

Maternally Inherited Essential Hypertension May Be Associated with the Mutations in Mitochondrial tRNA^{Glu} Gene

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Background: Mitochondrial DNA (mtDNA) mutations are associated with essential hypertension (EH), but the molecular mechanism remains largely unknown.

Objective: The aim of this study is to explore the association between mtDNA mutations and EH.

Methods: Two maternally inherited families with EH are underwent clinical, genetic and biochemical assessments. mtDNA mutations are screened by PCR-Sanger sequencing and phylogenetic, and bioinformatics analyses are performed to evaluate the pathogenicity of mtDNA mutations. We also generate cytoplasmic hybrid (cybrid) cell lines to analysis mitochondrial functions.

Results: Matrilinial relatives exhibit variable degree of clinical phenotypes. Molecular analysis reveals the presence of m.A14693G and m.A14696G mutations in two pedigrees. Notably, the m.A14693G mutation occurs at position 54 in the T ψ C loop of tRNA^{Glu}, a position which is critical for post-transcriptionally modification of tRNA^{Glu}. While the m.A14696G mutation creates a novel base-pairing (51C-64G). Bioinformatic analysis shows that these mutations alter tRNA^{Glu} secondary structure. Additionally, patients with tRNA^{Glu} mutations exhibit markedly decreased in mtDNA copy number, mitochondrial membrane potential (MMP) and ATP, whereas the levels of reactive oxygen species (ROS) increase significantly.

Conclusion: The m.A14696G and m.A14693G mutations lead to failure in tRNA^{Glu} metabolism and cause mitochondrial dysfunction that is responsible for EH.

Keywords: EH, mt-tRNA^{Glu} m.A14693G m.A14696G, mitochondrial dysfunction, Chinese families

Introduction

EH was a very common chronic disease which was becoming an urgent public health problem worldwide, it had been estimated that ~9.4 million hypertensive patients died each year.¹ EH was regarded as a risk factor for coronary heart disease (CHD), stroke and renal failure,² thus, understanding its pathophysiology had become a major research focus. Since the landmark discovery of the draft sequences of human genome, experts announced that within 10 years they expected to determine the important of the genome as related to EH.³ Some great advance had been made toward the molecular basis of EH, for example, nuclear genes such as *Nr2f2*,⁴ *CUL3*,⁵ and *EIF2AK4*⁶ had been identified to be associated with EH. However, the detailed molecular mechanism of EH was still unknown.

While the nuclear genome had been studied extensively with respect to EH, we noticed that mtDNA mutations also played active roles in EH. Mitochondrion was a small symbiotic organelle combined with aerobic bacteria and primordial eukaryotic cells.⁷ It had its own DNA, called mtDNA, encoding 37 genes spanning tRNAs, rRNAs and the subunits of the respiratory chain.⁸ As the adaptor that decoded the mRNA sequence into protein, the basic aspects of mt-tRNA structure and function were central to all studies of mitochondrial biomedicine. Unlike canonical tRNAs such as human cytosolic tRNAs, human mt-tRNAs had specific features such as non-classical G-C pairs and mismatches.⁹ In fact, most

mt-tRNAs from all domains of life had a highly conserved cloverleaf structures, consisting of acceptor arm, D-arm, anticodon stem, variable region and T ψ C loop, with an average length of 73 nucleotides. Mutations in mt-tRNAs had been reported in some cases of maternally inherited EH, such as tRNA^{Ile} A4263G,¹⁰ tRNA^{Ser(UCN)} 7471delC,¹¹ and tRNA^{Gln}/tRNA^{Met} A4401G mutations.¹² These mutations may reduce the steady-state level or aminoacylation ability of corresponding mt-tRNAs, affect the 5' or 3' end processing, CCA addition, cause the defects in chemical modification and lead to failure in mt-tRNA metabolism, and subsequently impair mitochondrial translation and function, which was involved in the progression and pathogenesis of EH.¹³

Most recently, with the aim of exploring the molecular basis of maternally transmitted EH, we carried out a genetic screening program for EH-associated mtDNA mutations in Anhui Province, P.R. China. Herein, we reported clinical, genetic and molecular characterizations of two Chinese pedigrees with EH, sequence analysis of the complete mitochondrial genomes led us to identify two potential pathogenic mutations: m.A14693G and m.A14696G in tRNA^{Glu} gene. To further explore the contributions of mt-tRNA^{Glu} mutations to EH, we analyzed mitochondrial functions in cybrids cell lines derived from six patients with tRNA^{Glu} mutations and four controls without these mutations. Subsequently, we noticed that m.A14693G and m.A14696G mutations affected mitochondrial functions, decreased mtDNA copy number, ATP and MMP, and enhanced ROS production. Therefore, mutations in tRNA^{Glu} caused mitochondrial dysfunction that was responsible for hypertension.

Materials and Methods

Subjects

We enrolled two genetically unrelated Chinese families with EH (Figure 1). These pedigrees were ascertained in the Department of Integrated TCM & Western Medicine, Mengcheng County Second People's Hospital in Anhui Province of China. This study was complied with the Declaration of Helsinki, and the methodologies for obtaining the blood samples, as well as the clinical examination of all participants from two pedigrees were approved by the Ethic Committees of Mengcheng County Second People's Hospital. Besides, 268 unrelated healthy controls were obtained from the volunteers in the same area, these subjects were consisted with 160 male and 108 females, aged from 25 to 48 years, with the average of 36 years. The inclusion criteria for these control subjects were as follows: healthy individuals without any diseases or had any family history of cardiovascular and neurological diseases. Exclusion criteria were ongoing maintenance dialysis, a grave acute infectious disease, neoplastic disease, severe liver dysfunction, major surgery, a chronic inflammatory disease and autoimmune disease. Patients who suffered severe life-threatening injury to other organs were also excluded. Both families provided written informed consent, and furthermore, the informed consent to have their case details published were obtained from all subjects enrolled in this study.

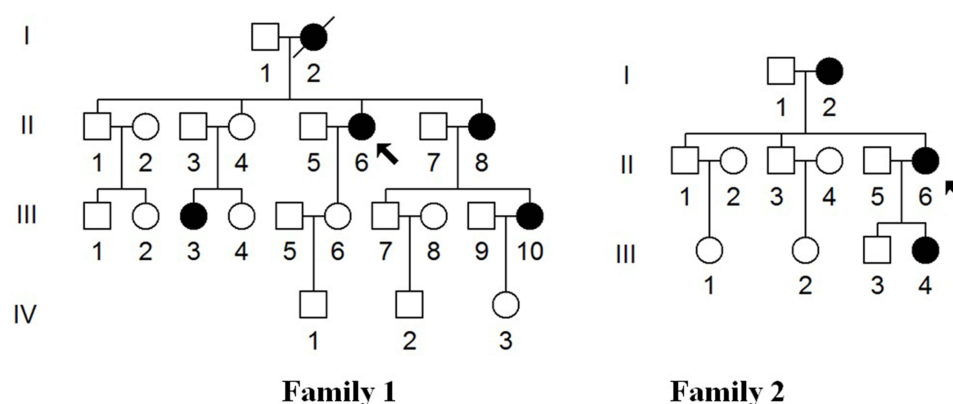


Figure 1 Two Han Chinese families with EH, arrows indicate the probands.

Assessment of BP

The BP of each patient from two families was measured by using a mercury column sphygmomanometer (CARENT Devices, China). The first and the fifth Korotkoff sounds were indicative of systolic and diastolic BP, respectively. The average of three measured BP readings was taken as the examination BP.¹⁴ Notably, EH was defined as systolic BP ≥ 140 mm Hg or diastolic BP ≥ 90 mm Hg on three consecutive days according to the Seventh Report of the Joint National Committee.¹⁵

mtDNA Analysis

To screen the EH-associated mtDNA mutations, we first isolated the genomic DNA from matrilineal relatives from two families, as well as 268 control subjects using the Puregene DNA Isolation Kits (Gentra Systems, Minneapolis, MN), according to the manufacturer's instructions. Briefly, 24 overlapping PCR fragments were generated and amplified the complete mitochondrial genome, according to a previous study.¹⁶ The PCR products were then purified and sequenced by the ABI 3700 automated DNA sequencer. The data were analyzed with SeqWeb program GAP (GCG) according to the revised Cambridge reference sequences (GenBank accession number: NC_012920.1).¹⁷

Analysis of the Conservation Index (CI)

To assess the potential pathogenic roles of mtDNA mutations, the CI of each variant was calculated by using the phylogenetic conservation analysis.¹⁸ The CI $\geq 75\%$ was considered to be functional potential.¹⁹

Bioinformatics Analysis

To see whether m.A14693G and m.A14696G mutations altered the secondary structure of tRNA^{Glu}, the RNA Fold Webserver (<http://rna.tbi.univie.ac.at/cgi-bin/RNAWebSuite/RNAfold.cgi>) was used to predict the minimum-free energy (MFE) of tRNA^{Glu} with and without these mutations.²⁰

Analysis of mtDNA Content

The mtDNA content was measured by using real-time PCR as suggested previously.²¹ Briefly, mtDNA content was normalized to a single copy nuclear β -globin gene. The following primers were used for real-time PCR analysis: for β -globin gene: forward: 5'-CTATgggACgCTTgATgT-3'; reverse: 5'-gCAATCATTcGtCTgTTT-3'. For mtDNA: forward: 5'-CACCAgCCTAACCAG ATTTC-3'; reverse: 5'-gggTTgTATTgATgAgATTAgT-3'. We first generated standard curves for both fragments and calculated their respective amplification efficiencies to test if using the $2^{-\Delta\Delta CT}$ method was appropriate. The real-time PCR was then conducted for the calibrator mtDNA content. All experiments were duplicated in three times.

Generation of Cybrid Cell Lines

Trans-mitochondrial cybrids were obtained by fusion of mtDNA-less ρ^0 human osteosarcoma 143B cells with platelets, which were isolated from the blood of six affected individuals (Family 1: II-6; II-8 and III-10; Family 2: II-1, II-6 and III-4), together with four controls (C1, C2, C3 and C4), as described previously.²² The transformant clones were cultured in DMEM (Sigma-Aldrich) containing 10% FBS (Sigma-Aldrich) at 37°C in a humidified CO₂ incubator.

ATP Analysis

The Cell Titer-Glo[®] Luminescent Cell Viability Assay kit (Promega, Madison, WI, USA) was used for the measurement of ATP levels with some modifications.²³ Luminescence intensity was analyzed by using a fluorescence microplate reader (Molecular Devices, CA, USA), and the amount of ATP was calculated from an ATP standard curve. Each experiment was repeated in three times.

Qualification of ROS Levels

The fluorogenic marker 2', 7'-Dichlorodihydrofluorescein diacetate (H₂DCFDA) was live-cell-permeable acetate ester, and upon entry, it was then cleaved by cellular esterases, reacted with cellular ROS and emitted green fluorescence. To

analyze the ROS level, a total of 2×10^6 cells were first incubated with H₂DCFDA for 30 min, after which the cells were analyzed using a fluorescence plate reader, as described previously.²⁴ Each experiment was repeated in three times.

MMP Analysis

To determine whether mt-tRNA^{Glu} mutations affected mitochondrial function, the MMP of mutant and control cell lines was performed using JC-1 Assay Kit-Microplate (Abcam). JC-1 was a cationic carbocyanine dye; when it accumulated in the mitochondria of low MMP in monomer form, it produced green fluorescence. At high concentrations in high MMP, it produced red fluorescence.²⁵ Each experiment was repeated in three times.

Evaluation of the Pathogenicity

We further utilized the pathogenicity scoring system to evaluate the potential pathogenic roles of m.A14693G and m.A14696G mutations.²⁶ This pathogenicity scoring system employed a number of weighted criteria covering a range of molecular and genetic data, from which an overall pathogenicity score can be obtained. In particular, a variant was classified as “definitely pathogenic” with a score >11 points, whereas variant was defined as “possible pathogenic” with a core of 7–10 points and a “neutral polymorphism” with a score of <6 points.

Statistical Analyses

Statistical analyses were performed using SPSS 22.0 (SPSS Inc., Chicago, IL, USA). Student's *t*-test was used to assess the statistical significance between unpaired samples. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

Clinical and Genetic Characterizations of Two Chinese Families with EH

In Family 1, the proband (II-6) was a 75-year-old woman who came from Mengcheng City of Anhui Province. She developed EH at the age of 70, her average BP was 150/100 mmHg. She went to our hospital for regular treatment of hypertension. After comprehensive history examination, we noticed that several members (II-8, III-3 and III-10) were also hypertensive persons. Therefore, the transmission of EH was a typical maternally inheritance. In addition, all members in Family 1 did not suffer from vision or hearing loss, cancer, diabetes, neurological disorders, suggested that they manifested hypertension as a sole clinical phenotype.

In Family 2, the proband (II-6) was a 66-year-old woman who lived in Mengcheng City of Anhui Province. After the genetic counseling, we found that she suffered from EH when she was 60, and her BP was 180/100 mmHg. In addition, her mother (I-2) and daughter (III-4) were hypertensive individuals. Moreover, all members in Family 2 did not exhibit other mitochondrial disorders such as hearing impairment, vision loss, neurological diseases, cancer and other endocrine diseases, indicating that they expressed the EH as sole clinical phenotypes. The clinical characterizations of these members from two families were listed in [Table 1](#).

Screening for mtDNA Mutations

As these pedigrees were maternally transmitted, which suggested the involvement of mtDNA mutations in the phenotypic manifestation of EH. For this purpose, we amplified the whole mitochondrial genomes from the matrilineal relatives of Family 1 (II-6, II-8, III-3 and III-10) and Family 2 (I-2, II-6 and III-4). After PCR amplification, the products were sequenced by automatic DNA sequencer and subsequently analyzed by using DNA Star software to detect mtDNA variants. The data were listed in [Table 2](#), sequence analysis revealed a set of genetic polymorphisms, in addition to the tRNA^{Glu} A14693G and A14696G mutations ([Figures 2 and 3](#)). Note that, there were 14 variants in D-loop region, four known mutations in 12S rRNA and three mutations in 16S rRNA, whereas other genetic polymorphisms were mainly localized at oxidative phosphorylation (OXPHOS)-related genes. Moreover, 12 missense mutations were identified, including the *ND2* G4924A (Ser to Asn) and C5178A (Leu to Met), *A8* C8414T (Leu to Phe), *A6* A8701G (Thr to Ala) and A8860G (Thr to Ala), *CO3* A9327G (Thr to Ala), *ND3*

Table 1 Summary of Clinical Data for Some Members in Two Families with EH

Subjects	Gender	Age at Test	Age at Onset	Diastolic BP (mmHg)	Systolic BP (mmHg)
II-6 (Family 1)	Female	75	70	100	150
II-8 (Family 1)	Female	72	69	90	145
III-3 (Family 1)	Female	50	48	95	130
III-10 (Family 1)	Female	45	41	90	160
I-2 (Family 2)	Female	88	71	95	145
II-6 (Family 2)	Female	66	60	100	180
III-4 (Family 2)	Female	40	35	90	155

Abbreviation: BP, blood pressure.

Table 2 mtDNA Sequence Variations in Two Families with EH

Gene	Position	Replacement	Conservation (H/B/M/X) ^a	rCRS ^b	Family 1	Family 2	Previously Reported ^c
D-loop	73	A to G		A	G	G	Yes
	146	T to C		T	C		Yes
	150	C to T		C		T	Yes
	195	T to C		T	C	C	Yes
	263	A to G		A	G	G	Yes
	310	T to C		T	C		Yes
	489	T to C		T	C	C	Yes
	524	Del C		C		Del C	Yes
	573	C to CCC		C	CCC		Yes
	16,189	T to C		T	C	C	Yes
	16,223	C to T		C	T	T	Yes
	16,304	T to C		T	C		Yes
	16,362	T to C		T		C	Yes
16,519	T to C		T	C		Yes	
12S rRNA	709	G to A	G/G/A/-	G		A	Yes
	750	A to G	A/G/G/-	A	G	G	Yes
	1107	T to C	T/C/T/T	T		C	Yes
	1438	A to G	A/A/A/G	A	G	G	Yes
16S rRNA	2706	A to G	A/G/A/A	A	G	G	Yes
	3010	G to A	G/G/A/A	G		A	Yes

(Continued)

Table 2 (Continued).

Gene	Position	Replacement	Conservation (H/B/M/X) ^a	rCRS ^b	Family 1	Family 2	Previously Reported ^c
	3107	Del N		N	Del N		Yes
ND1	3348	A to G		A	G		Yes
	3970	C to T		C	T	T	Yes
ND2	4685	A to G		A	G		Yes
	4769	A to G		A		G	Yes
	4883	C to T		C		T	Yes
	4924	G to A (Ser to Asn)	S/N/T/P	G	A		Yes
	4985	G to A		G		A	Yes
	5178	C to A (Leu to Met)	L/T/T/T	C		A	Yes
	5301	A to G		A	G		Yes
CO1	6026	G to A		G		A	Yes
	6284	A to G		A	G		Yes
	7028	C to T		C	T	T	Yes
	7196	C to A		C	A		Yes
CO2	7768	A to G		A	G	G	Yes
	8020	G to A		G	A		Yes
A8	8414	C to T (Leu to Phe)	L/F/M/W	C		T	Yes
A6	8701	A to G (Thr to Ala)	T/S/L/Q	A	G	G	Yes
	8860	A to G (Thr to Ala)	T/A/A/T	A	G	G	Yes
	8964	C to T		C		T	Yes
CO3	9327	A to G (Thr to Ala)	T/T/T/I	A	G		Yes
	9540	T to C		T	C	C	Yes
	9950	T to C		T		C	Yes
ND3	10,398	A to G (Thr to Ala)	T/T/T/A	A	G		Yes
	10,400	C to T		C	T		Yes
ND4	10,873	T to C		T	C	C	Yes
	11,719	G to A		G	A	A	Yes
ND5	12,705	C to T		C	T		Yes
	13,708	G to A (Ala to Thr)	A/L/A/A	G		A	Yes
	13,928	G to C (Ser to Thr)	S/T/S/T	G	C		Yes
ND6	14,455	C to T		C		T	Yes
tRNA^{Glu}	14,693	A to G	A/A/A/A	A	G		Yes

(Continued)

Table 2 (Continued).

Gene	Position	Replacement	Conservation (H/B/M/X) ^a	rCRS ^b	Family 1	Family 2	Previously Reported ^c
	14,696	A to G	A/A/A/A	A		G	Yes
CytB	14766	C to T (Thr to Ile)	T/S/T/S	C	T	T	Yes
	15,301	G to A		G		A	Yes
	15,326	A to G (Thr to Ala)	T/M/I/I	A	G	G	Yes
	15,508	C to T		C		T	Yes
	15,851	A to G (Ile to Val)	I/A/S/M	A		G	Yes

Notes: ^aConservation of amino acid for polypeptide or nucleotide for rRNAs in human (H), bovine (B), mouse (M), and *Xenopus laevis* (X). ^brCRS: revised Cambridge Reference Sequences. ^cSee the online mitochondrial genome database <http://www.mitomap.org>.

A10398G (Thr to Ala), *ND5* G13708A (Ala to Thr) and G13928C (Ser to Thr), *CytB* A15326G (Thr to Ala) and A15851G (Ile to Val).

We used the following criteria to classify mtDNA pathogenic mutations: 1) occurred in <1% of the controls; 2) evolutionary conservation (CI \geq 75%); 3) potential structural and functional alterations; 4) pathogenicity scoring system. Phylogenetic conservation analysis including the mtDNA sequences from mouse,²⁷ bovine²⁸ and *Xenopus laevis*²⁹ was also performed. We found that besides the m.A14693G and m.A14696G mutations (Figure 4), others were not well conserved from different species. Moreover, further genetic analysis revealed that the m.A14693G and m.A14696G mutations were not detected in 268 control subjects ($P < 0.05$ for all), suggesting that the m.A14693G and m.A14696G mutations may be involved in the pathogenesis of EH in these families.

In fact, the m.A14693G mutation occurred at position 54 in T Ψ C loop of tRNA^{Glu}, which was extremely conserved from bacteria to human mitochondria.³⁰ In addition, the m.A14696G mutation was localized at position 51 in T Ψ C loop of tRNA^{Glu}, creating a novel base pairing (51C-64G).³¹

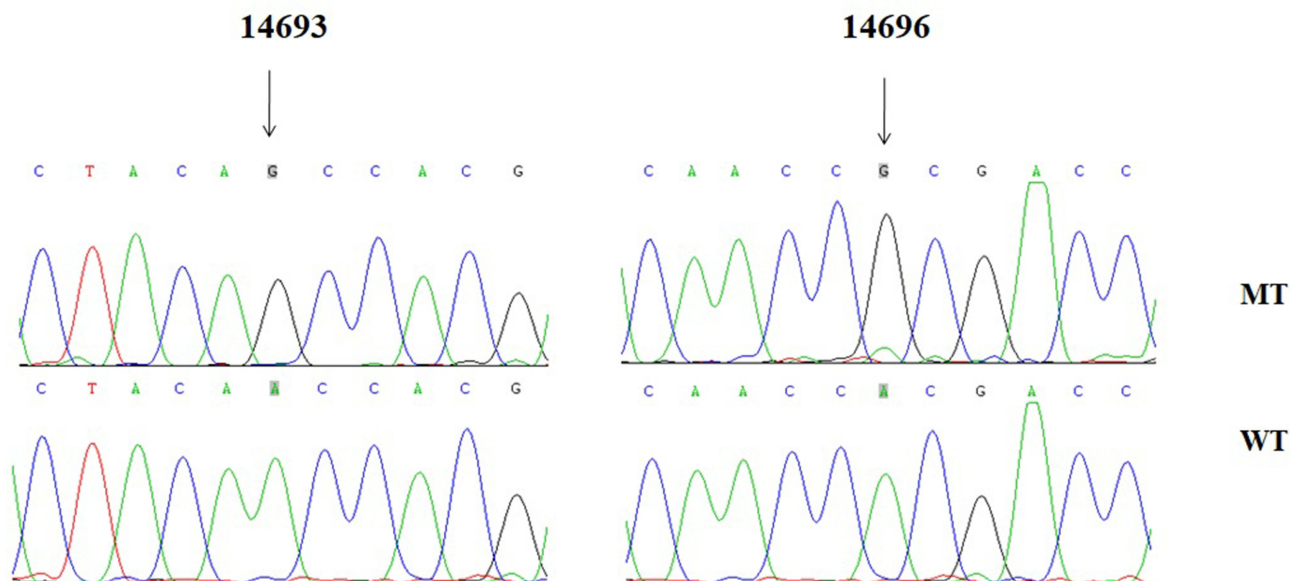


Figure 2 Identification of tRNA^{Glu} A14693G and A14696G mutations by direct sequencing.

Abbreviations: MT, mutant; WT, wild type.

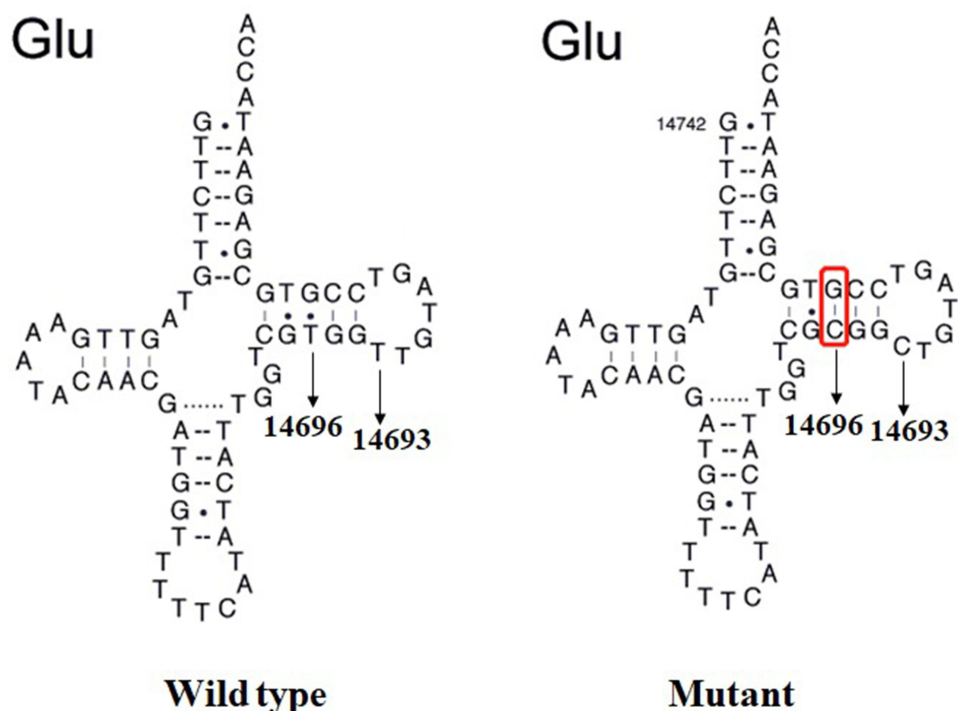


Figure 3 Secondary structure of tRNA^{Glu} gene with and without the m.A14693G and m.A14696G mutations.

14696 14693
↓ ↓

Organism	Acc-stem	D-stem	D-loop	D-stem	Ac-stem	Anticd-loop	Ac-stem	V-region	T-stem	T-loop	T-stem	Acc-stem			
	1	8	10	22	26	27	32	39	44	49 51	54	61	66	73	
<i>Cavia porcellus</i>	GTTTCTG	TA	GTTG	AATTA	CAAC	A	GTGGT	TTTTTCAT	AOCAC	TAGT	CATGG	TTAAACT	CCATG	TAGGAAT	T
<i>Macaca mulatta</i>	GCTTTTA	TA	GTTG	AAATA	CAAC	G	ATGGT	TTTTTCGT	ACCAT	TAGT	CATGG	TTGGAGT	CCATG	TGAAAGC	A
<i>Gorilla gorilla</i>	GTTCTTG	TA	GTTG	AAGTA	CAAC	G	ATGGT	TTTTTCAT	ATCAT	TAGT	CGCGG	TCGTGGT	CCGTG	CGAGAAT	G
<i>Homo sapiens</i>	GTTCTTG	TA	GTTG	AAATA	CAAC	G	ATGGT	TTTTTCAT	ATCAT	TGGT	CGTGG	TTGTAGT	CCGTG	CGAGAAT	A
<i>Panthera tigris</i>	GTTCTTA	TA	GTTG	AAATA	CAAC	G	ATGGT	TTTTTCAT	ATCAT	TGGT	CGTGG	TTAGAAT	CCATG	TGAGAAT	T
<i>Pongo pygmaeus</i>	GTTCTTG	TA	GTTG	AGATA	CAAC	G	ATGTT	TTTTTCAT	ATCAT	TAGT	CACAG	TTACAGT	CTATG	CGAGAAC	A
<i>Tarsius bancanus</i>	GTTCTTA	TA	GTTG	AATAA	CAAC	G	GTGAT	TTTTTCAG	GTCAT	TAAT	CATGG	TTTAAAT	CCATG	TAGGAAC	T
<i>Lemur catta</i>	ATTTTTA	TA	GTTG	AAGTA	CAAC	G	ATGGT	TTTTTCAT	ATCAT	TAGT	CATGG	TTTAAAT	CCATG	TAAGAAT	T
<i>Macaca mulatta</i>	GCTTTTA	TA	GTTG	AAATA	CAAC	G	ATGGT	TTTTTCGT	ACCAT	TAGT	CATGG	TTGGAGT	CCATG	TGAAAGC	A
<i>Crocidura russula</i>	GTTTCTG	TA	GTTG	AAGAA	CAAC	G	ATGAT	TTTTTCAT	GTCAT	AGGT	CGTGG	TTATAC	CCATG	CGGAAAT	T
<i>Mogera wogura</i>	ATTCCTG	TA	GTTG	AATTA	CAAC	G	ATGGT	TTTTTCAT	GTCAT	TGGT	CATGG	TTAAAT	CCATG	TGAGAAT	A
<i>Talpa europaea</i>	ATTCCTG	TA	GTTG	AATAA	CAAC	G	ATGAT	TTTTTCAT	GTCAT	TGGT	CATGG	TTAAAT	CCATG	TGGGAAT	A
<i>Bos taurus</i>	GTTCTTG	TA	GTTG	AATGA	CAAC	G	ATGGT	TTTTTCAT	ATCAT	TAGT	CATGG	TTAGATT	CCATG	TAAGAAT	A
<i>Uncia uncia</i>	GTTCTTA	TA	GTTG	AAATA	CAAC	A	ATGGT	TTTTTCAT	ATCAT	TAGT	CATGG	TTAGATT	CCATG	TGAGAAT	T

Figure 4 Alignment of tRNA^{Glu} gene from various species, arrows indicated the positions of 51 and 54, corresponding to the m.A14696G and m.A14693G mutations.

The m.A14693G and m.A14696G Mutations Altered tRNA^{Glu} Structure

The secondary structures of tRNA^{Glu} with and without A14696G and A14693G mutations were predicted by RNA Fold Webserver.²⁰ As shown in Figure 5, we noticed that m.A14696G and m.A14693G mutations significantly altered tRNA^{Glu} structure, highlighting the impact of these mutations on tRNA functions.³²

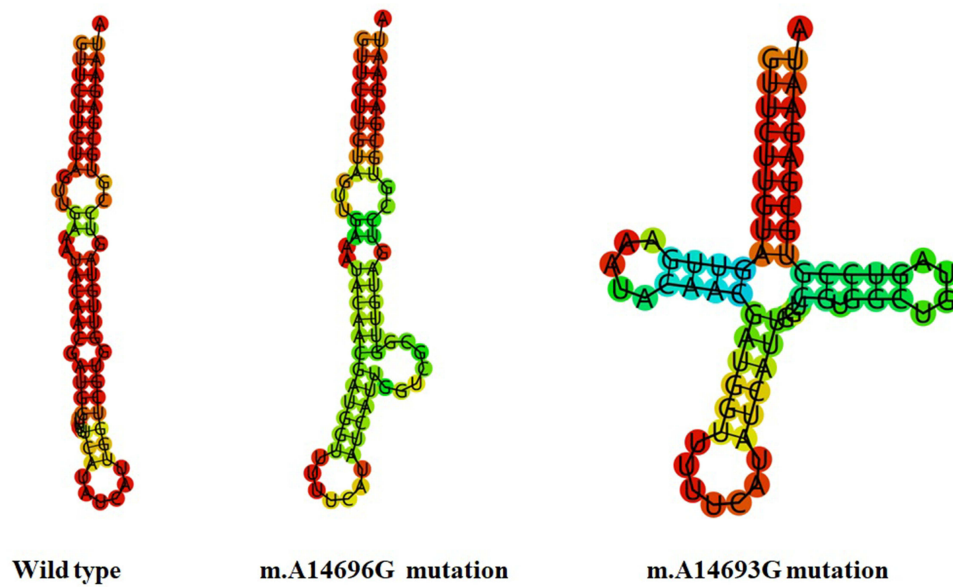


Figure 5 Prediction of tRNA^{Glu} secondary structure with and without the m.A14696G and m.A14693G mutations.

The m.A14693G and m.A14696G Mutations Decreased mtDNA Copy Number

As shown in [Figure 6A](#), patients with m.A14696G and m.A14693G mutations exhibited much lower levels of mtDNA copy number when comparing with the controls ($P=0.0030$).

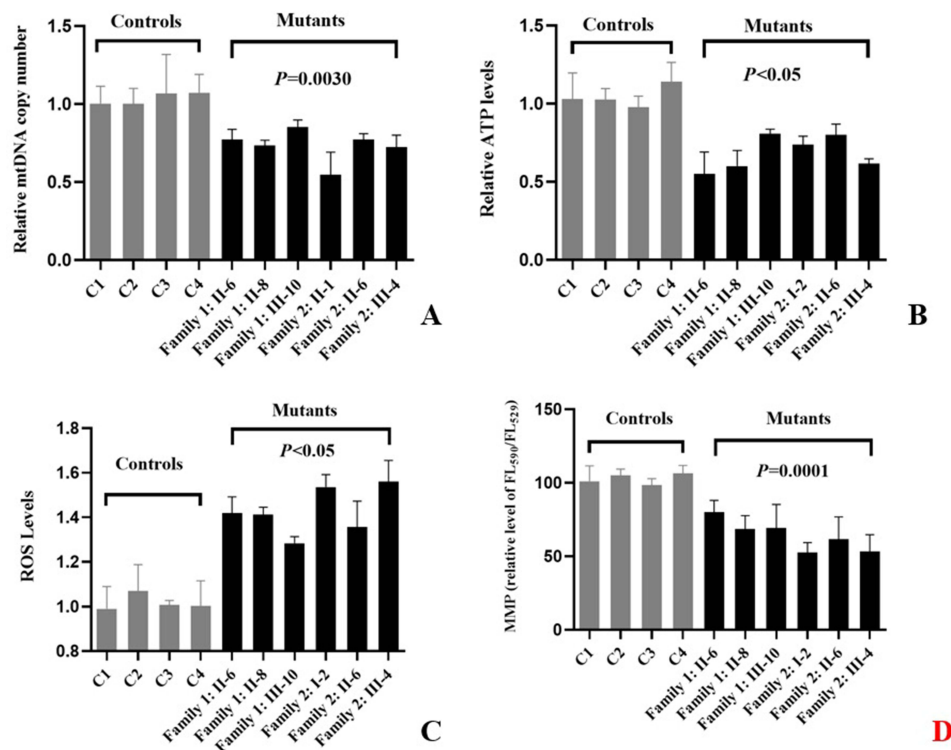


Figure 6 Analysis of mitochondrial functions in cybrids. (A) mtDNA copy number; (B) ATP analysis; (C) ROS analysis; (D) MMP analysis.

ATP Decreased in Cells Carrying tRNA^{Glu} Mutations

Since mitochondria generated ATP through OXPHOS, impairment in ATP synthesis was thought to be involved in mitochondrial function.³³ As shown in Figure 6B, approximately 33% reduction of ATP was observed in patients carrying the mt-tRNA^{Glu} mutations when compared with the healthy subjects ($P<0.05$).

Enhancement of ROS Production

As shown in Figure 6C, patients with m.A14693G or m.A14696G mutations showed an approximately 37% increased in ROS production as compared with controls without these mutations ($P<0.05$).

Decreased in MMP

Decreased in MMP was an early event for apoptosis and critical for mitochondrial function.³⁴ As shown in Figure 6D, subjects with mt-tRNA^{Glu} mutation showed an approximately 31% reduction in MMP as compared with controls without these mutations ($P=0.0001$).

The m.A14693G and m.A14696G Mutations May Be Risk Factors for EH

According to the revised pathogenicity scoring system by Yarham et al,²⁶ the total scores of m.A14693G and m.A14696G mutations were 11 and 9 points, respectively (Table 3), suggesting that they belonged to “definitely pathogenic” and “possibly pathogenic” at this stage.

Discussion

In the current study, we carried out clinical and genetic assessments of two Chinese families with EH, and further investigated the contributions of mtDNA mutations to EH. Notably, members of these pedigrees expressed variable degrees of BP and different age at onset of EH. Interestingly, the age of onset of EH in Family 1 ranged from 41 to 70 years, with the average of 57 years. Meanwhile, the age of onset of EH in Family 2 ranged from 35 to 71 years, with the average of 55 years. In addition, compared with the first and second generations of this pedigree, members in the third generation had an earlier age of onset of EH, indicating that screening for the mtDNA pathogenic mutations was necessary for early diagnosis and prevention of EH.

Table 3 The Pathogenicity Scoring System for m.A14693G and m.A14696G Mutations

Scoring Criteria	m.A14693G Mutation	Score	m.A14696G Mutation	Score	Classification
More than one independent report	Yes	2	Yes	2	≤6 points: neutral polymorphisms; 7–10 points: possibly pathogenic; ≥11 points (including trans-mitochondrial cybrid studies): definitely pathogenic.
Evolutionary conservation of the base pair	No changes	2	No changes	2	
Variant heteroplasmy	No	0	Yes	2	
Segregation of the mutation with disease	Yes	2	Yes	2	
Histochemical evidence of mitochondrial disease	No evidence	0	No evidence	0	
Biochemical defect in complex I, III or IV	No	0	No	0	
Evidence of mutation segregation with biochemical defect from single-fiber studies	No	0	No	0	
Mutant mt-tRNA steady-state level or evidence of pathogenicity in trans-mitochondrial cybrid studies	Strong evidence	5	Weak evidence	3	
Maximum score	Definitely pathogenic	11	Possibly pathogenic	9	

Analysis of the entire mitochondrial genomes from the matrilineal relatives revealed the presence of homoplasmic mt-tRNA^{Glu} A14693G and A14696G mutations, as well as sets of polymorphisms belonging to human mitochondrial haplogroup Y2 and D4a,³⁵ respectively. Notably, nucleotide at position 54 was more prone to be modified than other positions of tRNA.³⁶ The nucleotide modification at this position had been shown to have a pivotal role in the stabilization of tertiary structure and the biochemical function of tRNA^{Glu}.³⁷ Therefore, the m.A14693G mutation caused an impairment of tRNA modification and mitochondrial translation.³⁸ Moreover, the m.A14693G mutation had been reported to be related to Leber's hereditary optic neuropathy (LHON) and acted as a modifier for increasing the penetrance and expressivity of m.A1555G-induced deafness.^{39,40}

While the homoplasmic m.A14696G mutation occurred at position 51 in the T ψ C loop of tRNA^{Glu}, which was also very conserved from various species.⁴¹ Interestingly, the m.A14696G mutation created a novel base pairing (51C-64G) and may result a failure in tRNA metabolism. Importantly, the stem of T ψ C loop harbored a wobble composed of nucleotides 50 to 51 and 63 to 64 in human mt-tRNA^{Glu},⁴² therefore, the mutant 14,696 reduced the wobble and may lead to a failure in tRNA metabolism. Furthermore, bioinformatic analysis revealed that the m.A14693G and m.A14696G mutations caused obviously change of tRNA^{Glu} secondary structure, indicating that the alternation of tRNA structure may affect its steady-state level, as well as its aminoacylation ability, as in the case of tRNA^{Leu(UUR)} A3243G and ND6 T14502C mutations.⁴³

To see the contributions of m.A14693G and m.A14696G mutations to EH, we analyzed mitochondrial functions including mtDNA content, ATP, MMP and ROS in hypertension patients and healthy controls. As a result, markedly decreased in mtDNA copy number, ATP and MMP were observed in patients carrying tRNA^{Glu} mutations, whereas ROS increased significantly. In fact, mtDNA copy number was a mitochondrial function marker that reflected its depletion, energy reserves and oxidative stress.⁴⁴ Recent experimental studies indicated that decreased peripheral mtDNA copy number was associated with the risk of heart failure and long-term outcomes.⁴⁵ MMP reflected the pumping of hydrogen ions across the inner membrane during the process of electron transport and OXPHOS.⁴⁶ The defects in MMP may be due to strongly decreased efficiency of respiratory chain-mediated proton extrusion for the matrix, as in the case of tRNA^{His} T12201C mutation.⁴⁷

Additionally, a decreased mtDNA copy number had been demonstrated to lead to increased ROS levels; ROS induced by mitochondrial dysfunction can increase mitochondrial Ca²⁺ accumulation and may act as potential pathophysiological mechanism in hypertension.⁴⁸ Furthermore, the ATP dropped significantly indicated that the OXPHOS complexes were impaired in subjects with m.A14693G and m.A14696G mutations. Through the application of the pathogenicity scoring system,²⁶ the total scores of m.A14693G and m.A14696G mutations were 11 and 9 points, respectively, belonged to "definitely" and "possibly" pathogenic at this stage (Table 3).

However, the homoplasmic form, late onset and incomplete penetrance of EH observed in these Chinese families carrying the tRNA^{Glu} mutations suggested that the m.A14693G and m.A14696G mutations were involved in the development of EH but may be insufficient to produce a clinical phenotype; hence, other factors such as nuclear genes (*ADD1*; *ALDH1A3*),^{49,50} environmental components (air pollution), epigenetic modification (histone modification, DNA methylation)^{51,52} and personal lifestyle (smoking or high salt intake) may contribute to EH expression in these two families.^{53,54} In particular, mito-nuclear communication played a putative role in the pathogenesis of cardiovascular disease.⁵⁵ The main limitations of this study were the relatively small sample size, further studies including more EH patients and controls, as well as the examinations of tRNA functions in the cybrids were needed to verify this conclusion.

Conclusion

Our study indicated that mt-tRNA^{Glu} A14693G and A14696G mutations altered the tRNA structure and functions, led to mitochondrial dysfunction that was involved in EH, screening for tRNA mutations was recommended for early diagnosis and detection of EH.

Abbreviations

EH, essential hypertension; mtDNA, mitochondrial DNA; cybrid, cytoplasmic hybrid; MMP, mitochondrial membrane potential; ROS, reactive oxygen species; mt-tRNA, mitochondrial tRNA; CHD, coronary heart disease; BP, blood

pressure; CI, conservation index; MFE, minimum-free energy; H₂DCFDA, 2', 7'-Dichlorodihydrofluorescein diacetate; OXPHOS, oxidative phosphorylation; LHON, Leber's hereditary optic neuropathy.

Data Sharing Statement

The datasets used and analyzed during the current study are available from corresponding author (Chun Wang, E-mail: chunwang0424@163.com) on reasonable request.

Ethics Approval and Consent to Participate

This study was approved by the Ethics Committee of Mengcheng County Second People's Hospital and conformed to the ethical principles of the Declaration of Helsinki. All participants from two pedigrees signed informed consent forms, as well as their cases to be published were obtained before participating in this study.

Acknowledgments

We are grateful to the members of two pedigrees for participating in this study.

Author Contributions

All authors contributed to data analysis, drafting or revising the article, have agreed on the journal to which the article will be submitted, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Funding

There is no funding to report.

Disclosure

No potential conflict of interest was reported by the author(s).

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