Nuclear Modifier \textit{MTO2} Modulates the Aminoglycoside-Sensitivity of Mitochondrial 15S rRNA C1477G Mutation in \textit{Saccharomyces cerevisiae}

Xiangyu He, Xiaoyu Zhu, Xuexiang Wang, Wei Wang, Yu Dai, Qingfeng Yan*

College of Life Science, Zhejiang University, Hangzhou, Zhejiang, China

Abstract

The phenotypic manifestations of mitochondrial DNA (mtDNA) mutations are modulated by mitochondrial DNA haplotypes, nuclear modifier genes and environmental factors [1,2]. A typical example is the human mtDNA 12S rRNA A1555G mutation, which is well known as a primary determinant of aminoglycoside-induced nonsyndromic deafness [3,4,5]. However, individuals carrying the A1555G mutation exhibit diverse clinical phenotypes ranging from normal hearing to severe deafness. This suggests that the clinical symptom may be also under the regulation of nuclear genes and environmental factors [3,5,6,7]. So far, most studies on the mitochondrial A1555G mutation have had their main focus upon one factor, or the interaction between two factors. The combined effect of mtDNA mutation and nuclear modifier genes have been presented in several studies where environmental influences appear to play a role but remain poorly understood [7,8].

The yeast mitochondrial 15S rRNA C1477G mutation corresponds to the human 12S rRNA C1494T and A1555G mutations (Figure 1). Yeast carrying this mutation is often used as a model system to investigate aminoglycoside-sensitivity. Here we report that the deletion of the nuclear modifier gene \textit{MTO2} suppressed the aminoglycoside-sensitivity of mitochondrial 15S rRNA C1477G mutation in \textit{Saccharomyces cerevisiae}. First, the strain with a single mtDNA C1477G mutation exhibited hypersensitivity to neomycin. Functional assays indicated that the steady-state transcription level of mitochondrial DNA, the mitochondrial respiratory rate, and the membrane potential decreased significantly after neomycin treatment. The impaired mitochondria could not produce sufficient energy to maintain cell viability. Second, when the \textit{mto2} null and the mitochondrial C1477G mutations co-existed (\textit{mto2}(\textit{PS})), the oxygen consumption rate in the double mutant decreased markedly compared to that of the control strains (\textit{MTO2}(\textit{PS})), \textit{mto2}(\textit{PS}) and \textit{MTO2}(\textit{PS})). The expression levels of the key glycolytic genes \textit{HXK2}, \textit{PFK1}, and \textit{PYK1} in the \textit{mto2}(\textit{PS}) strain were stimulated by neomycin and up-regulated by 89%, 112% and 55%, respectively. The enhanced glycolysis compensated for the respiratory energy deficits, and could be inhibited by the glycolytic enzyme inhibitor. Our findings in yeast will provide a new insight into the pathogenesis of human deafness.

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* E-mail: qfyan@zju.edu.cn

Introduction

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The yeast mitochondrial 15S rRNA C1477G mutation corresponds to the human 12S rRNA C1494T and A1555G mutations (Figure 1). Yeast carrying this mutation is often used as a model system to investigate aminoglycoside-sensitivity. Here we report that the deletion of the nuclear modifier gene \textit{MTO2} suppressed the aminoglycoside-sensitivity of mitochondrial 15S rRNA C1477G mutation in \textit{Saccharomyces cerevisiae}. First, the strain with a single mtDNA C1477G mutation exhibited hypersensitivity to neomycin. Functional assays indicated that the steady-state transcription level of mitochondrial DNA, the mitochondrial respiratory rate, and the membrane potential decreased significantly after neomycin treatment. The impaired mitochondria could not produce sufficient energy to maintain cell viability. Second, when the \textit{mto2} null and the mitochondrial C1477G mutations co-existed (\textit{mto2}(\textit{PS})), the oxygen consumption rate in the double mutant decreased markedly compared to that of the control strains (\textit{MTO2}(\textit{PS})), \textit{mto2}(\textit{PS}) and \textit{MTO2}(\textit{PS})). The expression levels of the key glycolytic genes \textit{HXK2}, \textit{PFK1}, and \textit{PYK1} in the \textit{mto2}(\textit{PS}) strain were stimulated by neomycin and up-regulated by 89%, 112% and 55%, respectively. The enhanced glycolysis compensated for the respiratory energy deficits, and could be inhibited by the glycolytic enzyme inhibitor. Our findings in yeast will provide a new insight into the pathogenesis of human deafness.

In \textit{Saccharomyces cerevisiae}, the mitochondrial 15S rRNA C1477G mutation combining with the nuclear gene \textit{mss1}, \textit{mto1} or \textit{mto2} null mutations generated a respiratory deficient phenotype. In yeast mitochondria, Mto2p, along with Mto1p and Mss1p, participates in the same pathway catalyzing the formation of the hypermodi-
Figure 1. Secondary structure of small rRNA decoding sites in yeast and human mitochondria. A, secondary structure of *E. coli* small ribosome rRNA decoding site. B, wild type and \(P^\beta\) mutant forms of yeast 15S rRNA decoding sites, and the base-pair affected by \(P^\beta\) mutation are

### E. coli 16S rRNA

A

|   | C−G | G−C |
|---|-----|-----|
| U | U   |     |
| C−G | A   |
| A | A   |

1409 \(\rightarrow\) C−G \(\leftarrow\) 1491

A−U
C−G
C−G
A−G
U−G
G−U
G−C
G−A

**y**

### yeast 15S rRNA wild-type

B

|   | C−G | C−G |
|---|-----|-----|
| A−U | A−U |
| U | U   |     |
| C−G | C−G |
| A | A   |

A

1477 \(\rightarrow\) C−G−1583

G−C
C−G
G−U
U−A
U−A
G−U
A−U
A−A
A−A

**y**

### yeast 15S rRNA \(P^\beta\) 454 mutant

B

|   | C−G | C−G |
|---|-----|-----|
| A−U | A−U |
| U | U   |     |
| C−G | C−G |
| A | A   |

A

C

1477 \(\rightarrow\) G \(\leftarrow\) 1583

G−C
G−C
G−U
U−A
U−A
G−U
A−U
A−A
A−A

**y**

### human 12S rRNA wild-type

C

|   | C−G |
|---|-----|
| G−C | G−C |
| U | U   |
| C−G | C−G |
| A | A   |

A

C

1494 \(\rightarrow\) C−A−1555

C−G
C−G
C−G
C−G
C−G
C−G
C−G
A−A
A−A
A−A

**A**

### human 12S rRNA \(A^{1555}\) and \(C^{1494}\) mutants

C

|   | A−A |
|---|-----|
| A−A | A−A |

\(C^{1494}\) mutant

|   | A−A |
|---|-----|
| A−A | A−A |

\(A^{1555}\) mutant

|   | A−A |
|---|-----|
| A−A | A−A |
fied base 5-methyl-aminomethyl-2-thio-uridine (mm\textsuperscript{5}s\textsuperscript{2}U34) in the wobble position of tRNA\textsuperscript{Lys}, tRNA\textsuperscript{Glu} and tRNA\textsuperscript{Gln} [16,17,18]. MTO2 encodes a mitochondrial rRNA-specific 2-thioribidylase, which is responsible for 2-thiolation of the U34 nucleotide in the rRNA anti-codon loop. This is the initial step of the mnm\textsuperscript{5}s2U34 modification pathway [17]. This kind of modification is crucial to the mitochondrial translational fidelity and the efficiency of protein synthesis, because the modified uridine base increases both the stability of these tRNAs and the capacity of codon recognition at the ribosomal A-site. Meanwhile, TRMU (human MTO2) was the first identified nuclear modifier gene regulating the phenotypic manifestation of the mitochondrial A1555G mutation [19]. These studies indicated a complex interaction between the MTO2 (TRMU) gene and the mitochondrial small rRNA.

The aim of the present study is to determine the interactions among mitochondrial rRNA mutations, nuclear genes, and environmental factors, as well as their combined effects on cellular function. For the first time, we found that the nuclear gene MTO2 could modulate aminoglycoside antibiotic sensitivity induced by the mitochondrial C1477G mutation. Yeast carrying the 15S rRNA C1477G mutation alone exhibited hypersensitivity to aminoglycosides, while MTO2 deletion suppressed aminoglycoside-sensitivity in yeast carrying the mitochondrial C1477G mutation. The underlying molecular mechanism was also analyzed. Biochemical activities of cells and mitochondria, including those of respiratory rate, mitochondrial membrane potential, and expression levels of mitochondrial and glycolytic genes, were determined.

Materials and Methods

Yeast Strains and Culture Conditions

The genotypes and sources of yeast S. cerevisiae strains had been described elsewhere [8,9]. All yeast strains were cultured in YPD complete medium (1% yeast extract, 1% peptone and 2% glucose). Aminoglycoside antibiotic-containing media were prepared by adding 100 mg/ml stock solutions of antibiotics into the YPD medium after sterilization. The final working concentration of each antibiotic was determined according to the data from a minimal inhibitory concentration assay. 100 mM 2-deoxy-glucose (2-DG) stock solution was prepared in DMSO, and the final concentration of 2-DG in YPD was 2.5 mM. (2-DG) stock solution was prepared in DMSO, and the final concentration of 2-DG in YPD was 2.5 mM. Yeast cells were cultured in liquid YPD medium at 30°C overnight. Then cells were harvested and subcultured at the starting density of 2 × 10\textsuperscript{5} cells/ml in both antibiotic-containing and antibiotic-free media for 16 hours. After validation of the phenotypes, cells were harvested in the mid log-phase and subsequently seeded in pre-coated Poly-D Lysine (30 μg/ml) XF 96-well microplates (Sea-horse Bioscience) at 4 × 10\textsuperscript{5} cells per well, then spun down, and inoculated at 30°C for 30 min. Subsequently the oxygen consumption rate was measured according to manufacturer’s instructions on a Seahorse XF96 Extracellular Flux Analyzer. The OCR was shown as picomole oxygen per minute per cell (pmol/min/cell).

Mitochondrial Membrane Potential Assay

For the mitochondrial membrane potential assay, yeast cells were harvested from both antibiotic-free and antibiotic-containing media. 2 × 10\textsuperscript{8} cells were resuspended in 1 ml supernatant and incubated with Rhodamine 123 (5 μg/ml) for 20 min at 30°C in a shaker. The cells were then centrifuged and washed with a phosphate-buffered saline (PBS) 3 times. Cell pellets were resuspended in 20 μl PBS and visualized with a Carl Zeiss 710 LSM microscope [24].

Northern Blot Analysis

Total cellular RNA was obtained from yeast cultures (2.0 × 10\textsuperscript{7} cells) using TRIzol Reagent (Life Technologies) according to the manufacturer’s instructions. Equal amounts (10 μg) of total RNA were separated by electrophoresis through a 1.5% formaldehyde denaturing agarose gel, transferred onto a positively charged nylon membrane (Amersham) and hybridized with a DIG-labeled ATP6-specific antisense RNA probe. The blot was then stripped with stripping buffer (50% formamide, 30 mM Tris/HCl, pH 7.5, 5% SDS) and hybridized with the COX1, CITB, ATP9, 15S rRNA and 218 rRNA probes, respectively. Finally, the blot was hybridized with a nuclear encoded 23S rRNA probe as an internal control. For the transcriptional assay of glycolytic genes, probes specific to HXK2, PFK2 and PFK1 were used in a northern blot, and 23S rRNA was hybridized as an internal control.

Hexokinase Protein Expression Assay

Antibodies were purchased from Santa Cruz Biotechnologies (Santa Cruz, CA, U.S.A.). Tubulin was used as an internal control. The strains were cultured for 16 hours in YPD at 30°C with or without antibiotic, before processing. Protein samples were prepared, and 50 μg of total protein for each sample was loaded and SDS-PAGE was performed as described elsewhere, followed by transfer to a PVDF membrane [25]. Immunoblotting was performed as previously described with the secondary antibody.
goat anti-rabbit (or rabbit anti-goat) IgG conjugated with horseradish peroxidase, followed by development with ECL solution (Santa Cruz).

Statistical Analysis
All experiments were repeated at least three times and the representative data were presented as means±SD. One-way analysis of variance (ANOVA) was performed to determine the significance between groups. *P<0.05 was considered as statistically significant: #P<0.01.

Results
Growth Activities of Yeast Strains in the Presence of Neomycin
Our previous studies had revealed that the mitochondrial 15S rRNA C1477G combination with mto2 null mutation impaired mitochondrial function in yeast [8]. Here we have further determined the effects of aminoglycoside antibiotics and the interrelationships among nuclear modifier gene, mitochondrial rRNA mutation and antibiotics. To examine the functional consequence of antibiotics on yeast, yeast cells carrying mitochondrial 15S rRNA C1477G mutation and/or nuclear mto2 null mutation were cultured in YPD medium containing neomycin. Cellular growth activities were then measured. Yeast of wild type 15S rRNA and MTO2 genotypes were used as positive controls.

The phenotypes of yeast with different genetic backgrounds indicated that a mutation in mitochondrial 15S rRNA leads to neomycin sensitivity. All four strains grew well on the YPD medium and no difference in growth was observed (Figure 2A). However, when grown on medium containing 300 μg/ml neomycin, the growth of the strains carrying mitochondrial 15S rRNA mutations were inhibited compared to wild type P S strains (Figure 2C). These phenotypes were consistent with yeast growth curves in liquid media (Figure 2B, D). When cultured in media containing neomycin, the two mitochondrial mutant strains exhibited delayed logarithmic phases, while the P S strains entered rapidly into logarithmic growth at approximately 6 hours after inoculation.

Although the deletion of the nuclear gene MTO2 had no effect on P S strains, it significantly altered neomycin susceptibility for strains carrying a mitochondrial 15S rRNA mutation. The MTO2(P S) strain was totally inhibited by neomycin after 3 days' incubation (Figure 2C). However, the double mutant mto2(P S) displayed a much better growth activity and was only partially inhibited. Nevertheless, in the aspect of growth curves, the two 15S rRNA mutant strains presented quite different log-phases. The cell density of MTO2(P S) was 0.3 OD600 after 20 hours' incubation in the neomycin medium, while mto2(P S) reached a final cell density of 1.01 OD600. These data strongly indicates that the nuclear gene MTO2 regulates aminoglycoside sensitivity in yeast carrying the mitochondrial 15S rRNA C1477G mutation, and that deletion of MTO2 suppresses this sensitivity.

Mitochondrial Respiratory Rates
To determine the functional impacts of neomycin on yeast mitochondria in vivo, we examined the respiratory rates of the strains by measuring the rate of oxygen consumption. In the absence of neomycin, the wild-type strain MTO2(P S) had a basal respiratory rate of 0.43 fmol/min/cell (Figure 3). Mitochondrial function was slightly disturbed by the mitochondrial 15S rRNA mutation, and the basal respiratory rate of MTO2(P S) was 11.63% lower than that of MTO2(P S). Deletion of the MTO2 gene had a significant influence on mitochondrial respiration under both P S and P R mitochondrial genetic backgrounds, and 60.47% (p = 0.0203) and 71.05% (p = 0.0122) declines were observed in mto2 null strains, respectively.

We further analyzed the respiratory rates of yeast in the presence of neomycin. The respiratory rates of strains carrying the mitochondrial 15S rRNA mutation were significantly inhibited by neomycin as compared to the control group. After neomycin treatment, the respiratory rate of MTO2(P S) and the double mutant mto2(P S) declined by 65.79% and 58.33%, respectively. However, in the P S strains, only 23.26% and 29.41% declines were observed, respectively, suggesting that the wild type mitochondria were much less sensitive to neomycin. These data indicated that neomycin was more toxic to yeast cells carrying the mitochondrial 15S rRNA mutation.

Mitochondrial Membrane Potential Assay
The rhodamine 123 fluorescent dye was used as an indicator of yeast mitochondrial membrane potential as the uptake of the dye was dependent on the mitochondrial inner membrane potential [24]. The fluorescent signal strength was altered by different genetic and antibiotic factors (Figure 4). In the control group without neomycin, MTO2(P S), mto2(P S) and MTO2(P S) all had relatively high basal membrane potentials. However, the double mutant mto2(P S) strain had a much lower membrane potential, equivalent to 29% of that of MTO2(P S). This indicates that the mto2 null mutation and mitochondrial 15S rRNA point mutation have synergistic suppressive effects on mitochondrial function.

When cultured in media containing neomycin, the mitochondrial membrane potentials in yeast cells were inhibited to different extents. The membrane potentials of the two strains with a wild type mitochondrial genetic background declined by 11% and 19% after neomycin treatment, respectively. This suggests that neomycin has only a slight effect on wild type mitochondria. In contrast, the mitochondrial membrane potential of MTO2(P S) was significantly decreased by 86%, and the green fluorescent signal was almost invisible compared to the untreated control. Moreover, the mitochondrial membrane potential in the mto2(P S) strain was further attenuated by neomycin, and in a manner somewhat parallel to the change of its respiratory rate.

Mitochondrial DNA Transcription Assay
The mitochondrial 15S rRNA C1477G mutation affected steady levels of mitochondrial rRNAs. In the control group, MTO2(P S) and mto2(P S) had significantly lower 15S rRNA transcripts than wild type P S strains (Figure 5A). This suggested that the C1477G mutation had impaired the stability of 15S rRNA. However, the expression patterns of 21S rRNA between P S strains were quite different, suggesting that the large subunit and small subunit of mitochondrial ribosomes were differentially affected by the C1477G mutation. Unexpectedly, neomycin had no significant inhibitory effect on 15S rRNA and 21S rRNA in the P R strains.

Previous studies have revealed that the mto2 null mutation has effects on the expression of mitochondrial genes and the maturation of COX1 and CYTB primary transcripts [8,26]. We further performed transcription assays in the presence of neomycin. The mitochondrial 15S rRNA C1477G mutation affected the expression levels of COX1 and ATP6, but CYTB and ATP9 were less affected (Figure 5C). This result suggests that the transcription of mitochondrial genes in yeast may be differentially regulated. A novel phenomenon was observed that neomycin significantly affected precursor maturation of COX1 and CYTB in the mto2(P S) strain. However, the accumulation of precursors in the MTO2(P S) was less observed. This may indicate that MTO2
interacts functionally with the mitochondrial RNA processing machinery.

Expression of Regulatory Genes in Glycolytic Pathway

Hexokinase (HXK), phosphofructokinase (PFK) and pyruvate kinase (PYK) are key regulators in the glycolytic pathway, where PFK and PYK are enzymes involved exclusively in glycolysis [27]. Steady-state levels of HXK2, PFK1 and PYK1 mRNAs were analyzed by northern blot using DIG-labeled anti-sense RNA probes. The blot was then stripped and re-hybridized with a 25S rRNA probe as an internal control. In the absence of neomycin, the transcription levels of glycolytic genes differed slightly (Figure 6A). The MTO2(PR) strain had a lower HXK2 mRNA level, but the PFK1 and PYK1 transcription levels were higher than those in other three strains. However, after neomycin treatment, all of the three genes in MTO2(PR) were further down regulated. In mto2(PR) strains, HXK2, PFK1 and PYK1 transcription levels were simultaneously stimulated by neomycin and up-regulated by 89%, 112% and 55%, respectively. In contrast, transcription levels of these three genes in MTO2(PR) were further impaired by 14%, 54% and 32%, respectively.

The expression level of hexokinase was also analyzed by western blotting with an anti-hexokinase antibody. Hexokinase catalyzes the initial step of glycolysis, which irreversibly converts glucose to glucose-6-phosphate, a step which also plays an important regulatory role in glycolysis [28]. When cultured in media without neomycin, the hexokinase levels in mto2(PS), MTO2(PS) and mto2(PR) strains were 83%, 84% and 64% compared to wild type MTO2(PS) (Figure 6C). After neomycin treatment, only the double mutant mto2(PR) had an increased hexokinase level, which was 82% higher than its control. However, the other three strains had 12%, 5%, and 30% declines compared to the non-treated controls. These data indicated that the glycolytic pathway in the mto2(PR) strain was up-regulated after neomycin treatment. In summary, we concluded that the MTO2(PR) strain was significantly inhibited by neomycin because both aerobic and anaerobic metabolisms were suppressed. The up-regulated glycolysis in the mto2(PR) strain may

Figure 2. Growth activities of different yeast strains. A Series dilutions of each strain were spotted onto a 2% glucose medium (YPD) and the plate was incubated at 30°C for 72 hours. B Growth curves analysis of yeast strains in the absence of neomycin in 20 hours. C Growth activities of each strain when grown on medium containing neomycin after 72 hours incubation. D Growth curves of strains cultured in YPD containing 32 μg/ml neomycin.
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Figure 3. Assay of Oxygen consumption rates. Cells were harvested in the mid log-phase. The oxygen consumption rate of each yeast strain was measured at the density of 4×10^5 cells/well by FX-96 oxygraph (SeaHorse Biosciences) in the absence or presence of 32 μg/ml neomycin. The results are shown as means±SD of triplicate.
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compensate for mitochondrial dysfunction induced by neomycin and thus this strain was less sensitive to neomycin.

Effects of Glycolytic Inhibitor on Growth Activities of Yeast Strains

2-DG, as a glucose analogue, competitively inhibits hexokinase and blocks the glycolytic pathway [29]. On a YPD medium supplemented with 2.5 mM 2-DG, the double mutant mto2(PR) strain cannot grow because of its dependence on the glycolytic metabolism, but MTO2(P), mto2(P) and MTO2(PR) strains can maintain their growth through mitochondrial respiration (Figure 7). On a YPD medium supplemented with 2.5 mM 2-DG and 300 μg/ml neomyocin, the mto2(PR) strain exhibits slight growth (Figure 7). These data were consistent with the assays of oxygen consumption rates and steady-state level of glycolytic genes.

Discussion

In this study, we analyzed the interactions among the mitochondrial 15S rRNA C1477G mutation, the nuclear modifier gene MTO2, and neomycin, as well as their combined effects on mitochondrial function. The results suggest that yeast cells carrying the C1477G mutation in the mitochondrial 15S rRNA gene exhibit hypersensitivity to neomycin, while deletion of MTO2 gene suppresses this sensitivity. Kutzleb first described paromomycin-resistance (PR) as a consequence of 15S rRNA C1477G mutation, because yeast with this mutation could survive on 2 mg/ml paromomycin-containing medium [12]. In contrast, Weiss-Brummer and colleagues reported that the C1477G mutation alone was not sufficient to form a paromomycin-resistant phenotype [12,13]. Our results supported Weiss-Brummer’s study. When cultured in a YPD medium containing a low dosage of neomycin, yeast cells with the mitochondrial C1477G mutation were significantly inhibited compared to those of the wild type strain. Moreover, PR strains were also sensitive to other aminoglycosides, such as paromomycin and ribostamycin (Figure S1). The MIC90 values of the neomycin of the wild type PS strains were both 128 μg/ml, while the MTO2(PR) and mto2(PR) had the MIC90 values of 32 μg/ml and 64 μg/ml, respectively (Table S1). A spot assay displayed a totally inhibited phenotype of MTO2(PR) by neomycin, while the wild type MTO2(PR) remained unaffected. Functional assays revealed that both the mitochondrial respiratory rate and membrane potential were aggravated after neomycin treatment. Meanwhile, the transcription levels of mitochondrial genes in C1477G mutant strains were sharply attenuated compared to those of the 15S rRNA wild type strains. Neomycin and other aminoglycosides are known to exert antibacterial effects by disrupting the elongation step in protein synthesis with ribosomes [30]. These antibiotics interfere with translation by inhibiting the translocation of the tRNA-mRNA complex from the ribosomal A site to the P site [31]. Several mutations in eukaryotic mitochondrial rRNA genes, such as those of human A1555G and C1494T mutations, alter the binding activity of aminoglycosides to ribosomes [3,15]. The binding of antibiotics to mutant rRNA may further interfere with the movement of the tRNA-mRNA complex on the ribosome [32]. The yeast mitochondrial C1477G mutation is well known to form an identical secondary structure to that of

![Figure 4. In vivo staining of the mitochondrion to measure the membrane potential by Rhodamine 123 dyes.](image-url)
human A1555G and C1477G mutations, and the structure of A1555G and C1494T mutant forms in humans is highly susceptible to neomycin binding [14]. Therefore, the yeast mitochondrial ribosome with the C1477G mutation may exhibit a comparable affinity to neomycin [8,9].

Another interesting finding is that the nuclear modifier gene MTO2 regulates the antibiotic sensitivity of yeast carrying the C1477G mutation. The MIC90 value of neomycin in mto2(PR) was elevated 2-fold compared to that in MTO2(PR), suggesting that the mto2 null mutant was less sensitive to neomycin in the mitochondrial C1477G mutation. The growth curves clearly indicated that the mto2(PR) strain had a more vigorous growth activity than MTO2(PR) in the presence of neomycin. We also studied the neomycin intake and the expression levels of neomycin-resistant genes in these strains. No difference was observed (data unpublished). Expression assays of genes involved in the glycolytic pathway displayed distinct patterns between the two C1477G mutant strains when treated with neomycin. The transcription levels of HXK2, PFK1 and PK1 in MTO2(PR) strains were further suppressed after neomycin exposure, while in the mto2(PR) strain these genes were up-regulated as compared to neomycin-free controls. This result indicates that, although deletion of the MTO2 gene reduces oxidative phosphorylation in the presence of a C1477G mutation, it might also activate the glycolytic metabolic pathway and make the strain more dependent on glycolysis for its energy supply. Thus, the double mutant strain mto2(PR) exhibits a partially restored phenotype in a neomycin-containing medium. We also confirmed that the up-regulated glycolytic pathway in mto2(PR) cells could be suppressed by the hexokinase inhibitor 2-DG. This suppression again made the mto2(PR) strain sensitive to neomycin.

MTO2 is highly conserved evolutionally from E.coli to humans, and in all eukaryotes this gene encodes a mitochondrial-specific tRNA 2-thiouridylase, catalyzing the formation of cmnm5s2U34 in mitochondrial tRNAGlu, tRNAGln and tRNALys [17,33]. According to our data, we suspect that the human homolog of MTO2 may also regulate the aminoglycoside sensitivity of the A1555G mutation. Indeed, according to some epidemiological studies, the aminoglycoside sensitivity of A1555G mutation carriers varies between different ethnic populations where Asian people, for example, developed deafness more rapidly and severely after exposure to aminoglycosides. This could be associated with
different nuclear genetic background, such as those of the various TRMU genotypes. Recently, a series of reports regarding TRMU mutation has been published, and it seems that these mutations are much more common in European and Latin American populations [19,34,35,36]. In contrast, no mutation in the TRMU gene has thus far been identified in Asian populations, such as Chinese or Korean [37,38,39]. Our study may therefore provide a novel

**Figure 6. Steady-state level of key glycolytic genes.**  
**A**, northern blot analysis of HXX2, PFK1 and PYK1 transcription levels. The nuclear encoded 25S rRNA was hybridized as an internal control. **B**, relative expression levels of these genes were calculated by three determinations. **C** translational levels of hexokinase in each strain, tubulin was as an internal control. **D**, calculation of the relative expression level of hexokinase.  
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**Figure 7. Properties of yeast cells on YPD plate in the presence of glycolytic inhibitor 2-Deoxy-D-glucose (2-DG).** Series dilutions of each strain were spotted on medium and the plate was incubated at 30°C for 72 hours. **A** on YPD medium. **B** on YPD medium supplemented with 2.5 mM 2-DG. **C** on YPD medium supplemented with 2.5 mM 2-DG and 300 μg/ml neomycin.  
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insight into the pathogenesis of non-syndromic deafness induced by aminoglycoside antibiotics.

Supporting Information

Figure S1 Effects of paromomycin and ribostamycin on yeast carrying mitochondrial C1477G mutation and wild type allele. 10-fold dilutions of each strain were spotted onto YPD or YPD containing indicated antibiotic, and the plates were incubated for 3 days at 30°C. (TIF)

Table S1 MIC90 values (μg/ml) of neomycin and paromomycin on yeast. (DOC)

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