Generation and bioenergetic analysis of cybrids containing mitochondrial DNA from mouse skeletal muscle during aging

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ABSTRACT
Mitochondrial respiratory chain defects have been associated with various diseases and normal aging, particularly in tissues with high energy demands including skeletal muscle. Muscle-specific mitochondrial DNA (mtDNA) mutations have also been reported to accumulate with aging. Our understanding of the molecular processes mediating altered mitochondrial gene expression to dysfunction associated with mtDNA mutations in muscle would be greatly enhanced by our ability to transfer muscle mtDNA to established cell lines. Here, we report the successful generation of mouse cybrids carrying skeletal muscle mtDNA. Using this novel approach, we performed bioenergetic analysis of cells bearing mtDNA derived from young and old mouse skeletal muscles. A significant decrease in oxidative phosphorylation coupling and regulation capacity has been observed with cybrids carrying mtDNA from skeletal muscle of old mice. Our results also revealed decrease growth capacity and cell viability associated with the mtDNA derived from muscle of old mice. These findings indicate that a decline in mitochondrial function associated with compromised mtDNA quality during aging leads to a decrease in both the capacity and regulation of oxidative phosphorylation.

INTRODUCTION
Mitochondria are ubiquitous organelles in eukaryotic cells whose primary function is to generate energy through oxidative phosphorylation (1), the step-wise transfer of electrons to oxygen coupled with ATP production. High-energy electrons, which leave the electron transport chain (ETC) prematurely, generate reactive oxygen species (ROS) (2). Disruption of ETC has been suggested to alter the production of ROS (3). Oxidative damage caused by ROS to DNA, proteins and lipids in animal tissues has been reported, and the quantity of such damage increases during aging (4,5).

Skeletal muscles, which contain long fibers formed from a large number of fused cells, support or move the skeleton. The performance of skeletal muscle, speed and strength, deteriorates with aging due to fiber loss and atrophy (6), a condition known as sarcopenia. Cytochrome c oxidase (COX), the terminal component of ETC, -negative muscle fibers have been detected in human, monkey, mouse and Drosophila, and their quantity increases with age (7). In addition, studies carried out with different ages of rhesus monkeys indicate regions of muscle fibers exhibiting high levels of ETC defects also contain a large accumulation of mtDNA deletions (8). Further, mtDNA deletions in old animals could focally accumulate to high levels in certain muscle fibers, and that could directly contribute to sarcopenia (9,10). Interestingly, in calorie-restricted mice, a consistently proven animal model of extended mean and maximum life span and attenuated development of age-related diseases (11,12), both the accumulation of deleted mtDNAs and the number of COX-negative fibers, were decreased (13).

Mitochondrial function is under dual genome control. Although the majority of structural genes and all of the regulatory genes are encoded by the nuclear genome, the mitochondrial genome contains the sequences for 37 genes, including 13 for essential polypeptides of the oxidative phosphorylation apparatus, 22 tRNAs and...
two rRNAs for mitochondrial protein synthesis (14,15). It is believed that since mtDNA is close to the ROS generation site, it undergoes relatively more cycles of replication (especially in postmitotic cells such as muscles), and is subject to a reduced breadth of DNA repair activities. Consistent with this hypothesis, mtDNA exhibits a relatively high mutation frequency (16).

The successful development of mitochondrial-mediated transformation has opened the way to elucidate the role of mtDNA (17). However, in-depth studies of the contributions of mtDNA mutations to tissue-specific pathogenesis are limited as the transfer of mitochondria from tissues to cell lines has only successfully been carried out with platelets (18) and synaptosomes (19,20). A noticeable missing technique is a method to establish cybrids with mtDNA derived from muscle. In the present study, we developed a new approach to generate cybrids containing mtDNA from mouse skeletal muscle. Using this method, we further analyzed respiratory activity and growth capacity in glucose and galactose media of cybrids with mtDNA from old and young mice.

MATERIALS AND METHODS

Animals and sample preparation

C57BL/6 and DBA/2 mice were purchased from Charles River Laboratories. Among them, six each of C57BL/6 were aged to 6 and 26 months at the UTHSCSA animal facility. Skeletal muscles were dissected from the hind legs. Each muscle was cut into small pieces on a 10-mm dish on ice, and rinsed several times with Hank’s Balanced Salt Solution (HBSS, Invitrogen). The muscle samples were resuspended with 6 vol hypotonic buffer (RSB) (10.0 mM KCl, 0.15 mM MgCl2, pH 6.7) and further processed with a Dounce A type homogenizer for about 30 strokes. The processed samples were then centrifuged at 2300 r.p.m. for 3 min and the supernatant was then subjected to another centrifugation at 1000 r.p.m. for 10 min. Finally, muscle mitochondria enclosed by plasma membrane were collected by centrifugation at 6500 r.p.m. for 10 min.

Cell lines and media

All cell lines used in the present work were grown in monolayer culture. The mtDNA-less pB LL/2-m21 cell line was a derivative of mouse cell line LL/2 as described previously (21). Cells were grown in Dulbecco’s Modified Eagle Medium (DMEM) supplemented with 10% fetal bovine serum (FBS).

mtDNA analysis

For mtDNA sequencing, total DNA was extracted with the Easy-DNA™ kit (Invitrogen, Carlsbad, CA, USA) according to the manufacturer’s instructions, and then subjected to polymerase chain reaction (PCR) amplification of the fragments containing part of the tRNAArg and ND3 genes. Amplified mtDNA fragments were purified using the AllStar PCR Clean Up kit (AllStar Sci. Inc., Sunnyvale, CA, USA), and purified PCR products were sequenced.

The sequences of primers used were:

ND3-5: (9312-9331) GTGAAGCCGGCACGTGACTACT
ND3-3: (9900-9881) GTTGAAGAAGGTAGATGGCA

O2 consumption measurement

The measurements were carried out in two chambers of a YSI Model 5300 Biological Oxygen Monitor as described earlier (22). Briefly, determination of the O2 consumption rate was carried out in about 5×10⁶ cells in Tris-based, Mg2+, Ca2+-deficient (TD) buffer (0.137 M NaCl, 5 mM KCl, 0.7 mM Na2HPO4, 25 mM Tris–HCl, pH 7.4). After recording the base respiration rate, 2.5 μg/ml oligomycin was added to measure uncoupled respiration, and then 0.5 μM FCCP was added to determine maximal respiration.

Cell growth and viability analysis

Multiple identical samples of 5×10⁴ cells were grown for 4 days on six-well plates in the appropriate medium (DMEM, which contains 4.5 mg/ml glucose and 0.11 mg/ml pyruvate, or DMEM lacking glucose and containing 0.9 mg/ml galactose and 0.11 mg/ml pyruvate, both supplemented with 10% dialyzed FBS), and counted on a daily basis. Doubling time = 72 h×Ln (2)/[Ln (total cell number at 96 h)−Ln (total cell number at 24 h)].

Cell viability was analyzed with a Vi-Cell™ XR cell viability analyzer (Beckman Coulter).

mtDNA content analysis

The relative mtDNA levels were measured by a real-time PCR and normalized by simultaneous measurement of the nuclear DNA. Primers and probes for quantitative PCR (qPCR) were designed using Primer Express (Applied Biosystems). QPCR was carried out using an ABI 7900HT Fast Real-Time PCR System (Applied Biosystems) in 20-μl reaction in different tubes containing 0.3 μM each of the forward and reverse primers, 0.1 μM for each probe (ND3 and β-actin genes) and 50 ng DNA sample. The PCR conditions were 95°C for 15 min, followed by 40 cycles of 95°C for 15 s, and 60°C for 60 s. The threshold cycle number (Ct) values of the β-actin gene and the mitochondrial ND3 gene were determined. Each measurement was carried out four times and normalized against a serial dilution of a control DNA sample. Ct values can be used as a measure of the input DNA contents and Ct value differences were used to quantify mtDNA levels relative to the β-actin gene with the following equation: Relative copy number (R) = 2ΔCt, where ΔCt is the Ctβ-actin − CtND3.

The primer and probe information is as follows:

For mtDNA ND3: forward, TCTGACTCCCCCAAACTGACT; reverse, GGGTCAATCCGGCATTCA.

For nuclear β-actin: forward, AAATCGTGCGTGAC; reverse, GGCCATCTCCCTGCTCAG.

Probes: Cox III, FAM- TCAGAAAAGCAAATCC; β-actin, FAM- AGCTGTGCTATGTGCT.
Statistical analysis

$P$, probability, values were calculated by using the unpaired Student’s $t$-test contained in the Microsoft Excel program. The difference is considered as statistically significant when $P < 0.05$. $r$, Pearson product-moment correlation coefficient, is a measure of the strength of linear dependence between two variables. It ranges from $-1$ to $1$. A value of $1$ implies that a linear equation describes the relationship perfectly; a value of $-1$ implies a perfect negative correlation; a value of $0$ implies that there is no linear correlation between the variables.

RESULTS

Generation of cybrids with mitochondria from mouse skeletal muscle

As shown in Figure 1, muscle samples were obtained from hind legs of healthy mice right after the animals were euthanized. In particular, muscles were dissected out; fatty and nerve cells were removed. After being rinsed with HBSS, the muscle samples were cut into small pieces on ice. Fine muscle pieces were subjected to treatment with hypotonic buffer and then homogenized. The processed muscle was constantly checked under the

Figure 1. Generation of cybrids containing mtDNA from skeletal muscle. Flowchart of how mtDNA from the skeletal muscle was transferred to $\rho^0$ cells to produce cybrids.
microscope after 20 strokes with the homogenizer until the nuclei were released and plasma membrane-enclosed mitochondria formed. The processed samples were then centrifuged at 2300 r.p.m. for 3 min and the supernatant was then subjected to another centrifugation at 1000 r.p.m. for 10 min. Finally, muscle mitochondria enclosed by plasma membrane were collected by centrifugation at 6500 r.p.m. for 10 min.

Fusion solution was prepared as 9 g of polyethylene glycol (PEG) (Sigma P-5402; MW: 1450) was autoclaved with slow exhaust for 15 min, and then 2 ml DMSO and 9 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-m21 β0 cell line was established previously (23). Approximately 65 ml DMEM were added when it was cooled to about 65°C. The solution was mixed gently and then kept in 37°C incubator. The mouse mtDNA-less LL/2-
observed. In particular, in regular glucose medium there was no significant difference in the average doubling time between cells derived from the young and old groups (Figure 4A). However, in the galactose medium, where ATP production predominantly relies on mitochondrial oxidative phosphorylation, the average doubling time of old group cells was markedly longer than that of young group cells (Figure 4B). In order to reduce the background variability caused by other factors, the ratio of doubling time in galactose medium to that in glucose medium was calculated as indicated by the growth index. As shown in Figure 4C, the difference between the two age groups persisted, although with somewhat reduced statistical significance ($P < 0.05$ versus $P < 0.01$).
As mitochondrial dysfunction has been associated with apoptosis, we further investigated cell viability in glucose- and galactose-containing media following 3 days of culturing in the respective media. As shown in Figure 5A, in the regular glucose medium, there was no significant difference in cell viability between the young group and the old group. However, in the galactose medium, the cell viability of the old group was significantly lower than that of the young group ($P < 0.05$) (Figure 5B). The ratio of viability in galactose to that in glucose medium, indicated as viability index, was significantly lower for the old group relative to the young group cells ($P < 0.05$) (Figure 5C).

mtDNA content alteration and its implication

To investigate the alteration of mtDNA content as a result of changes in mtDNA sequences associated with replication or turnover, the mtDNA copy number of the transformants was measured by quantitative real-time PCR method. While a considerable variability among the clones derived from the same age group was revealed, there is a clear tendency of decrease of mtDNA content in the cybrids from old mice (Figure 6A). To further determine if the decrease in mtDNA copy number has any effects on the compromised mitochondrial function we observed, we plotted the

Figure 3. Oxygen consumption measurement. Six and 26 months represent 24 and 22 cybrids per age group, respectively. Endogenous respiration was measured in intact cells (A), after treatment with the ATP synthase inhibitor oligomycin at 2.5 $\mu$g/ml (B), and with the uncoupler FCCP at 0.5 $\mu$M (C). The activities were determined with about 5 x 10^6 cells. Three determinations were made for each cybrid and the average values are shown. Ratios of oligomycin-resistant to native endogenous respiration rates (D) and FCCP-treated fully uncoupled to oligomycin-resistant respiration rates (E) of each age group were also calculated.

Figure 4. Growth capacity measurement. Six and 26 months represent 24 and 25 cybrids per age group, respectively. Cells were grown in glucose-containing DMEM for 24 h, then in glucose medium (A) or in galactose medium (B) for another 72 h. Doubling times were calculated as: $72 \times \log(2)/\log([\text{total cell number at the end}]-[\text{total cell number when media were changed}])$. Three determinations were made for each cybrid. Growth index was calculated as the ratio of doubling time in galactose-containing medium to that in glucose medium (C).
maximal respiration rate measured by oxygen consumption with FCCP treatment (Figure 6B), growth index as defined by the ratio of doubling time of cybrids in galactose medium to that in glucose medium (Figure 6C) and cell viability index as the ratio of cell viability assayed in galactose medium to that in glucose medium (Figure 6D), to mtDNA copy number, respectively. We found that the maximal respiration rate declined as the mtDNA copy number reduced (Figure 6B). Similarly, the normalized growth capacity in galactose medium also reduced with the decline of the mtDNA copy number (Figure 6C).

However, no significant effect was observed in the range of the mtDNA copy number changes in this study on the cell viability in galactose medium (Figure 6D), while we did observe a decrease in cell viability with the same assay in cybrids carrying mtDNA from muscle of old mice (Figure 5C).

These results indicate that the declined mtDNA copy number might contribute to the compromised respiration rate and oxidative phosphorylation capacity associated with the mtDNA from the skeletal muscle of the old mice. In addition, other age-related mtDNA mutations...
in mouse skeletal muscle could also play a role in age-related pathogenesis.

**DISCUSSION**

The successful generation of muscle cybrids provided a unique opportunity to recover and analyze somatic mtDNA mutations directly from fresh muscle samples. This new technology also provided an opportunity to further our ability to investigate diseases or age-related mtDNA mutations. In addition, since low-occurrence mutations within the heteroplasmic mtDNA populations could be enriched in culture cells (24), one great advantage of this new approach is the potential to study the low level pathogenic mutations which could otherwise be very difficult to carry out sophisticated studies on at the molecular level in tissues.

Studies of the role of mitochondrial dysfunction and mtDNA mutations in pathogenesis associated with diseases and aging in skeletal muscle is of particular interest since: (i) muscle is one of the most targeted tissues by mitochondrial deficiency. Actually, one of the earliest identified hallmarks of mitochondrial disease is the characterization of ragged red fibers (RRG) in biopsied skeletal muscle, and interestingly most have been associated with abnormal mtDNA (28). (ii) Skeletal muscle predominantly relies on oxidative phosphorylation and at the same time, suffers rather marked age-related degeneration (29). (iii) Skeletal muscle is arguably the best system where the causative relationship among age-related degeneration, mitochondrial dysfunction, and mtDNA alteration, mtDNA deletions in particular, has been established, owing largely to a collection of excellent work from Judd Aiken and colleagues (30–34).

Our results revealed that respiration measured with FCCP was more sensitive in detecting in mitochondrial function decline associated with aging. The other significant result in the present study was the demonstration that the loss of coupling and regulation of oxidative phosphorylation associated with mtDNA alteration during aging. Such an age-related compromise in coupling efficiency of respiration was reported previously in a large scale investigation with fibroblasts, mostly skin-derived, from human subjects with a large range of age, by measurement of P:O ratios (35).

As previously reported with human skeletal muscle (36) and mitochondrial transformants with mtDNA derived from human fibroblasts (37), an age-related decline of mtDNA contents was revealed with mouse skeletal muscle in this study. However, while we observed the declines in respiration and growth capacity associated with the decrease in mtDNA copy number, we did not see a significant correlation between mtDNA content and cell viability in galactose medium. As shown with the mutant mice, a premature aging mouse model with an increase in mtDNA mutations due to the expression of a proof-reading-deficient version of mtDNA polymerase (38,39), increased mtDNA mutations have been associated with enhanced apoptosis. Our results also suggested that there was an age-related increase in mtDNA mutations which might also play a role in the decreased viability in galactose medium associated with the cybrids of old mice.

As a significant difference between young and old age groups was only detected when cybrids were grown in galactose medium, but not in the regular glucose medium where cells can obtain ATP through both oxidative phosphorylation and glycolysis, these findings suggested that the decrease in mitochondrial ATP production in old mitochondria could be compensated for by an upregulation in the glycolytic pathway. It is interesting to note that such a bioenergetic switch is also a signature of cancer cells (40), as aging is the biggest risk factor for cancer occurrence.

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