 cells of each cell line under basal condition and then sequentially added to oligomycin (1.5 μ M), carbonyl cyanide p-(trifluoromethoxy) phenylhydrazone (FCCP) (0.5 μ M), rotenone (1 μ M) and antimycin A (1 μ M) at indicated times to determine different parameters of mitochondrial functions. (B) Graphs presented the ATP-linked OCR, proton leak OCR, maximal OCR, reserve capacity and non-mitochondrial OCR in mutant and control cell lines. Non-mitochondrial OCR was determined as the OCR after rotenone/antimycin A treatment. Basal OCR was determined as OCR before oligomycin minus OCR after rotenone/antimycin A. ATP-linked OCR was determined as OCR before oligomycin minus OCR after oligomycin. Proton leak was determined as Basal OCR minus ATP-linked OCR. Maximal was determined as the OCR after FCCP minus non-mitochondrial OCR. Reserve Capacity was defined as the difference between Maximal OCR after FCCP minus Basal OCR. The average of 4 determinations for each cell line is shown, the horizontal dashed lines represent the average value for each group. Graph details and symbols are explained in the legend to Figure 2.

OCR, proton leak OCR, maximal OCR, reserve capacity and non-mitochondrial OCR. As shown in Figure 5, the ATP-linked OCR, proton leak OCR, maximal OCR, reserve capacity and non-mitochondrial OCR in mutant cell lines were $\sim 74\%$, 38% , 61% , 117% and 137% , relative to the mean value measured in the control cell lines ($P = 0.065$, 0.045 , 0.011 , 0.330 and 0.084), respectively.

Reduced activities of respiratory chain complexes

To investigate the effect of the m.7551A > G mutation on the oxidative phosphorylation, we measured the activities of respiratory chain complexes by isolating mitochondria from mutant and control cell lines. Complex I (NADH ubiquinone oxidoreductase) activity was determined by following the oxidation of NADH with ubiquinone as the electron acceptor (47,48). Complex III (ubiquinone cytochrome c oxidoreductase) activity was measured as the reduction of cytochrome c (III) using d-ubiquinol-2 as the electron donor. The activity of complex IV (cytochrome c oxidase) was monitored by following the oxidation of cytochrome c

(II). As shown in Figure 6, the activity of complexes I, II, III and IV in the mutant cells carrying m.7551A > G mutation were 49%, 104%, 71% and 69% of the mean value measured in three control cell lines, respectively.

Reduced levels in mitochondrial ATP production

The capacity of oxidative phosphorylation in mutant and wild-type cell lines was examined by measuring the levels of cellular and mitochondrial ATP using a luciferin/luciferase assay. Populations of cells were incubated in the media in the presence of glucose, and 2-deoxy-D-glucose with pyruvate (20,50). As shown in Figure 7A, the levels of ATP production in mutant cells in the presence of glucose (total cellular levels of ATP) were 80% of those measured in the control cell lines. By contrast, as shown in Figure 7B, the levels of ATP production in mutant cell lines, in the presence of pyruvate and 2-deoxy-D-glucose to inhibit glycolysis (mitochondrial levels of ATP) ranged from 54 to 63%, with an average of 57% relative to the mean value measured in the control cell lines ($P = 0.0012$).

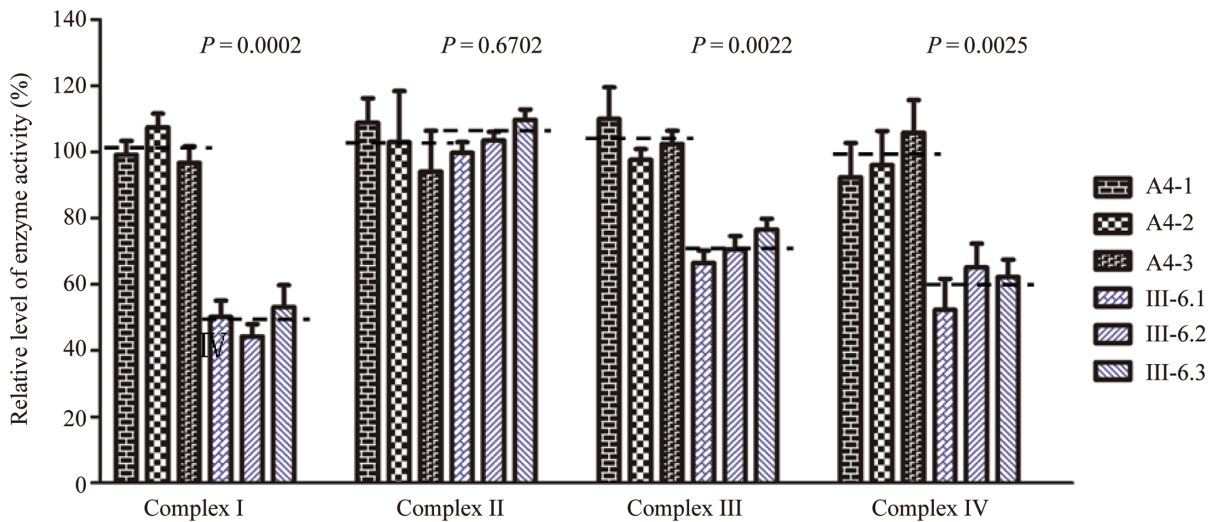


Figure 6. Enzymatic activities of respiratory chain complexes. The activities of respiratory chain complexes were investigated by enzymatic assays on complexes I, II, III and IV in mitochondria isolated from three mutant and three control cybrid cell lines. The calculations were based on four independent determinations. Graph details and symbols are explained in the legend to Figure 2.

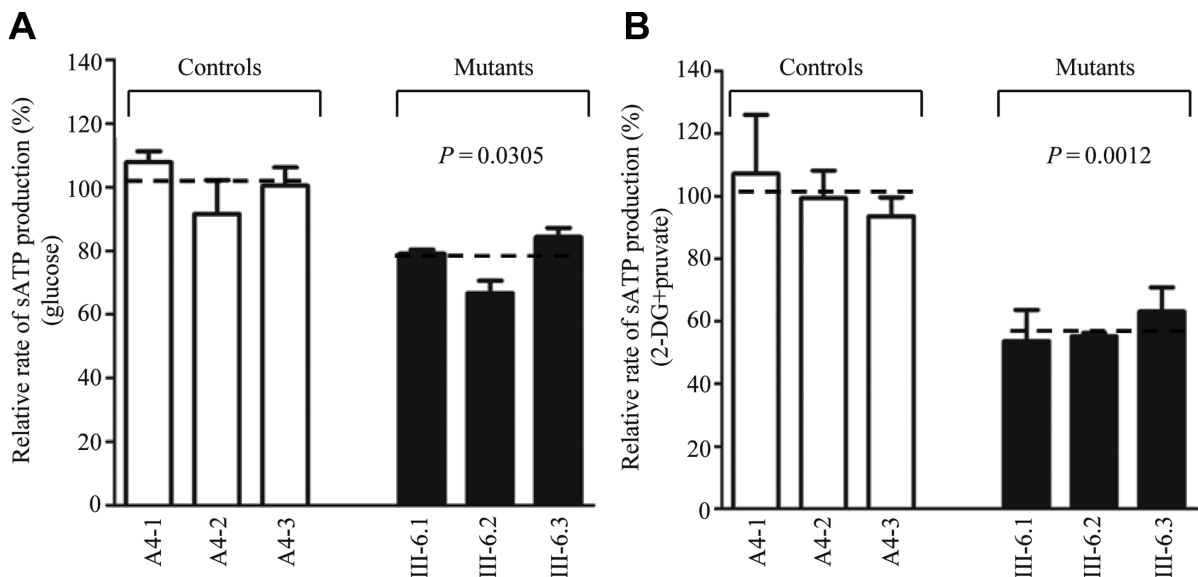


Figure 7. Measurement of cellular and mitochondrial ATP levels using bioluminescence assay. Cells were incubated with 10 mM glucose or 5 mM 2-deoxy-d-glucose plus 5 mM pyruvate to determine ATP generation under mitochondrial ATP synthesis. Average rates of ATP level per cell line and are shown: (A) ATP level in total cells; (B) ATP level in mitochondria. Six to seven determinations were made for each cell line. Graph details and symbols are explained in the legend to Figure 2.

Mild decreases in mitochondrial membrane potentials

The mitochondrial membrane potential ($\Delta\Psi_m$) changes were measured in mutant and control cell lines using the fluorescence probe JC-10 assay system. The ratios of fluorescence intensities $Ex/Em = 490/590$ and $490/530$ nm (FL590/FL530) were recorded to delineate the $\Delta\Psi_m$ level of each sample. The relative ratios of FL590/FL530 geometric mean between mutant and control cell lines were calculated to represent the level of $\Delta\Psi_m$. As shown in Figure 8, the levels of the $\Delta\Psi_m$ in the mutant cell lines carrying the m.7551A > G mutation ranged from 77.2% and 96.8%, with an average 86.2% ($P = 0.182$) of the mean value measured in the control cell lines. In contrast, the levels of

$\Delta\Psi_m$ in mutant cells in the presence of carbonyl cyanide-p-(trifluoromethoxy)phenylhydrazone were comparable with those measured in the control cell lines ($P = 0.766$).

The increase of ROS production

The levels of the ROS generation in the vital cells derived from three mutant cybrid cell lines carrying the m.7551A > G mutation and three control cybrid cell lines lacking the mutation were measured with flow cytometry under normal conditions and following H_2O_2 stimulation (20,49). Geometric mean intensity was recorded to measure the rate of ROS of each sample. The ratio of geometric mean intensity between unstimulated and stimulated with H_2O_2 in each cell

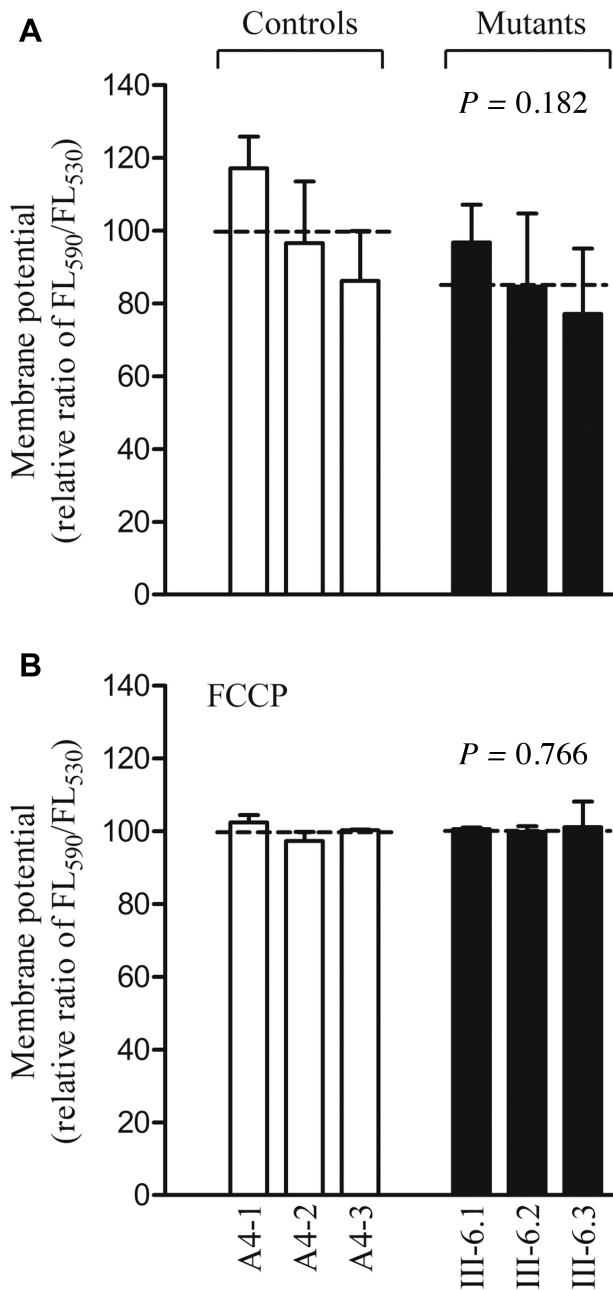


Figure 8. Mitochondrial membrane potential analysis. The mitochondrial membrane potential ($\Delta\Psi_m$) was measured in three mutant and three control cell lines using a fluorescence probe JC-10 assay system. The ratio of fluorescence intensities $Ex/Em = 490/590$ nm and $490/530$ nm (FL₅₉₀/FL₅₃₀) were recorded to delineate the $\Delta\Psi_m$ level of each sample. The relative ratios of FL₅₉₀/FL₅₃₀ geometric mean between mutant and control cell lines were calculated to reflect the level of $\Delta\Psi_m$. Relative ratio of JC-10 fluorescence intensities at $Ex/Em = 490/530$ nm and $490/590$ nm in (A) absence and (B) presence of $10 \mu\text{M}$ of FCCP. The average of 3–5 determinations for each cell line is shown. Graph details and symbols are explained in the legend to Figure 2.

line was calculated to delineate the reaction upon increasing level of ROS under oxidative stress. As shown in Figure 9, the levels of ROS generation in the mutant cell lines carrying the m.7551A > G mutation ranged from 126 and 140%,

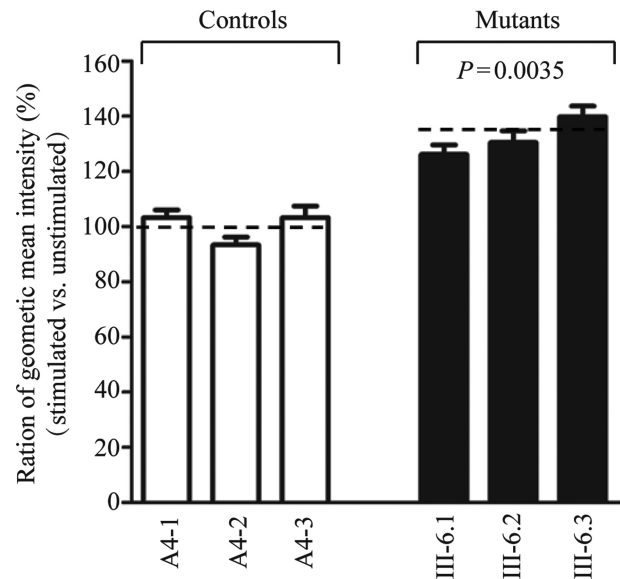


Figure 9. Ratio of geometric mean intensity between levels of the ROS generation in the vital cells with or without H_2O_2 stimulation. The rates of production in ROS from three mutant cell lines and three control cell lines were analyzed by BD-LSR II flow cytometer system with or without H_2O_2 stimulation. The relative ratio of intensity (stimulated versus unstimulated with H_2O_2) was calculated. The average of three determinations for each cell line is shown. Graph details and symbols are explained in the legend to Figure 2.

with an average 132% ($P = 0.0035$) of the mean value measured in the control cell lines.

DISCUSSION

In the present study, we investigated the pathogenetic mechanism of the deafness-associated m.7551A > G mutation in the mt-tRNA^{Asp} gene. The mutation is localized at a highly conserved nucleotide (A37), adjacent to the 3' end of the anticodon of the mt-tRNA^{Asp} (22,23). The nucleotide at position 37 of the tRNAs are often modified by methylthiolation (28,29). Modification at position A37 of some mitochondrial tRNAs such as mt-tRNA^{Lys} and mt-tRNA^{Ser(UCN)} was catalyzed by the modifying enzymes TRIT1 (MiaA in bacterial ortholog), while the modification of G37 in other tRNAs such as mt-tRNA^{Leu(CUN)} and mt-tRNA^{Gln} was catalyzed by the TRMT5 (TrmD in bacterial ortholog) (28–30). Modification at position 37 contributes to the high fidelity of codon recognition and to the structural formation and stabilization of functional tRNAs (28–31). However, the modification of A37 was not detected in human mt-tRNA^{Asp}, mt-tRNA^{Met} and mt-tRNA^{Leu(UUR)} (27,28). Thus, it was anticipated that the m.7551A > G mutation created the m¹G37 modification, thereby altering the structure and function of mt-tRNA^{Asp}. In this study, the primer extension experiment demonstrated that the m.7551A > G mutation produced the m¹G37 modification of mt-tRNA^{Asp}. The primary defect in this mutation appeared to affect the stability and aminoacylation of mutant mt-tRNA^{Asp} (50,51). Unlike the m.12201T > C mutation in the mt-tRNA^{His} and m.15927A > G mutation in the mt-tRNA^{Thr} (20,50), the m.7551A > G muta-

tion did not change conformation of mt-tRNA^{Asp}. However, the aminoacylation level of mt-tRNA^{Asp} but no other tRNAs were mildly decreased in mutant cell lines, as compared with controls. In fact, the A-to-G substitution at position 37 resulted in a 10-fold reduction in the amount of bacterial tRNA at the aminoacyl-tRNA binding site (52). A failure to aminoacylate tRNA properly then makes the mutant mt-tRNA^{Asp} to be metabolically less stable and more subject to degradation, thereby lowering the level of this tRNA, as in the case of m.3243A > G mutation in the mt-tRNA^{Leu(UUR)} (42,53). Alternatively, the m.7551A > G mutation likely affects the formation of functional structure of tRNAs and thus makes the tRNA be more unstable (22,23). In the present study, 42% reduction in the steady-state level of mt-tRNA^{Asp} observed in mutant cybrids was consistent with the previous observations in the lymphoblastoid cell lines carrying the m.4435A > G mutation in the tRNA^{Met} gene (32,33). However, the reduced level of mt-tRNA^{Asp} in mutant cells harboring the m.7551A > G mutation is indeed above the proposed threshold to produce a clinical phenotype associated with a mitochondrial tRNA mutation (20,53,54). This suggest that the m.7551A > G mutation alone is not sufficient to produce a clinical phenotype, as in the case of the deafness-associated m.1555A > G and m.1494C > T mutations in the 12S rRNA gene (37,55).

A failure in tRNA metabolism including inefficient aminoacylation or shortage of mt-tRNA^{Asp} could be the source of impaired mitochondrial translation of mtDNA encoding polypeptides and subsequent respiratory chain defects. In the present study, reduced levels of mitochondrial proteins (an average decrease of ~35%) were observed in mutant cybrid cell lines, as compared to the average levels in control cell lines. The reduced levels of variable mitochondrial protein detected in cybrids carrying the m.7551A > G mutation were comparable with the reduced rates of mitochondrial protein synthesis observed in lymphoblastoid cell lines carrying the m.4435A > G mutation (32,33). In mutant cell lines, a variable decrease in level of 7 mtDNA-encoded polypeptides was observed in each protein. However, polypeptide levels in mutant cell lines, relative to those in control cell lines, did not significantly correlate with either the number or proportion of aspartic acid codons, in contrast to what was previously shown in cell lines carrying the m.7445A > G mutation in the precursor of mt-tRNA^{Ser(UCN)} (24) or the m.8344A > G mutation in mt-tRNA^{Lys} (54). The lower level of mt-tRNA^{Asp} may contribute to the variable decrease of each polypeptide in mutant cell lines. Thus, the impaired synthesis of these mtDNA-encoding polypeptides may affect the activities of complex I, complex III and complex IV, respectively. Furthermore, impairment of mitochondrial protein synthesis were apparently responsible for the reduced rates in the basal OCR, or ATP-linked OCR, reserve capacity and maximal OCR among the control and mutant cell lines. These observations were clearly consistent with the critical role of mt-tRNA^{Asp} metabolic failure in producing their respiration defects, as in the cases of mutant cell lines carrying the deafness-associated mt-tRNA^{Ser(UCN)} 7445A > G, 7511T > C and mt-tRNA^{His} 12201T > C mutations (20,24,25).

The respiratory deficiency then affects the efficiency of mitochondrial ATP synthesis. In this investigation, 43% drop in mitochondrial ATP production in mutant cybrids carrying the m.7551A > G mutation may be caused by the defective activities of complexes I, III and IV. The reducing level was comparable with those in cells carrying the deafness-associated m.12201T > C and LHON-associated m.11778G > A mutations (20,43), but the decreasing level was much lower than those in cells carrying the mt-tRNA^{Lys} 8344A > G and mt-tRNA^{Leu(UUR)} 3243A > G mutations (56,57). Alternatively, the reduction in mitochondrial ATP production in mutant cells was likely a consequence of the decrease in the proton electrochemical potential gradient of mutant mitochondria (56). As a result, the hair cells carrying the mtDNA mutation may be particularly sensitive to increased ATP demand (3,4,58). Furthermore, the deficient activities of respiratory chain complexes caused by tRNA mutations often alter mitochondrial membrane potentials (59). In this study, only relatively mild reductions in mitochondrial membrane potential were observed in mutant cell lines carrying the m.7551A > G mutation, in contrast with those in the cell lines carrying the m.12201T > C mutation (20). The impairment of oxidative phosphorylation can lead to more electron leakage from electron transport chain, and in turn, elevate the production of ROS in mutant cells (60), thereby damaging mitochondrial and cellular proteins, lipids and nuclear acids (61). The hair cells and cochlear neurons may be preferentially involved because they are somehow exquisitely sensitive to subtle imbalance in cellular redox state or increased level of free radicals (61–63). However, the incomplete penetrance of deafness and mild biochemical defects indicated that the m.7551A > G mutation may be necessary evident but not sufficient to produce a clinical phenotype. The other genetic or epigenetic factors may contribute to the development of clinical phenotype in subjects carrying the m.7551A > G mutation (64,65). In particular, the hearing specific phenotype of this tRNA mutation may be attributed to the tissue-specificity of OXPHOS via tRNA modification or involvement of nuclear modifier genes (64,66–68).

In summary, our findings convincingly demonstrate the pathogenic mechanism leading to an impaired oxidative phosphorylation in cybrid cell lines carrying the deafness-associated m.7551A > G mutation in the mt-tRNA^{Asp} gene. The m.7551A > G mutation created the m¹G37 modification of mt-tRNA^{Asp}, but reduced the efficiency of aminoacylation and stability of this tRNA. A failure in tRNA^{Asp} metabolism led to the decreased synthesis of mtDNA encoding polypeptides and respiration. As a result, this respiratory deficiency reduced mitochondrial ATP production and increased the production of oxidative reactive species. The subsequent hearing loss may be involved in the participation of other genetic, epigenetic and environmental modifier factors. Thus, our findings may provide the new insights into the understanding of pathophysiology of maternally inherited deafness that was manifested by altered nucleotide modification of mitochondrial tRNA.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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REFERENCE

- Jacobs,H. (2003) Disorders of mitochondrial protein synthesis. *Hum. Mol. Genet.*, **12**, R293–R301.
- Rötig,A. (2011) Human diseases with impaired mitochondrial protein synthesis. *Biochim. Biophys. Acta*, **1807**, 1198–1205.
- Fischel-Ghodsian,N. (1999) Mitochondrial deafness mutations reviewed. *Hum. Mutat.*, **13**, 261–270.
- Guan,M.X. (2011) Mitochondrial 12S rRNA mutations associated with aminoglycoside ototoxicity. *Mitochondrion*, **11**, 237–245.
- Zheng,J., Ji,Y. and Guan,M.X. (2012) Mitochondrial tRNA mutations associated with deafness. *Mitochondrion*, **12**, 406–413.
- Andrews,R.M., Kubacka,I., Chinnery,P.F., Lightowlers,R.N., Turnbull,D.M. and Howell,N. (1999) Reanalysis and revision of the Cambridge reference sequence for human mitochondrial DNA. *Nat. Genet.*, **23**, 147.
- Calvo,S.E. and Mootha,V.K. (2010) The mitochondrial proteome and human disease. *Annu. Rev. Genomics Hum. Genet.*, **11**, 25–44.
- Area-Gomez,E. and Schon,E.A. (2014) Mitochondrial genetics and disease. *J. Child Neurol.*, **29**, 1208–1215.
- Simon,M., Richard,E.M., Wang,X., Shahzad,M., Huang,V.H., Qaiser,T.A., Potluri,P., Mahl,S.E., Davila,A., Nazli,S. *et al.* (2015) Mutations of human NARS2, encoding the mitochondrial asparaginyl-tRNA synthetase, cause nonsyndromic deafness and Leigh syndrome. *PLoS Genet.*, **11**, e1005097
- Pierce,S.B., Gersak,K., Michaelson-Cohen,R., Walsh,T., Lee,M.K., Malach,D., Klevit,R.E., King,M.C. and Levy-Lahad,E. (2013) Mutations in LARS2, encoding mitochondrial leucyl-tRNA synthetase, lead to premature ovarian failure and hearing loss in Perrault syndrome. *Am. J. Hum. Genet.*, **92**, 614–620.
- Prezant,T.R., Agopian,J.V., Bohlman,M.C., Bu,X., Oztas,S., Qiu,W.Q., Arnos,K.S., Cortopassi,G.A., Jaber,L., Rotter,J.I. *et al.* (1993) Mitochondrial ribosomal RNA mutation associated with both antibiotic-induced and non-syndromic deafness. *Nat. Genet.*, **4**, 289–294.
- Zhao,H., Li,R., Wang,Q., Yan,Q., Deng,J.H., Han,D., Bai,Y., Young,W.Y. and Guan,M.X. (2004) Maternally inherited aminoglycoside-induced and non-syndromic deafness is associated with the novel C1494T mutation in the mitochondrial 12S rRNA gene in a large Chinese family. *Am. J. Hum. Genet.*, **74**, 139–152.
- Goto,Y., Nonaka,I. and Horai,S. (1990) A mutation in the tRNA^{Leu(UUR)} gene associated with the MELAS subgroup of mitochondrial encephalomyopathies. *Nature*, **348**, 651–653.
- Mancuso,M., Orsucci,D., Angelini,C., Bertini,E., Carelli,V., Comi,G.P., Minetti,C., Moggio,M., Mongini,T., Servidei,S. *et al.* (2013). Phenotypic heterogeneity of the 8344A>G mtDNA 'MERRF' mutation. *Neurology*, **80**, 2049–2054.
- Gutierrez Cortes,N., Pertuiset,C., Dumon,E., Borlin,M., Hebert-Chatelain,E., Pierron,D., Feldmann,D., Jonard,L., Marlin,S., Letellier,T. *et al.* (2012) Novel mitochondrial DNA mutations responsible for maternally inherited nonsyndromic hearing loss. *Hum. Mutat.*, **33**, 681–689.
- Fischel-Ghodsian,N., Prezant,T.R., Fournier,P., Stewart,I.A. and Maw,M. (1995) Mitochondrial mutation associated with nonsyndromic deafness. *Am. J. Otolaryngol.*, **16**, 403–408.
- Tiranti,V., Chariot,P., Carella,F., Toscano,A., Soliveri,P., Girlanda,P., Carrara,F., Fratta,G.M., Reid,F.M., Mariotti,C. *et al.* (1995) Maternally inherited hearing loss, ataxia and myoclonus associated with a novel point mutation in mitochondrial tRNA^{Ser(UCN)} gene. *Hum. Mol. Genet.*, **4**, 1421–1427.
- Sue,C.M., Tanji,K., Hadjigeorgiou,G., Andreu,A.L., Nishino,I., Krishna,S., Bruno,C., Hirano,M., Shanske,S., Bonilla,E. *et al.* (1999) Maternally inherited hearing loss in a large kindred with a novel T7511C mutation in the mitochondrial DNA tRNA^{Ser(UCN)} gene. *Neurology*, **52**, 1905–1908.
- Tang,X., Zheng,J., Ying,Z., Cai,Z., Gao,Y., He,Z., Yu,H., Yao,J., Yang,Y., Wang,H. *et al.* (2015) Mitochondrial tRNA^{Ser(UCN)} variants in 2651 Han Chinese subjects with hearing loss. *Mitochondrion*, **23**, 17–24.
- Gong,S., Peng,Y., Jiang,P., Wang,M., Fan,M., Wang,X., Zhou,H., Li,H., Yan,Q., Huang,T. *et al.* (2014) A deafness-associated tRNA^{His} mutation alters the mitochondrial function, ROS production and membrane potential. *Nucleic Acids Res.*, **42**, 8039–8048.
- Liu,H., Li,R., Li,W., Wang,M., Ji,J., Zheng,J., Mao,Z., Mo,J.Q., Jiang,P., Lu,J. *et al.* (2015) Maternally inherited diabetes is associated with a homoplasmic T10003C mutation in the mitochondrial tRNA^{Gly} gene. *Mitochondrion*, **21**, 49–57.
- Suzuki,T., Nagao,A. and Suzuki,T. (2011) Human mitochondrial tRNAs: Biogenesis, function, structural aspects, and diseases. *Annu. Rev. Genet.*, **45**, 299–329.
- Zhou,X.L. and Wang,ED. (2013) Transfer RNA: a dancer between charging and mis-charging for protein biosynthesis. *Sci. China Life Sci.*, **56**, 921–932.
- Guan,M.X., Enriquez,J.A., Fischel-Ghodsian,N., Puranam,R.S., Lin,C.P., Maw,M.A. and Attardi,G. (1998) The deafness-associated mitochondrial DNA mutation at position 7445, which affects tRNA^{Ser(UCN)} precursor processing, has long-range effects on NADH dehydrogenase subunit ND6 gene expression. *Mol. Cell. Biol.*, **18**, 5868–5879.
- Li,X., Fischel-Ghodsian,N., Schwartz,F., Yan,Q., Friedman,R.A. and Guan,M.X. (2004) Biochemical characterization of the mitochondrial tRNA^{Ser(UCN)} T7511C mutation associated with nonsyndromic deafness. *Nucleic Acids Res.*, **32**, 867–877.
- Wu,Y., Liang,L.Z., Xiao,H.L., Yang,Y.L., Yu,X., Zheng,J., Fang,F., Zheng,B.J., Tang,X.W., Jin,L.J. *et al.* (2013) Hearing loss may be associated with the novel mitochondrial tRNA^{Asp} A7551G mutation in a Chinese family. *Zhonghua Er Bi Yan Hou Tou Jing Wai Ke Za Zhi*, **48**, 978–984.
- Messmer,M., Pütz,J., Suzuki,T., Suzuki,T., Sauter,C., Sissler,M. and Catherine,F. (2009) Tertiary network in mammalian mitochondrial tRNA^{Asp} revealed by solution probing and phylogeny. *Nucleic Acids Res.*, **37**, 6881–6895.
- Suzuki,T. and Suzuki,T. (2014) A complete landscape of post-transcriptional modifications in mammalian mitochondrial tRNAs. *Nucleic Acids Res.*, **42**, 7346–7357.
- Allnér,O. and Nilsson,L. (2011) Nucleotide modifications and tRNA anticodon-mRNA codon interactions on the ribosome. *RNA*, **17**, 2177–2188.
- Björk,G.R. and Hagervall,T.G. (2014) Transfer RNA modification: presence, synthesis, and function. *EcoSal Plus.*, **6**, doi:10.1128/ecosalplus.ESP-0007-2013.
- Wei,F.Y., Zhou,B., Suzuki,T., Miyata,K., Ujihara,Y., Horiguchi,H., Takahashi,N., Xie,P., Michiue,H., Fujimura,A. *et al.* (2015) Cdk5rap1-mediated 2-methylthio modification of mitochondrial tRNAs governs protein translation and contributes to myopathy in mice and humans. *Cell Metab.*, **21**, 428–442.
- Qu,J., Li,R., Tong,Y., Lu,F., Qian,Y., Hu,Y., Mo,J.Q., West,C.E. and Guan,M.X. (2006) The novel A4435G mutation in the mitochondrial tRNA^{Met} may modulate the phenotypic expression of the LHON-associated ND4 G11778A mutation. *Invest. Ophthalmol. Vis. Sci.*, **47**, 475–483.
- Liu,Y., Li,R., Li,Z., Wang,X.J., Yang,L., Wang,S. and Guan,M.X. (2009) Mitochondrial transfer RNA^{Met} 4435A>G mutation is associated with maternally inherited hypertension in a Chinese pedigree. *Hypertension*, **53**, 1083–1090.

34. King, M.P. and Attardi, G. (1996) Mitochondria-mediated transformation of human rho(0) cells. *Methods Enzymol.*, **264**, 313–334.
35. Jiang, P., Wang, M., Xue, L., Xiao, Y., Yu, J., Wang, H., Yao, J., Liu, H., Peng, Y., Liu, H. *et al.* (2016) A hypertension-associated tRNA^{Ala} mutation alters the tRNA metabolism and mitochondrial function. *Mol. Cell. Biol.*, **36**, 1920–1930.
36. Miller, G. and Lipman, M. (1973) Release of infectious Epstein-Barr virus by transformed marmoset leukocytes. *Proc. Natl. Acad. Sci. U.S.A.*, **70**, 190–194.
37. Guan, M.X., Fischel-Ghodsian, N. and Attardi, G. (2001) Nuclear background determines biochemical phenotype in the deafness-associated mitochondrial 12S rRNA mutation. *Hum. Mol. Genet.*, **10**, 573–580.
38. Guan, M.X., Fischel-Ghodsian, N. and Attardi, G. (1996) Biochemical evidence for nuclear gene involvement in phenotype of non-syndromic deafness associated with mitochondrial 12S rRNA mutation. *Hum. Mol. Genet.*, **5**, 963–971.
39. Ofengand, J., Del Campo, M. and Kaya, Y. (2001) Mapping pseudouridines in RNA molecules. *Methods*, **25**, 365–373.
40. Qian, Y. and Guan, M.X. (2009) Interaction of aminoglycosides with human mitochondrial 12S rRNA carrying the deafness-associated mutation. *Antimicrob. Agents Chemother.*, **53**, 4612–4618.
41. King, M.P. and Attardi, G. (1993) Post-transcriptional regulation of the steady-state levels of mitochondrial tRNAs in HeLa cells. *J. Biol. Chem.*, **268**, 10228–10237.
42. Li, R. and Guan, M.X. (2010) Human mitochondrial leucyl-tRNA synthetase corrects mitochondrial dysfunctions due to the tRNA^{Leu(UUR)} A3243G mutation, associated with mitochondrial encephalomyopathy, lactic acidosis, and stroke-like symptoms and diabetes. *Mol. Cell. Biol.*, **30**, 2147–2154.
43. Jiang, P., Jin, X., Peng, Y., Wang, M., Liu, H., Liu, X., Zhang, Z., Ji, Y., Zhang, J., Liang, M. *et al.* (2016) The exome sequencing identified the mutation in YARS2 encoding the mitochondrial tyrosyl-tRNA synthetase as a nuclear modifier for the phenotypic manifestation of Leber's hereditary optic neuropathy-associated mitochondrial DNA mutation. *Hum. Mol. Genet.*, **25**, 584–596.
44. Enriquez, J.A. and Attardi, G. (1996) Analysis of aminoacylation of human mitochondrial tRNAs. *Methods Enzymol.*, **264**, 183–196.
45. Li, Y., D'Aurelio, M., Deng, J.H., Park, J.S., Manfredi, G., Hu, P., Lu, J. and Bai, Y. (2007) An assembled complex IV maintains the stability and activity of complex I in mammalian mitochondria. *J. Biol. Chem.*, **282**, 17557–17562.
46. Zhang, J., Jiang, P., Jin, X., Liu, X., Zhang, M., Xie, S., Gao, M., Zhang, S., Sun, Y.H., Zhu, J. *et al.* (2014) Leber's hereditary optic neuropathy caused by the homoplasmic ND1 m.3635G>A mutation in nine Han Chinese families. *Mitochondrion*, **18**, 18–26.
47. Dranka, B.P., Benavides, G.A., Diers, A.R., Giordano, S., Zelickson, B.R., Reily, C., Zou, L., Chatham, J.C., Hill, B.G., Zhang, J. *et al.* (2011) Assessing bioenergetic function in response to oxidative stress by metabolic profiling. *Free Radic. Biol. Med.*, **51**, 1621–1635.
48. Zhou, X., Qian, Y., Zhang, J., Tong, Y., Jiang, P., Liang, M., Dai, X., Zhou, H., Zhao, F., Ji, Y. *et al.* (2012) Leber's hereditary optic neuropathy is associated with the T3866C mutation in mitochondrial ND1 gene in three Han Chinese Families. *Invest. Ophthalmol. Vis. Sci.*, **53**, 4586–4594.
49. Mahfouz, R., Sharma, R., Lackner, J., Aziz, N. and Agarwal, A. (2009) Evaluation of chemiluminescence and flow cytometry as tools in assessing production of hydrogen peroxide and superoxide anion in human spermatozoa. *Fertil. Steril.*, **92**, 819–827.
50. Jia, Z., Wang, X., Qin, Y., Xue, L., Jiang, P., Meng, Y., Shi, S., Wang, Y., Qin Mo, J. and Guan, M.X. (2013) Coronary heart disease is associated with a mutation in mitochondrial tRNA. *Hum. Mol. Genet.*, **22**, 4064–4073.
51. Jones, C.N., Jones, C.I., Graham, W.D., Agris, P.F. and Spremulli, L.L. (2008) A disease-causing point mutation in human mitochondrial tRNA^{Met} results in tRNA misfolding leading to defects in translational initiation and elongation. *J. Biol. Chem.*, **283**, 34445–34456.
52. Yarus, M., Cline, S.W., Wier, P., Breeden, L. and Thompson, R.C. (1986) Actions of the anticodon arm in translation on the phenotypes of RNA mutants. *J. Mol. Biol.*, **192**, 235–255.
53. Chomyn, A., Enriquez, J.A., Micol, V., Fernandez-Silva, P. and Attardi, G. (2000) The mitochondrial myopathy, encephalopathy, lactic acidosis, and stroke-like episode syndrome-associated human mitochondrial tRNA^{Leu(UUR)} mutation causes aminoacylation deficiency and concomitant reduced association of mRNA with ribosomes. *J. Biol. Chem.*, **275**, 19198–19209.
54. Enriquez, J.A., Chomyn, A. and Attardi, G. (1995) MtDNA mutation in MERRF syndrome causes defective aminoacylation of tRNALys and premature translation termination. *Nat. Genet.*, **10**, 47–55.
55. Zhao, H., Young, W.Y., Yan, Q., Li, R., Cao, J., Wang, Q., Li, X., Peters, J.L., Han, D. and Guan, M.X. (2005) Functional characterization of the mitochondrial 12S rRNA C1494T mutation associated with aminoglycoside-induced and non-syndromic hearing loss. *Nucleic Acids Res.*, **33**, 1132–1139.
56. James, A.M., Sheard, P.W., Wei, Y.H. and Murphy, M.P. (1999) Decreased ATP synthesis is phenotypically expressed during increased energy demand in fibroblasts containing mitochondrial tRNA mutations. *Eur. J. Biochem.*, **259**, 462–469.
57. Pallotti, F., Baracca, A., Hernandez-Rosa, E., Walker, W.F., Solaini, G., Lenaz, G., Melzi D'Eril, G.V., Dimairo, S., Schon, E.A. and Davidson, M.M. (2004) Biochemical analysis of respiratory function in cybrid cell lines harbouring mitochondrial DNA mutations. *Biochem. J.*, **384**, 287–293.
58. Guan, M.X. (2004) Molecular pathogenetic mechanism of maternally inherited deafness. *Ann. N. Y. Acad. Sci.*, **1011**, 259–271.
59. de Andrade, P.B., Rubi, B., Frigerio, F., van den Ouweland, J.M., Maassen, J.A. and Maechler, P. (2006) Diabetes-associated mitochondrial DNA mutation A3243G impairs cellular metabolic pathways necessary for beta cell function. *Diabetologia*, **49**, 1816–1826.
60. Lenaz, G., Baracca, A., Carelli, V., D'Aurelio, M., Sgarbi, G. and Solaini, G. (2004) Bioenergetics of mitochondrial diseases associated with mtDNA mutations. *Biochim. Biophys. Acta*, **1658**, 89–94.
61. Hayashi, G. and Cortopassi, G. (2015) Oxidative stress in inherited mitochondrial diseases. *Free Radic. Biol. Med.*, **88**, 10–17.
62. Schieber, M. and Chandel, N.S. (2014) ROS function in redox signaling and oxidative stress. *Curr. Biol.*, **24**, R453–R462.
63. Raimundo, N., Song, L., Shutt, T.E., McKay, S.E., Cotney, J., Guan, M.X., Gilliland, T.C., Hohuan, D., Santos-Sacchi, J. and Shadel, G.S. (2012) Mitochondrial stress engages E2F1 apoptotic signaling to cause deafness. *Cell*, **148**, 716–726.
64. Guan, M. X., Yan, Q., Li, X., Bykhovskaya, Y., Gallo-Teran, J., Hajek, P., Umeda, N., Zhao, H., Garrido, G., Mengesha, E. *et al.* (2006) Mutation in TRMU related to transfer RNA modification modulates the phenotypic expression of the deafness-associated mitochondrial 12S ribosomal RNA mutations. *Am. J. Hum. Genet.*, **79**, 291–302.
65. Chen, C., Chen, Y. and Guan, M.X. (2015) A peep into mitochondrial disorder: multifaceted from mitochondrial DNA mutations to nuclear gene modulation. *Protein Cell*, **6**, 862–870.
66. Tischner, C., Hofer, A., Wulff, V., Stepek, J., Dumitru, I., Becker, L., Haack, T., Kremer, L., Datta, A.N., Sperl, W. *et al.* (2015) MTO1 mediates tissue specificity of OXPHOS defects via tRNA modification and translation optimization, which can be bypassed by dietary intervention. *Hum. Mol. Genet.*, **24**, 2247–2266.
67. Chen, D., Li, F., Yang, Q., Tian, M., Zhang, Z., Zhang, Q., Chen, Y. and Guan, M.X. (2016) The defective expression of gtpbp3 related to tRNA modification alters the mitochondrial function and development of zebrafish. *Int. J. Biochem. Cell. Biol.*, **77**, 1–9.
68. Dittmar, K.A., Goodenbour, J.M. and Pan, T. (2006) Tissue-specific differences in human transfer RNA expression. *PLoS Genet.*, **2**, e221.