

# High-Fat Diet-Induced Mitochondrial Dysfunction Promotes Genioglossus Injury – A Potential Mechanism for Obstructive Sleep Apnea with Obesity

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**Purpose:** Obesity is a worldwide metabolic disease and a critical risk factor for several chronic conditions. Obstructive sleep apnea (OSA) is an important complication of obesity. With the soaring morbidity of obesity, the prevalence of OSA has markedly increased. However, the underlying mechanism of the high relevance between obesity and OSA has not been elucidated. This study investigated the effects of obesity on the structure and function of the genioglossus to explore the possible mechanisms involved in OSA combined with obesity.

**Methods:** Six-week-old male C57BL/6J mice were fed high-fat diet (HFD, 60% energy) or normal diet (Control, 10% energy) for 16 weeks. The muscle fibre structure and electromyography (EMG) activity of genioglossus were measured. The ultrastructure and function of mitochondrial, oxidative damage and apoptosis in genioglossus were detected by transmission electron microscopy (TEM), qPCR, Western blotting, immunohistochemistry and TUNEL staining. We further studied the influence of palmitic acid (PA) on the proliferation and myogenic differentiation of C2C12 myoblasts, as well as mitochondrial function, oxidative stress, and apoptosis in C2C12 myotubes.

**Results:** Compared with the control, the number of muscle fibres was decreased, the fibre type was remarkably changed, and the EMG activity had declined in genioglossus. In addition, a HFD also reduced mitochondria quantity and function, induced excessive oxidative stress and increased apoptosis in genioglossus. In vitro, PA treatment significantly inhibited the proliferation and myogenic differentiation of C2C12 myoblasts. Moreover, PA decreased the mitochondrial membrane potential, upregulated mitochondrial reactive oxygen species (ROS) levels, and activated the mitochondrial-related apoptotic pathway in myotubes.

**Conclusion:** Our findings suggest that a HFD caused genioglossus injury in obese mice. The mitochondrial dysfunction and the accompanying oxidative stress were involved in the genioglossus injury, which may provide potential therapeutic targets for OSA with obesity.

**Keywords:** obesity, obstructive sleep apnea, genioglossus, muscle, mitochondria

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## Introduction

Obesity is a worldwide metabolic disease that has shown increasing morbidity.<sup>1,2</sup> Recent studies have found that obesity characterized by an excessive accumulation of lipids in the blood and subcutaneous and visceral compartments, leads to disorders in glucose and lipid metabolism and increases the risk of several chronic conditions, notably

cardiovascular and respiratory diseases.<sup>3–5</sup> Obstructive sleep apnea (OSA) is one of the most common respiratory diseases and is characterized by intermittent and recurrent collapse of the upper airway during sleep, inducing hypoxia or apnea and causing a significant threat to the health and quality of life of the affected individuals.<sup>6,7</sup> Obesity is highly correlated with the incidence of OSA.<sup>8,9</sup> According to a recent epidemiological study, the prevalence of OSA was four times higher in the obese population than in the general population.<sup>10</sup> The anatomic stenosis of the upper airway and the dysfunction of upper airway dilator muscles were reported to have important implications in the pathogenesis of OSA.<sup>11,12</sup> High-fat diet (HFD)-induced obesity increases the accumulation of fat around the upper airway, aggravates stenosis of the upper airway, and more easily results in airflow reduction or obstruction during sleep.<sup>13–16</sup> However, how the dysfunction of the upper airway dilator muscles in the obese population is involved in the mechanism of OSA has not been elucidated.

The genioglossus is the largest upper airway dilator muscle and plays a very important role in maintaining airway patency.<sup>17</sup> Compared to control subjects, OSA patients show an abnormal structure and function of the genioglossus, with a higher proportion of type II fibres (fast-twitch, glycolytic), which decreases fatigue resistance in the genioglossus.<sup>18</sup> HFD-induced obesity is a metabolic disease. The accumulation of excess free fatty acids (FFAs) in skeletal muscle could impair myogenesis, decrease skeletal muscle mass and reduce muscle strength.<sup>19,20</sup> However, there have been few studies addressing the effects of obesity on genioglossus structure and function.

Mitochondria are the primary energy source in muscle cells. Mitochondria produce adenosine triphosphate (ATP) via the oxidative metabolism of glucose and fatty acids, which is essential for the maintenance of skeletal muscle function.<sup>21</sup> In the context of HFD-induced obesity, excess lipids in skeletal muscle suppress glucose oxidative metabolism and lead to disturbances in mitochondrial function. A large body of studies have shown that mitochondrial damage is involved in obesity-induced skeletal muscle dysfunction, impairing mitochondrial biogenesis, lowering mitochondrial content and reducing the activity of mitochondrial enzymes.<sup>22–24</sup> Moreover, mitochondrial damage was also found in the genioglossus of chronic intermittent hypoxia (CIH) rats.<sup>25</sup> Therefore, it is essential for us to investigate the potential influence of obesity on the mitochondria in the genioglossus, which will be beneficial to better understand the underlying mechanism of OSA that occurs with obesity.

Mitochondrial dysfunction contributes to the excessive accumulation of reactive oxygen species (ROS) and oxidative stress, which causes oxidative damage to major cellular structures (in particular, the plasma membrane, proteins and DNA) and leads to apoptosis.<sup>26</sup> In HFD-induced obesity, oxidative damage accompanied by mitochondrial impairment has been found, which is a critical factor for skeletal muscle dysfunction.<sup>27,28</sup> Our previous work that focused on OSA demonstrated that ROS overproduction caused by hypoxia contributed to pyroptosis in myoblasts and might be a critical pathogenesis of OSA.<sup>29</sup>

Therefore, we hypothesized that mitochondrial dysfunction and oxidative stress may be involved in the effects of HFD-induced obesity in the genioglossus, which is an important pathogenesis of OSA combined with obesity. To address this hypothesis, we examined the changes in genioglossus activity and structure, as well as the mitochondrial ultrastructure and function, in HFD-induced obese mice. In addition, the underlying preliminary mechanisms have been investigated, including oxidative stress and apoptosis in the genioglossus. We also used palmitic acid (PA), one of the most abundant saturated free fatty acids (FFAs) in humans, to mimic HFD-induced obesity in an *in vitro* study.<sup>30</sup> We studied the influence of PA on the proliferation and myogenic differentiation of C2C12 myoblasts, as well as mitochondrial function, oxidative stress, and apoptosis in C2C12 myotubes.

## Materials and Methods

### Animals

Six-week-old male C57BL/6J mice were provided by Shanghai Model Organisms Center and housed under controlled environmental conditions ( $22 \pm 2$  °C,  $50\% \pm 5\%$  humidity, 12:12 h light/dark cycle). The mice were randomly divided into two groups: the control group (normal diet, 10% energy; Research diet D12450J) and the HFD group (60% energy; Research diet D12492). All mice had free access to food and water under specific pathogen-free (SPF) conditions, and the mice body weights were recorded weekly. The animal study was approved by the Animal Welfare and Ethics Group, Department of Laboratory Animal Science at Fudan University. All procedures were performed according to Laboratory animal – Guideline for ethical review of animal welfare (GB/T 35892–2018, standardization administration of the People's Republic of China). After 16 weeks, the mice were euthanized, and samples were collected. Before

sample collection, the mice were fasted overnight and anaesthetized with chloral hydrate. Blood was collected from the retro-orbital plexus and centrifuged at 3000 rpm for 10 minutes. Plasma was collected and used for clinical biochemical analyses. Additionally, each mouse genioglossus was harvested for subsequent analysis.

### Blood Biochemical Assay

Chemical analysis of the serum samples ( $n = 6$ ) was carried out with an automatic biochemistry analyser (Chemray 800, Rayto Life and Analytical Sciences Co., Shenzhen, China). The following parameters were evaluated: triglycerides (TGs) and cholesterol (CHOL). The levels of FFAs were measured with commercial kits (Nanjing Jiancheng Bioengineering Institute, Nanjing, China).

### Intraperitoneal Glucose Tolerance Test (IPGTT) and Insulin Tolerance Test (IPITT)

At the end of the 16-week experiment, the IPGTT and IPITT were performed. For the IPGTT, mice were intraperitoneally injected with glucose (Sigma–Aldrich, St. Louis, MO, USA; 1.5 g/kg body weight) after 12 h of fasting. For the IPITT, mice were intraperitoneally injected with insulin (Sigma–Aldrich, St. Louis, MO, USA; 0.5 U/kg body weight) after 4 h of fasting. Blood samples ( $n = 6$ ) were taken from the tail vein, and the blood glucose levels were measured at 0, 15, 30, 60, and 120 minutes using a glucometer (Roche, USA).

### Electromyography (EMG)

EMG of the genioglossus muscle was acquired and analysed as described in our previous study.<sup>31</sup> Briefly, six mice from each group were first anaesthetized with 1% pentobarbital, and then the genioglossus was exposed. EMG electrodes (AD Instrument, Australia) were inserted into the genioglossus, and the EMG signal was amplified and recorded (LabChart 8, AD Instrument, filter frequency was 1–1000 Hz, time constant: 1 s).

### Histological Examination and Immunohistochemical Assay

The genioglossus samples were harvested and immediately fixed in ice-cold 4% paraformaldehyde solution. The histopathological tests were performed using standard laboratory procedures. Briefly, the genioglossus samples were embedded in paraffin and cut into 5  $\mu\text{m}$  sections, which

were mounted onto glass slides for haematoxylin and eosin (H&E), Masson's trichrome, myosin heavy chain (MyHC)-slow, MyHC-fast, 4-hydroxynonenal (4-HNE) and 8-hydroxy-2'-deoxyguanosine (8-OHdG) staining. For histological examinations, genioglossus sections were stained with micrographs of H&E and Masson's trichrome staining (Solarbio, Beijing, China). For tissue immunohistochemistry, after being deparaffinized and rehydrated, the genioglossus sections were boiled in 10 mM citrate buffer for 10 minutes, processed with 0.25% Triton X-100 for 30 minutes and blocked with a 5% bovine serum albumin (BSA) solution at room temperature for 30 minutes. Next, the sections were incubated overnight at 4 °C with primary antibodies against MyHC-slow (1:1000, Abcam, UK), MyHC-fast (1:1000, Abcam, UK), 4-HNE (1:200, Abcam, UK) and 8-OHdG (1:200, Abcam, UK). Then, staining was carried out with an immunodetection kit and DAB peroxidase substrate kit according to the manufacturer's instructions (ZSGB-BIO, Beijing, China). Micrographs were acquired with a light microscope (DM2500, Leica, Wetzlar, Germany). Three to ten fields of view were randomly selected for quantification analysis.

### Transmission Electron Microscopy (TEM)

The genioglossus samples were sliced into several tissue pieces (each 1 × 1 × 1 mm) and fixed in 2.5% glutaraldehyde at 4 °C. Then, the samples were incubated in 0.1 M phosphate buffer with 1% osmium tetroxide for 2 h, dehydrated, embedded in epoxy resin, and polymerized for 48 h at 60 °C. Subsequently, the sections were cut into ultrathin sections (50–70 nm), stained with uranyl acetate-lead citrate, and examined with a transmission electron microscope (JEOL-1200, JEOL electron, Japan).

### Mitochondrial DNA (mtDNA) Copy Number Determination

The genomic DNA of genioglossus samples ( $n = 6$ ) was isolated with the DNA Extraction Kit (Tiangen, DP304, Beijing, China) according to the manufacturer's instructions. QPCR was used to detect the copy numbers of nuclear DNA and mtDNA with SuperReal PreMix Plus (Tiangen, FP205, Beijing, China) on a LightCycler 480 System (Roche, Basel, Switzerland). The relative copy number of mtDNA was determined by the comparative cycle threshold (Ct) value of nuclear DNA. The primers of nuclear DNA: (F) CGTGGGCTCCAGCATTCTA and (R) TCACCAGTCATTTCTGCCTTTG; and the primers

of mtDNA: (F) CGAAAGGACAAGAGAAATAGAG and (R) GAACAAGGTTTTAAGTCTTACGCA.

## Cell Culture and Palmitic Acid Treatment

C2C12 mouse myoblasts from the Cell Bank of Chinese Academy of Sciences Shanghai Institute of Cell Biology (Shanghai, China) were cultured in growth medium (GM) (high-glucose DMEM with 10% FBS (Gibco, MA, USA) and 1% penicillin/streptomycin (Gibco, MA, USA)) at 37 °C in 5% CO<sub>2</sub>/95% air. For myogenic differentiation, when C2C12 myoblasts were 100% confluent, GM was switched to high-glucose DMEM containing 2% horse serum (Gibco, Thermo Fisher Scientific, Waltham, MA, USA) and 1% penicillin/streptomycin for 5 days, with medium change every 2 days.

In vitro, PA was used to mimic the high-fat environment. PA (Sigma–Aldrich, St. Louis, MO, USA) was dissolved in absolute ethanol to generate a 100 mM stock solution after filtration through a 0.22 μm PES membrane. Then, the PA stock solution was added to serum-free DMEM containing 5% BSA (Sigma–Aldrich, St. Louis, MO, USA) at final concentrations of 100 μM, 300 μM and 500 μM at 37 °C with continuous shaking for 2 h. Serum-free DMEM containing 5% BSA with absolute ethanol was used as the control. To examine the effect of PA on the proliferation of C2C12, the C2C12 myoblasts were treated with different concentrations of PA for 6 h to 48 h. To assess the effect of PA on the myogenic differentiation of C2C12, the C2C12 myoblasts were treated with different concentrations of PA for 2 days during cellular differentiation. To investigate the effect of PA on the mitochondrial function of C2C12 myotubes, after 5 days of myogenic differentiation, the myotubes were treated with different concentrations of PA for 2 days. In some experiments, in order to detect the role of PA-induced mitochondrial ROS in the damage of C2C12 myotubes, Szeto-Schiller (SS) peptide SS-31 (Selleckchem, Houston, TX, USA) was used to treat C2C12 myotubes at 10 μM for 2 days. SS-31 is reported to target mitochondria and reduce mitochondrial ROS.<sup>32,33</sup>

## Cell Viability Assay, Lactic Dehydrogenase (LDH) Measurements and Cell Morphological Imaging

Cell viability was determined using a Cell Counting Kit-8 assay (CCK-8, Dojindo, Japan). In brief, C2C12 myoblasts were seeded in 96-well plates at a density of 5 × 10<sup>3</sup> cells per well and allowed to attach for 24 h. Then, the C2C12 myoblasts were treated with PA at different concentrations (100

μM, 300 μM and 500 μM). After treatment for 0 h, 6 h, 12 h, 24 h, and 48 h, the cells were incubated with CCK-8 for 1 h, and the absorbance values at 450 nm were measured with an Epoch 2 microplate spectrophotometer (BioTek, VT, USA). After treatment with PA for 24 h, the morphological changes in the C2C12 myoblasts were detected with a Leica DMI8 microscope (Leica, Wetzlar, Germany).

Cytoplasmic LDH was released into the medium following cell death. The amount of LDH in the extracellular medium was determined using a commercial LDH kit (Beyotime, Nanjing, China). Briefly, C2C12 myoblasts were seeded in 96-well plates and treated with PA for 24 h. Then, the culture supernatants were collected and centrifuged at 400 × g for 5 minutes. LDH levels were detected according to the manufacturer's protocol, and the absorbance at 495 nm was measured with an Epoch 2 microplate spectrophotometer.

## Flow Cytometry Detection of Apoptosis

Cell apoptosis was measured using an Annexin V-FITC/propidium iodide (PI) apoptosis detection kit (Beyotime, Nanjing, China). After treatment with PA for 24 h, C2C12 myoblasts were harvested and incubated with annexin V-FITC and PI for 15 minutes according to the manufacturer's instructions. The rate of apoptosis was measured using flow cytometry (ACEA NovoCyte, Hangzhou, China) and analysed with Novo Express software 1.4.1 (ACEA Biosciences).

## Immunofluorescence Cell Staining

C2C12 myoblasts were incubated with different concentrations of PA during cellular differentiation for 2 days. Then, the myotubes were washed with PBS, fixed in 4% paraformaldehyde for 15 minutes, permeabilized with 0.25% Triton X-100 for 10 minutes and blocked with 5% BSA for 1 h at room temperature. Subsequently, the cells were incubated overnight at 4 °C with primary antibodies against MyHC (1:10, MF20 DSHB, USA). The following day, the cells were incubated with the secondary antibody for 1 h, and the nuclei were counterstained with DAPI for 5 minutes at room temperature. Finally, fluorescent images were captured using a fluorescence microscope (DMI8, Leica, Wetzlar, Germany).

## MitoTracker Green and Mitochondrial Membrane Potential (MMP) Fluorescent Staining

After myogenic differentiation of C2C12 for 5 days, the myotubes were treated with different concentrations of PA

for 2 days. Then, C2C12 myotubes were incubated with MitoTracker green dye (Beyotime, Nanjing, China) at 37 °C for 30 minutes to observe changes in mitochondrial morphology. The nuclei were counterstained with DAPI for 5 minutes at 37 °C. Images were immediately acquired with a fluorescence microscope (DMI8, Leica, Wetzlar, Germany). Moreover, the MMP was evaluated using a tetraethylbenzimidazolylcarbocyanine iodide kit (JC-1, Beyotime, Nanjing, China) according to the manufacturer's instructions. C2C12 myotubes were incubated with different concentrations of PA for 2 days. Then, the myotubes were stained with 10  $\mu$ M JC-1 for 20 minutes at 37 °C. The nuclei were counterstained with DAPI for 5 minutes at 37 °C, and images were immediately acquired with a fluorescence microscope.

### Mitochondrial ROS Determination

Mitochondrial ROS was evaluated using the mitochondrial superoxide indicator MitoSOX Red (Invitrogen, Thermo Fisher Scientific, Waltham, MA, USA) according to the manufacturer's instructions. Briefly, after myogenic differentiation of C2C12 for 5 days, the myotubes were treated with PA or SS-31 for 2 days. Then, C2C12 myotubes were washed twice with PBS and incubated in 5  $\mu$ M MitoSOX Red for 30 minutes at 37 °C. The nuclei were counterstained with DAPI for 5 minutes at 37 °C, and images were immediately acquired with a fluorescence microscope.

### Apoptosis Assay

A TUNEL assay was performed to detect apoptotic cells in genioglossus tissues or C2C12 myotubes using a One-Step TUNEL Assay Kit (Beyotime, Shanghai, China) according to the manufacturer's instructions. Briefly, the fixed genioglossus sections or C2C12 myotubes were incubated with TUNEL reaction mixture for 30 minutes at 37 °C and then counterstained with 4',6-diamidino-2-phenylindole (1:10000, DAPI, Abcam, UK) for 5 minutes. Images were acquired with a fluorescence microscope (DM2500, Leica, Wetzlar, Germany).

### Western Blotting

Total protein was extracted from genioglossus tissues or C2C12 myotubes using RIPA buffer (Beyotime, Nanjing, China) with protease and phosphatase inhibitors. To detect the protein levels of Cyt-c in the mitochondria and cytoplasm of genioglossus tissues, mitochondrial and cytoplasm proteins were isolated using a mitochondria isolation kit

(Beyotime, Nanjing, China). Protein concentration was determined with a bicinchoninic acid (BCA) Protein Assay kit (Thermo Fisher Scientific, Waltham, MA, USA). Total protein was separated by 10% SDS polyacrylamide gel electrophoresis (SDS-PAGE) and transferred onto polyvinylidene fluoride (PVDF) membranes using a Bio-Rad system (Hercules, CA). The membranes were incubated overnight at 4 °C with primary antibodies. Antibodies against DRP1 (1:1000), cleaved caspase-3 (1:1000) and Bax (1:1000) were purchased from Cell Signaling Technology (BSN, USA). Antibodies against MFN2 (1:300), Bcl-2 (1:1000), Cyt-c (1:1000) and Cox IV (1:1000) were purchased from Proteintech Group (PA, USA). Antibodies against MyoG (1:500), MyoD1 (1:500) and GAPDH (1:500) were purchased from Santa Cruz Biotechnology (TX, USA). The MyHC antibody (1:60) was purchased from Developmental Studies Hybridoma Bank (IA, USA). The OXPHOS antibody (1:1000) was purchased from Abcam (Cambridge, UK). Horseradish peroxidase-conjugated secondary antibodies (1:10000, Cell Signaling Technology, BSN, USA) were applied the following day and incubated with the membranes at room temperature for 1.5 h. Then, the membranes were developed with a chemiluminescent substrate kit (Thermo Fisher Scientific, Waltham, MA, USA), and the bands were detected with Amersham Imager 600 (GE Health care, IL, USA). GAPDH was used as an internal control in the total protein and cytoplasm protein, Cox IV was used as an internal control in the mitochondria protein.

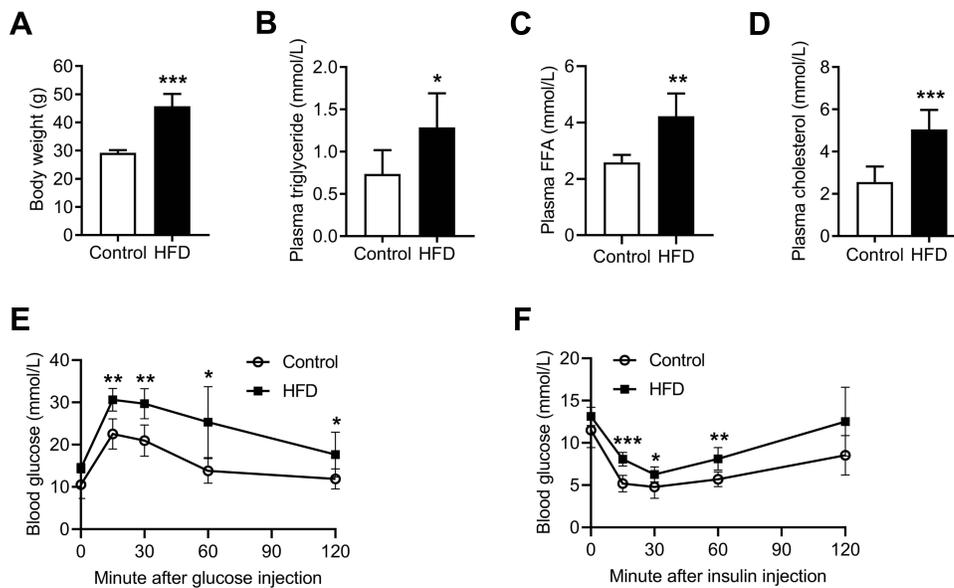
### Statistical Analysis

Data were analysed using GraphPad Prism 7.0 (GraphPad Software, San Diego, CA, USA) and expressed as the mean  $\pm$  SD. Differences between groups were determined using Student's *t*-test or one-way ANOVA. A *p* value < 0.05 was considered statistically significant.

## Results

### Effects of HFD on Glucose and Lipid Metabolism in Mice

The body weights of the HFD mice significantly increased after 16 weeks, and these mice were markedly heavier than the control group (*p* < 0.001) (Figure 1A). Moreover, the serum TG, FFA and CHOL levels were all elevated in the HFD group (Figure 1B–D). In addition, the IPGTT and the IPITT were performed, and the data showed that HFD induced systemic insulin resistance, with a significant reduction in glucose tolerance and insulin sensitivity (Figure 1E and F).



**Figure 1** HFD induced obesity and caused glucose and lipid metabolism disorders in C57BL/6j mice. **(A)** Body weights of control and HFD-fed mice at the end of 16 weeks. **(B)** Plasma triglyceride, **(C)** plasma FFA, and **(D)** plasma cholesterol of the control and HFD-fed mice at the end of 16 weeks. **(E and F)** IPGTT and IPITT of the control and HFD-fed mice at the end of 16 weeks. Data are represented as the mean  $\pm$  SD. \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$  compared to the control.

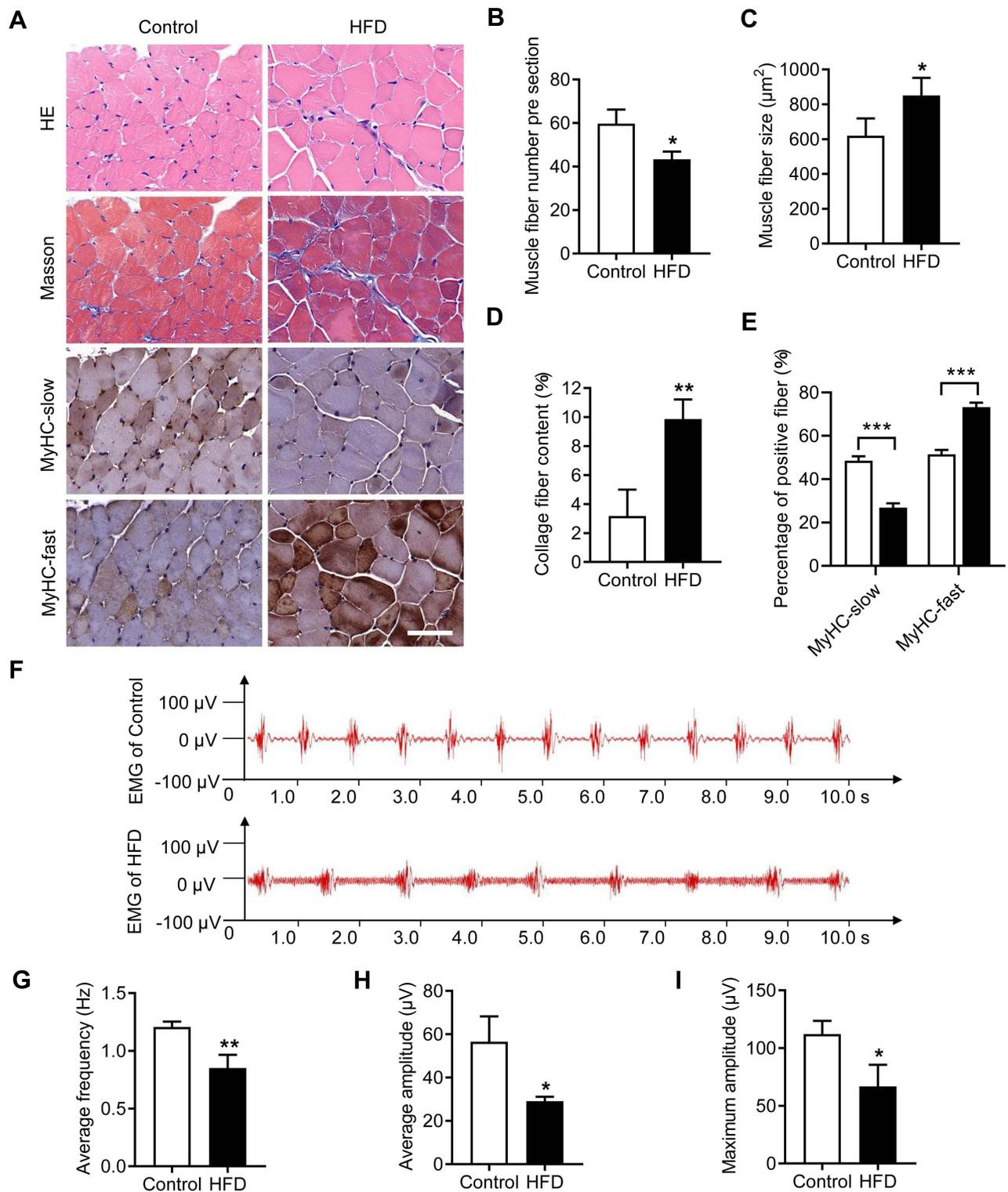
## Effects of HFD on Genioglossus Structure and Activity

The genioglossus is the major upper airway muscle and plays an important role in the maintenance of upper airway patency. The structure of the genioglossus was detected using H&E and Masson's trichrome staining and MyHC-slow or MyHC-fast immunohistochemistry assays (Figure 2A). H&E-stained sections of the genioglossus showed that the diameters of the muscle fibres in the HFD group were increased, while the number of muscle fibres was significantly less than that in the control group ( $p < 0.05$ ) (Figure 2B and C). Masson's trichrome staining showed that a HFD enhanced collagen content in the genioglossus compared with the control ( $p < 0.01$ ) (Figure 2D). Then, the genioglossus was subjected to immunohistochemical staining for the determination of muscle fibre types, which showed that the numbers of slow-twitch fibres decreased and the fast-twitch fibres increased in the HFD group ( $p < 0.001$ ) (Figure 2E).

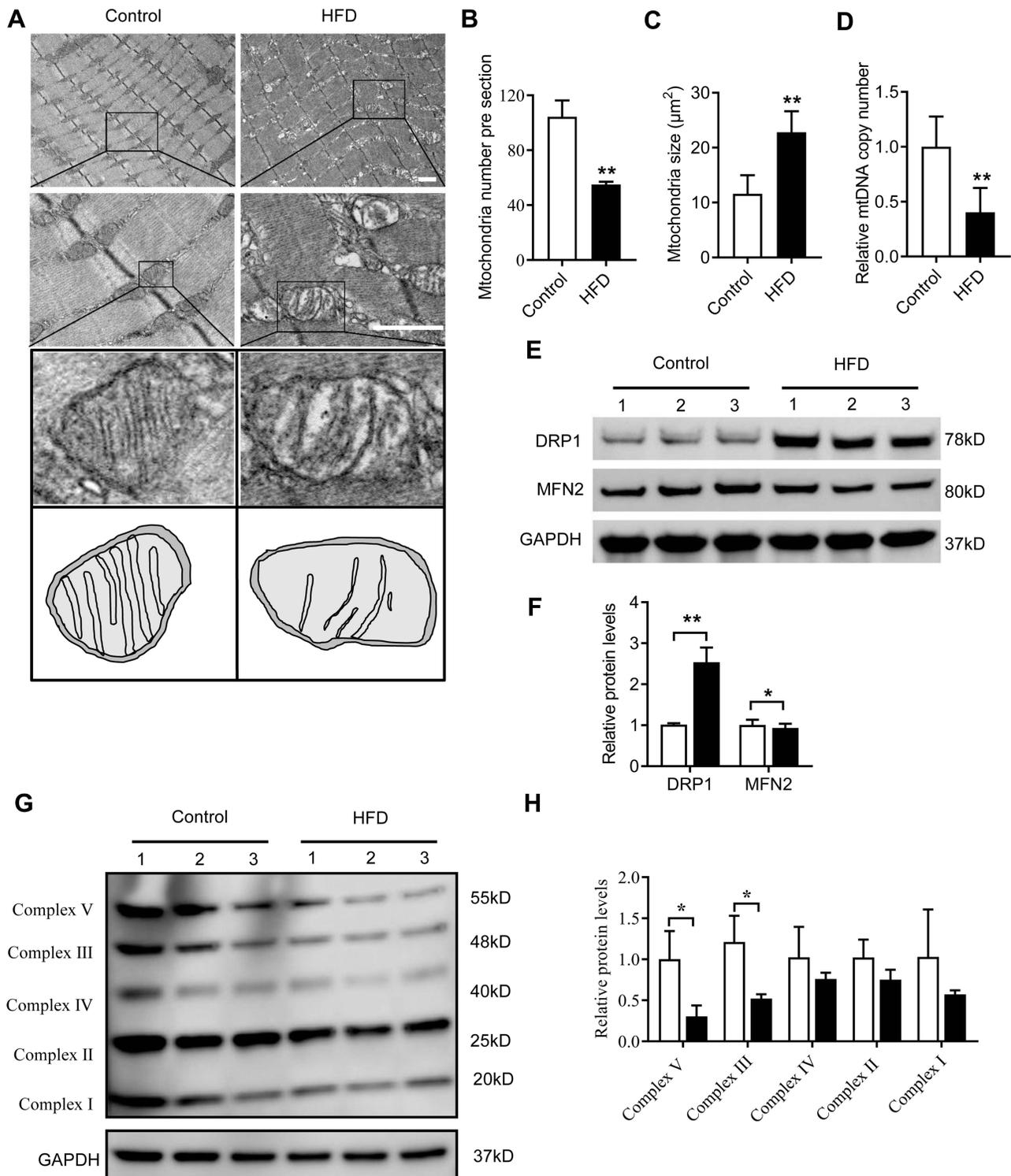
To study the influence of a HFD on the function of the genioglossus, we measured EMG activity, as depicted in Figure 2F. The average frequency of the genioglossus by EMG in the HFD group was approximately 0.85 Hz, which is significantly lower than that of the control (1.2 Hz,  $p < 0.01$ ) (Figure 2G). Moreover, both the average amplitude and maximum amplitude declined in the HFD group ( $p < 0.05$ ) (Figure 2H and I).

## Effects of HFD on Mitochondrial Function in the Genioglossus

Mitochondria are the main energy source in skeletal muscle. We investigated the effects of HFD-induced obesity on the mitochondria of the genioglossus by TEM. For the control group, the genioglossus showed regular myofibril structure and normal mitochondrial morphology, with a large number of cristae. The HFD group showed serious damage to the genioglossus ultrastructure, with vague myofibril structure and a decrease in the number of mitochondria (Figure 3A and B). The mitochondria in the HFD group were swollen, and the cristae were partially disintegrated (Figure 3A and C). We also evaluated the copy number of mtDNA in the genioglossus and the result showed that the relative mtDNA level was significantly decreased in the HFD group (Figure 3D). In addition, the expression levels of mitochondrial fission protein DRP1 and fusion protein MFN2 were measured by Western blot. As shown in Figure 3E and F, the basal expression level of DRP1 was elevated, and the expression of MFN2 was inhibited, indicating an imbalance in mitochondrial fusion and fission in the genioglossus of the HFD group. To further investigate the effect of HFD on the function of mitochondria in the genioglossus, the mitochondrial respiratory chain complexes were assessed by Western blotting. In the HFD mice, the expression of mitochondrial respiratory chain complexes III and V were downregulated (Figure 3G and H).



**Figure 2** HFD impaired muscle structure and function in the genioglossus. **(A)** Representative images of genioglossus sections from control and HFD-fed mice with H&E and Masson's trichrome staining and MyHC-slow or MyHC-fast immunohistochemical staining. **(B)** Quantification of muscle fibre number, **(C)** muscle fibre size, **(D)** collagen fibre content and **(E)** percentage of MyHC-slow and MyHC-fast per visual field. **(F)** Typical EMG in the genioglossus of control and HFD mice. **(G)** The average frequency, **(H)** average amplitude and **(I)** maximum amplitude of the genioglossus by EMG. Data are represented as the mean  $\pm$  SD. \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$  compared to the control. Scale bar is 50  $\mu\text{m}$ .



**Figure 3** HFD induced mitochondrial damage and dysfunction in the genioglossus. **(A)** Mitochondrial morphology and size in the genioglossus of control and HFD-fed mice determined by TEM. **(B)** Quantification of mitochondrial number and **(C)** mitochondrial size per visual field. **(D)** Relative mtDNA copy number in the genioglossus of control and HFD mice. **(E)** Protein expression of DRP1 and MFN2 measured by Western blot analysis in the genioglossus of control and HFD mice **(F)**. **(G and H)** Protein expression of mitochondrial respiratory chain complexes I–V measured by Western blot analysis in the genioglossus of control and HFD mice. Data are represented as the mean  $\pm$  SD. \* $p < 0.05$ , \*\* $p < 0.01$  compared to the control. Scale bars are 1  $\mu$ m.

## Effects of HFD on Oxidative Stress and Apoptosis in the Genioglossus

To study the effects of a HFD on oxidative stress in the genioglossus, the protein levels of 4-HNE and 8-OHdG, markers of oxidative stress-induced lipid peroxidation and DNA damage, were detected using immunohistochemical staining. The results showed that the numbers of 4-HNE-positive fibres (Figure 4A and B) and 8-OHdG-positive fibres (Figure 4C and D) were substantially elevated in the genioglossus of the HFD group, which indicated that the level of oxidative stress was elevated.

To investigate the effects of a HFD on apoptosis in the genioglossus, TUNEL staining was performed, which labelled the DNA fragmentation of apoptotic cells. The results showed that the number of TUNEL-positive cells in the HFD group significantly increased compared with those in the control group ( $p < 0.001$ ) (Figure 4E and F). Moreover, we also analysed the expression of the mitochondria-related apoptotic proteins Cyt-c, Bax (a proapoptotic protein) and Bcl-2 (an antiapoptotic protein) in the genioglossus by Western blotting. Compared with control mice, the protein level of cytoplasm Cyt-c (cyto Cyt-c) was increased and mitochondria Cyt-c (mito Cyt-c) was decreased in genioglossus of the HFD group, indicating that Cyt-c was released from the mitochondria into cytoplasm. A HFD also induced an elevated level of protein Bax/Bcl-2 (Figure 4G and H).

## Effects of PA on the Viability and Apoptosis in C2C12 Myoblasts

Myoblast proliferation is essential for skeletal muscle renewal and regeneration. To investigate the effects of PA on myoblast proliferation, C2C12 myoblasts were treated with different doses of PA for 6, 12, 24 and 48 hours, and CCK-8 assays were performed. As shown in Figure 5A, PA at 100  $\mu\text{M}$  did not affect the viability of C2C12 myoblasts. However, at 300  $\mu\text{M}$ , the cell morphologies changed and the percent of surviving cells was reduced to 62.31% and 54.48% that of untreated cells at 24 h and 48 h, respectively ( $p < 0.001$ ). More importantly, PA at 500  $\mu\text{M}$  exhibited a dramatic cytotoxic effect on C2C12 myoblasts, with a large portion of round and floating cells in the medium (Figure 5C), and the percent of surviving cells was only 23.57% and 11.93% that of untreated cells at 24 h and 48 h, respectively ( $p < 0.001$ ). The cytotoxic effects of PA on C2C12 myoblasts were also measured by an LDH assay. After treatment with PA at

300  $\mu\text{M}$  and 500  $\mu\text{M}$  for 24 h, the LDH leakage in the medium was elevated, exhibiting concentration-dependent cytotoxicity ( $p < 0.001$ ) (Figure 5B).

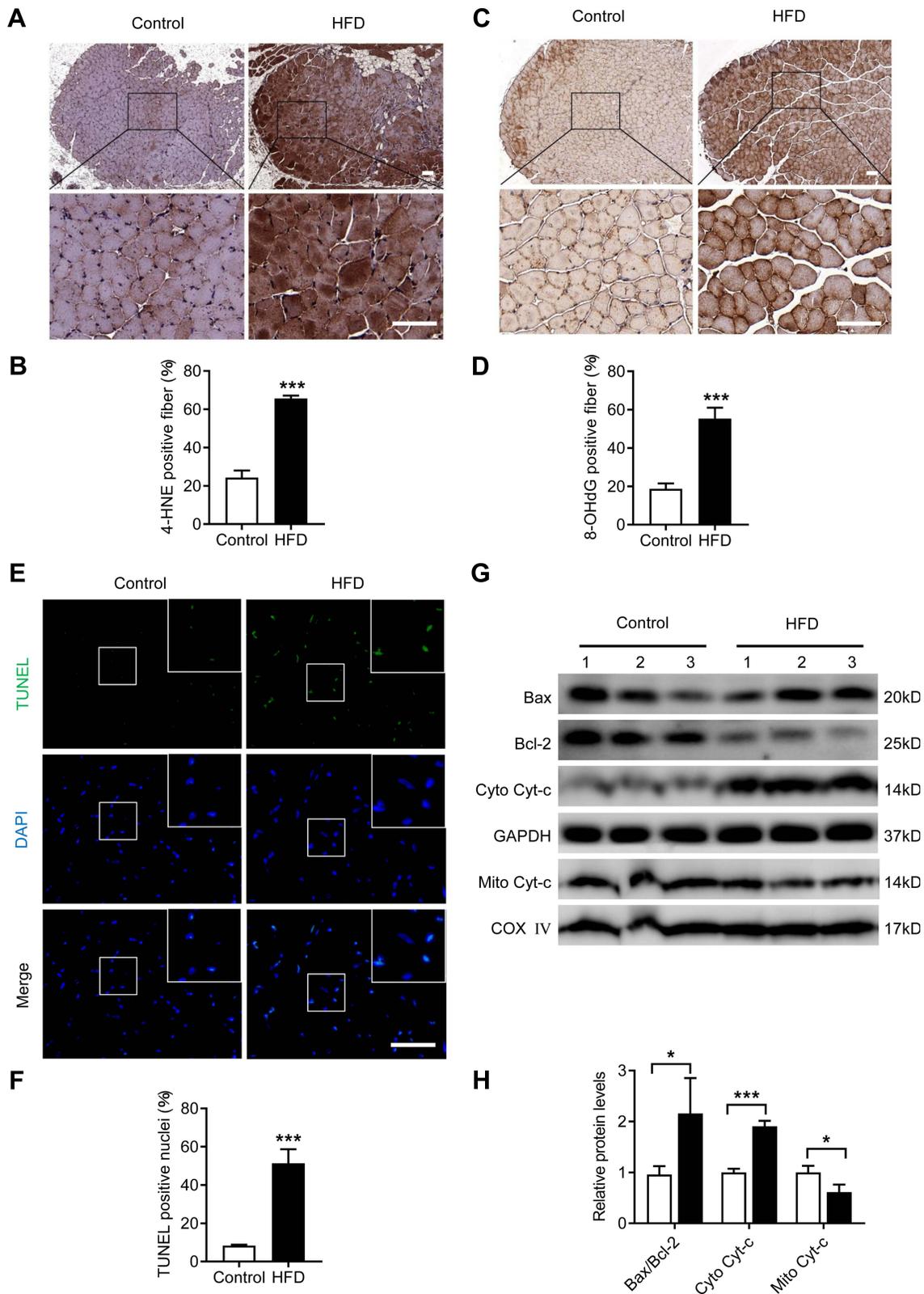
To further investigate the effects of a HFD on myoblast damage, apoptosis in C2C12 myoblasts was detected using Annexin V-FITC/PI staining and flow cytometry. Consistent with the findings of the CCK-8 and LDH assays, the results indicated that the number of apoptotic cells was significantly increased after treatment with PA at 300  $\mu\text{M}$  and 500  $\mu\text{M}$  for 24 h ( $p < 0.001$ ), and no changes were found with PA treatment at 100  $\mu\text{M}$  (Figure 5D and E).

## Effects of PA on the Myogenic Differentiation of C2C12 Myoblasts

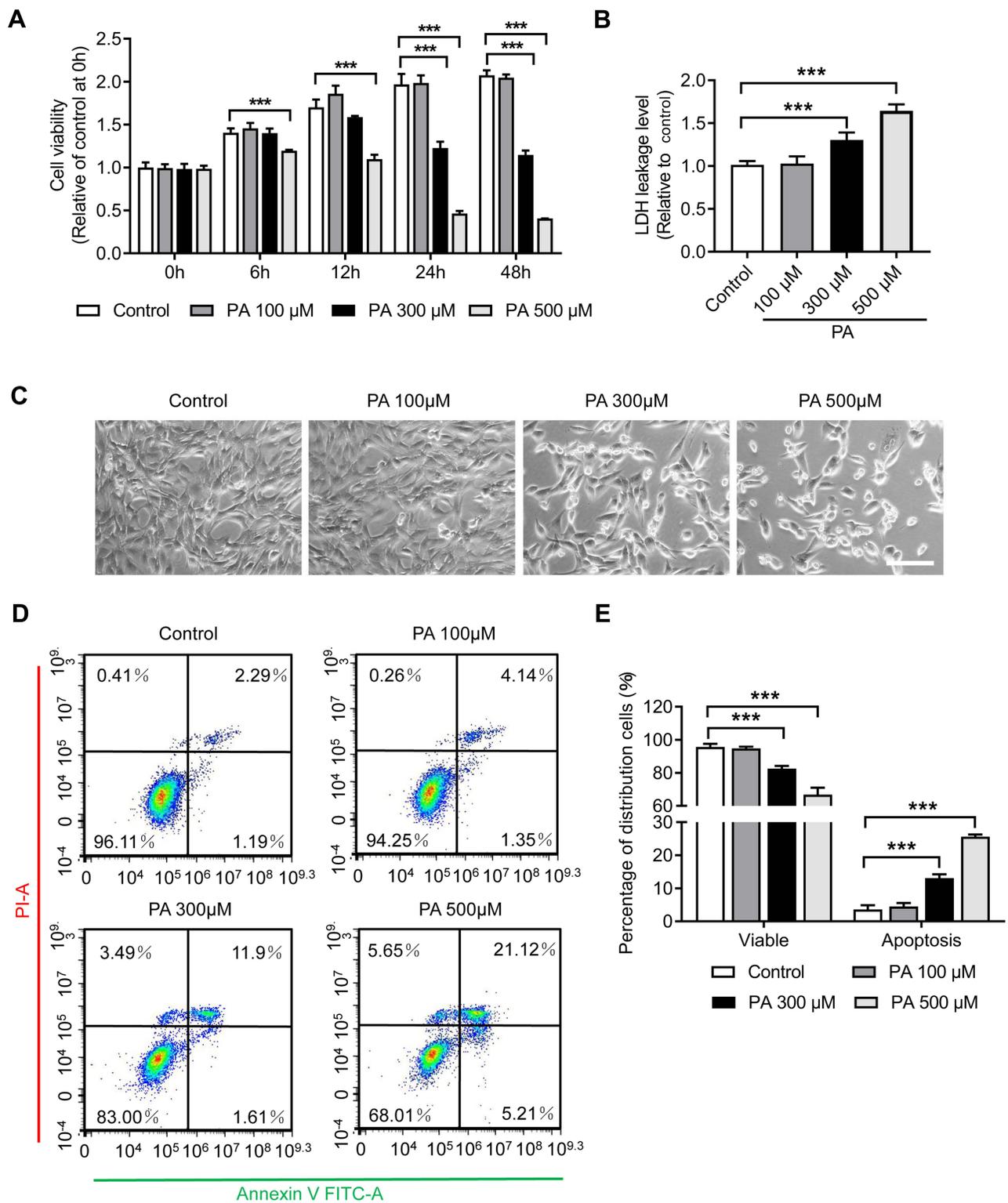
Myogenic differentiation is another important characteristic of myoblasts and plays a key role in the regeneration of skeletal muscles. To investigate whether PA affects myogenic differentiation, C2C12 myoblasts were incubated with different concentrations of PA during cellular differentiation for 2 days. The results of myotube immunofluorescence staining demonstrated that treatment with PA at 300  $\mu\text{M}$  and 500  $\mu\text{M}$  significantly prevented C2C12 differentiation, with the number of myotubes decreasing and the fusion of the myotubes becoming disrupted ( $p < 0.01$ ) (Figure 6A and B). Moreover, the relative protein levels of the key myogenic differentiation markers MyoD1, MyoG and MyHC were all downregulated significantly following treatment with 300  $\mu\text{M}$  PA (Figure 6C and D).

## Effects of PA on Mitochondrial Function in C2C12 Myotubes

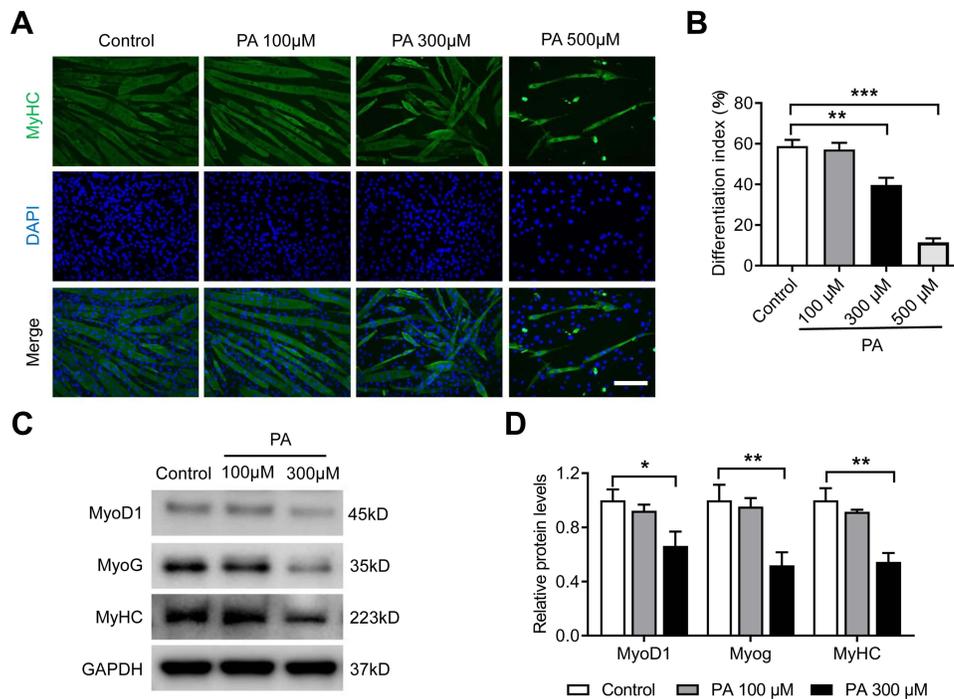
We evaluated the effects of PA on mitochondrial morphology in C2C12 myotubes using MitoTracker green. As shown in Figure 7A and C, the green fluorescence intensity of MitoTracker was markedly decreased after treatment with PA at 300  $\mu\text{M}$ , indicating decreased mitochondrial mass. Then, the expression of DRP1 and MFN2 was measured by Western blot. Consistent with the in vivo findings, the expression level of DRP1 was significantly elevated after 300  $\mu\text{M}$  PA treatment ( $p < 0.01$ ) (Figure 7E and F). Moreover, the MMP was measured with JC-1 kits, which reflect the function of mitochondrial energy metabolism. For the untreated myotubes, the red fluorescence of JC-1 aggregates represented potential-dependence in the mitochondria to produce adenosine triphosphate (ATP). Conversely, after treatment with 300  $\mu\text{M}$  PA, the green fluorescence of the JC-1 monomer was detected, and the ratio of red/green



**Figure 4** HFD led to oxidative stress and apoptosis in the genioglossus. **(A)** Representative images of genioglossus sections from control and HFD mice after 4-HNE immunohistochemical staining. **(B)** Quantification of the percentage of 4-HNE-positive fibres per visual field. **(C)** Representative images of genioglossus sections from control and HFD mice after 8-OHdG immunohistochemical staining. **(D)** Quantification of the percentage of 8-OHdG -positive fibres per visual field. **(E)** Representative images of TUNEL staining in the genioglossus. **(F)** Percentage of TUNEL-positive nuclei per visual field. **(G and H)** Protein expression of Bax, Bcl-2, Cyto Cyt-c and Mito Cyt-c measured by Western blot analysis in the genioglossus of control and HFD mice. Data are represented as the mean  $\pm$  SD. \* $p < 0.05$ , \*\*\* $p < 0.001$  compared to the control. Scale bars are 50  $\mu$ m.



**Figure 5** PA inhibited the proliferation and viability of C2C12 myoblasts. **(A)** The viability of C2C12 myoblasts was measured via CCK-8 assay after PA treatment. **(B)** LDH release in the medium of C2C12 myoblasts after PA treatment for 24 hours. **(C)** Optical micrographs of C2C12 myoblast morphology and quantity changes after PA treatment for 24 hours. **(D)** Apoptosis of C2C12 myoblasts was analysed using Annexin V-FITC/PI staining by flow cytometry after PA treatment for 24 hours. **(E)** Quantification of the percentage of viable (Annexin V-FITC negative and PI negative) cells and apoptotic (Annexin V-FITC positive) cells. Data are represented as the mean  $\pm$  SD. \*\*\* $p < 0.001$  compared to the control. Scale bars are 100  $\mu$ m.



**Figure 6** PA inhibited myogenic differentiation of C2C12 myoblasts. C2C12 myoblasts were treated with different concentrations of PA for 2 days during myogenic differentiation. **(A)** The differentiation of C2C12 myoblasts was assessed using immunofluorescence staining with MyHC (green) and DAPI (blue). **(B)** The differentiation index is presented as the percentage of MyHC-positive nuclei to the total number of nuclei. **(C and D)** Protein expression of MyoD1, MyoG and MyHC measured by Western blot analysis. Data are represented as the mean  $\pm$  SD. \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$  compared to the control. Scale bars are 50  $\mu$ m.

fluorescence intensity significantly decreased, which indicated mitochondrial membrane depolarization (Figure 7B and D). In addition, we observed a decrease in the expression of mitochondrial respiratory chain complexes III and V at 300  $\mu$ M PA treatment ( $p < 0.01$ ) (Figure 7G and H). Together, these findings suggest that PA can lead to mitochondrial damage and impair mitochondrial respiratory function.

## Effects of PA on Mitochondrial ROS and Apoptosis in C2C12 Myotubes

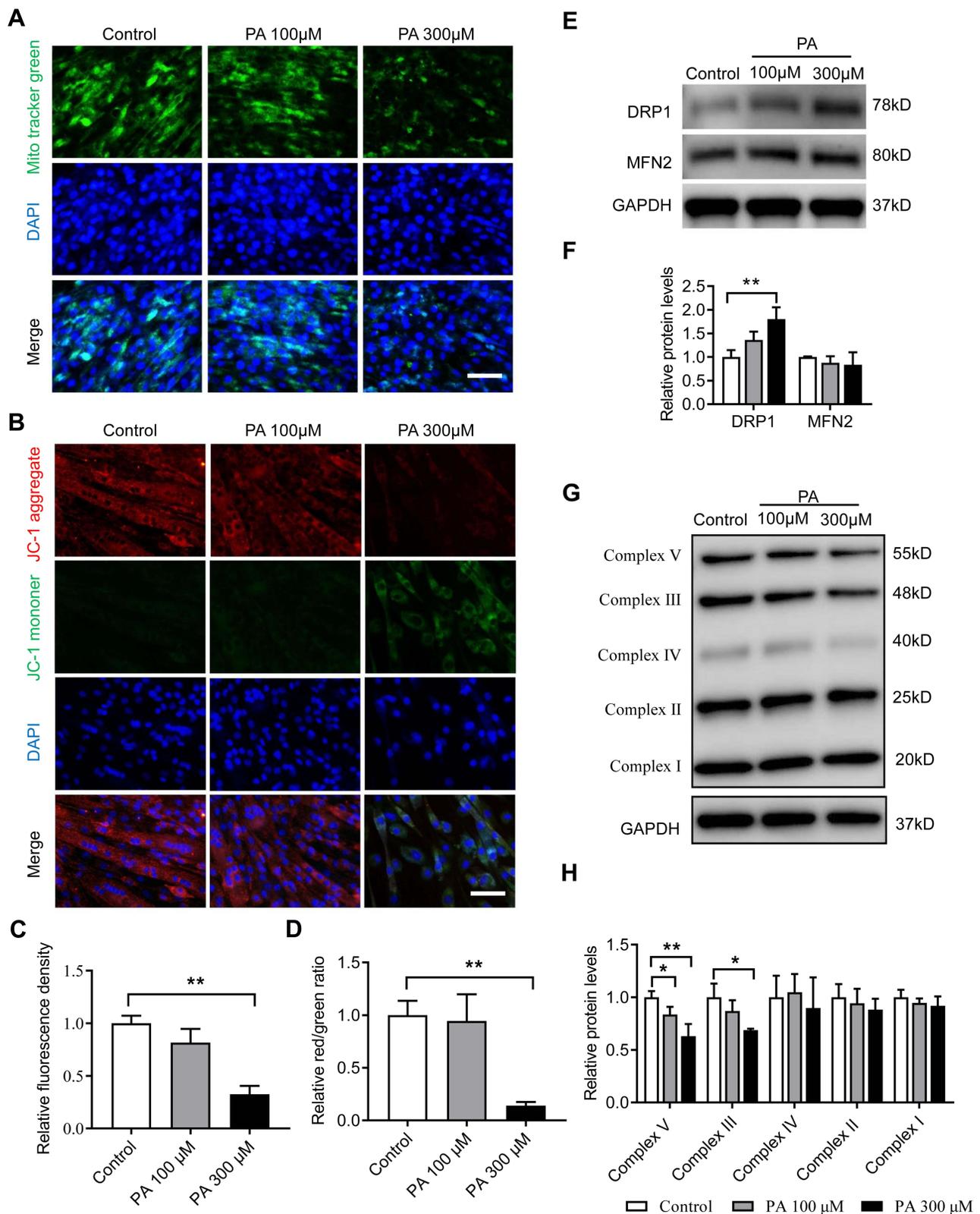
Mitochondrial dysfunction can lead to ROS overproduction, causing damage to cells and tissues.<sup>26</sup> To evaluate the effects of PA on mitochondrial ROS levels, MitoSOX Red (a mitochondrial superoxide indicator) was used for immunofluorescence staining. PA at 300  $\mu$ M significantly increased mitochondrial ROS accumulation in C2C12 myotubes ( $p < 0.001$ ) (Figure 8A and B). ROS are known to activate apoptotic proteins and promote cell apoptosis. After treatment with 300  $\mu$ M PA, the number of TUNEL-positive cells was significantly increased, and the expression of the mitochondria-related apoptotic proteins Cyt-c, Bax/Bcl-2 and cleaved caspase-3 was notably elevated in C2C12 myotubes (Figure 8C–F). SS-31 is reported to be an antioxidant targeting mitochondria.<sup>32,33</sup>

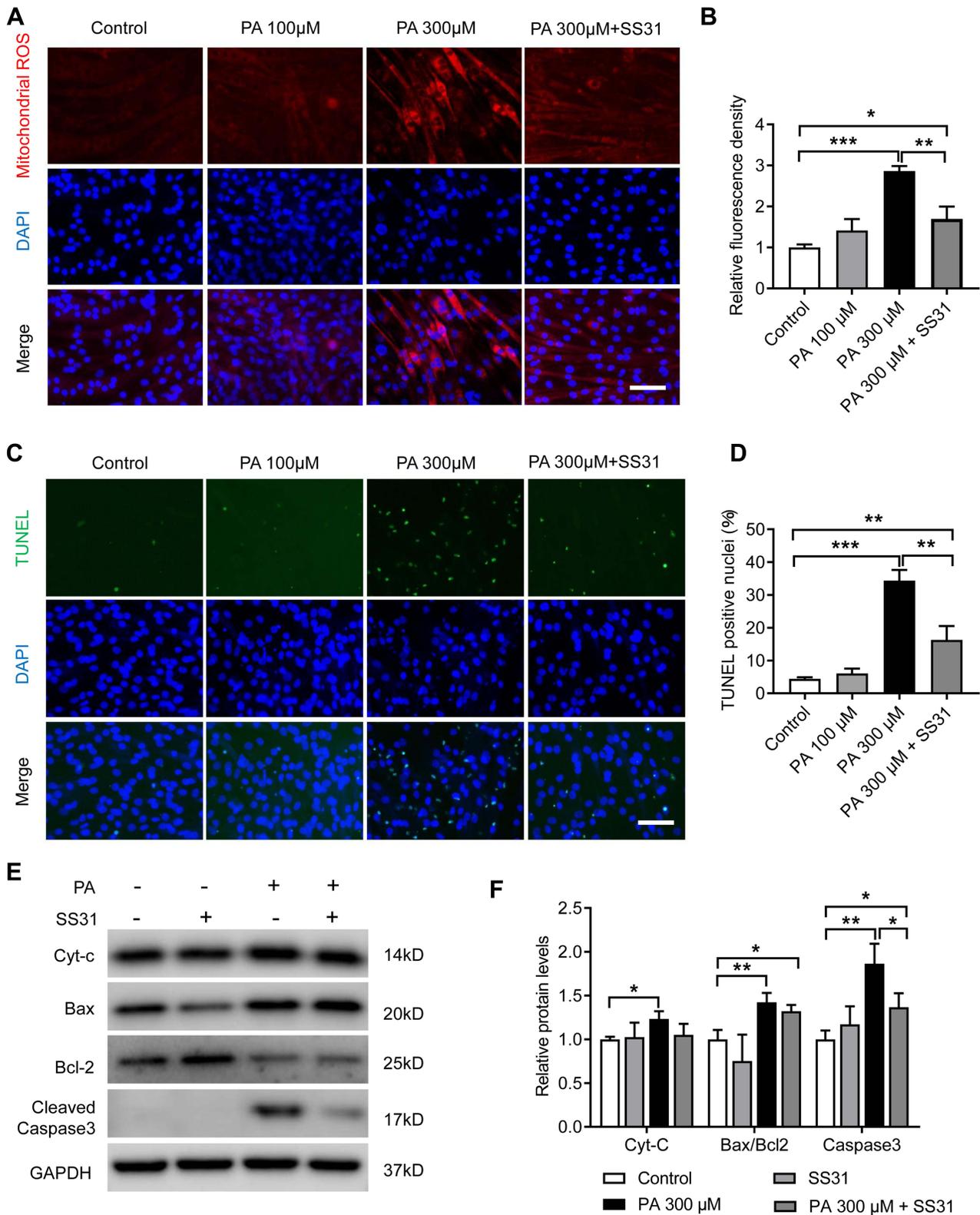
SS-31 treatment significantly reduced the mitochondrial ROS accumulation in C2C12 myotubes induced by PA (Figure 8A and B). In addition, compared with the 300  $\mu$ M PA-treated group, SS-31 treatment reduced the number of TUNEL-positive cells (Figure 8C and D) and down-regulated the protein level of cleaved caspase-3 (Figure 8E and F), indicating SS-31 could partly alleviate the apoptotic in C2C12 myotubes caused by PA.

## Discussion

The dysfunction of upper airway dilator muscles, especially the genioglossus, is thought to be the main pathogenesis of OSA. The HFD-induced obese mice in our study showed that genioglossus injury was accompanied by mitochondrial dysfunction, oxidative stress and apoptosis, which might be an important mechanism of OSA co-occurring with obesity.

In the present study, mice exhibited a significant increase in body weight and plasma levels of FFAs after 16 weeks of HFD administration. Skeletal muscles play an important role in lipid metabolism, while the excessive accumulation of lipids can impair oxidative capacities, promote lipotoxicity and lead to changes in the structure and function of skeletal muscles.<sup>34</sup> As predicted, we





**Figure 8** PA increased mitochondrial ROS and apoptosis in C2C12 myotubes. C2C12 myotubes were treated with PA or SS-31 for 2 days. **(A and B)** Fluorescence images and analyses of mitochondrial ROS with MitoSOX red in C2C12 myotubes. **(C)** Representative images of TUNEL staining in C2C12 myotubes. **(D)** Quantification of the percentage of TUNEL-positive nuclei per visual field. **(E and F)** Protein expression of Cyt-c, Bax, Bcl-2 and cleaved caspase-3 measured by Western blot analysis in C2C12 myotubes. Data are represented as the mean ± SD. \**p* < 0.05, \*\**p* < 0.01, \*\*\**p* < 0.001. Scale bars are 50 µm.

discovered changes in genioglossus muscle fibres of HFD mice, with a decrease in the number of muscle fibres and an increase in the collagen content.

In addition, the muscle fibre type, with a dynamic and plastic structure, is of importance to muscle development, metabolism and functional adaptations. Slow-twitch fibres have more mitochondria and exhibit high oxidative capacity and fatigue resistance. In contrast, fast-twitch fibres have lower mitochondria levels and oxidative capacity. Studies have demonstrated that obese individuals have a decreased proportion of slow-twitch fibres and an increased proportion of fast-twitch fibres in skeletal muscles.<sup>35,36</sup> The genioglossus in patients with OSA have also been reported to have a higher proportion of fast-twitch fibres and be more vulnerable to fatigue.<sup>37</sup> In this study, we confirmed that a HFD caused significant changes in fibre types in the genioglossus, with the percentage of slow-twitch fibres declining and the percentage of fast-twitch fibres increasing. Changes in muscle fibre structure could lead to impairment of muscle function. Consistently, we found that the EMG activity of the genioglossus was significantly diminished in HFD-induced obese mice.

Mitochondria are the main energy source in skeletal muscle and play a critical role in maintaining contractility.<sup>38</sup> The muscle fibre structure and fibre type have been reported to be significantly associated with mitochondrial dysfunction.<sup>39</sup> Overaccumulation of FFAs could induce excessive ceramide levels,<sup>40</sup> which were reported to ultimately cause mitochondrial dysfunction and lipotoxicity.<sup>41,42</sup> In this study, we found that HFD-induced obesity led to mitochondrial damage; as the number of mitochondria decreased, they became swollen, and the cristae ruptured.

Mitochondria in skeletal muscle are highly dynamic, and mitochondrial biogenesis, fusion, fission and mitophagy are all involved in the regulation of mitochondrial morphology and function. Recent studies have shown that disorder in the balance between mitochondrial fission and fusion is implicated in mitochondrial damage and dysfunction in skeletal muscle in obesity.<sup>43,44</sup> We detected a decrease in the mitochondrial fusion protein MFN2 and an increase in the fission protein DRP1 in the genioglossus of HFD-induced obese mice, indicating that obesity impaired mitochondrial fusion and induced excessive mitochondrial fission in the genioglossus. Our *in vitro* data support this conclusion. PA-treated C2C12 myotubes exhibited a decrease in mitochondrial mass and an increase in protein DRP1 expression, coupled with decreases in the MMP and mitochondrial oxidative phosphorylation (OXPHOS) complexes III and V, which are consistent with the observation *in vivo*. Mitochondrial dysfunction has been

suggested to be one of the key processes during obesity that leads to skeletal muscle injury.<sup>45</sup> Considering the importance of the mitochondria on oxidative metabolism and energy generation, we speculated that mitochondrial damage and dysfunction could ultimately lead to decreased oxidative capacities and fatigue resistance in the genioglossus.

Mitochondrial dysfunction can trigger apoptosis by regulating apoptosis-related genes, such as those in the Bcl-2 family.<sup>46</sup> In the genioglossus, more TUNEL-positive apoptotic cells were observed in HFD-induced obese mice. Moreover, the expression levels of pro-apoptosis proteins (Cyt-c and cleaved caspase-3) were upregulated, while the expression of an apoptosis-inhibiting protein (Bcl-2) was downregulated in the genioglossus of obese mice. In support of these findings, our *in vitro* study demonstrated that high levels of PA significantly inhibited cellular proliferation and myogenic differentiation in C2C12 myoblasts. Moreover, PA exposure stimulated cell apoptosis in both C2C12 myoblasts and myotubes.

Mitochondrial impairment contributes to excessive ROS generation and oxidative stress, which is an important mechanism of tissue injury and apoptosis.<sup>47</sup> Oxidative stress is a critical factor for skeletal muscle injury in obesity.<sup>48</sup> ROS overproduction was also found in the genioglossus of rats with chronic intermittent hypoxia.<sup>49</sup> In this study, we indeed detected the oxidative damage in the genioglossus of HFD-induced obese mice, manifested by the dramatic upregulation of 4-HNE-positive staining muscle fibres and 8-OHdG-positive fibres. Consistent with the *in vivo* findings, we also found increased production of mitochondrial ROS in C2C12 myotubes after PA treatment. Furthermore, SS-31, an antioxidant targeting mitochondria, reduced the accumulation of mitochondrial ROS and ultimately alleviated the apoptotic in C2C12 myotubes caused by PA.

In conclusion, our data indicate that a HFD could decrease EMG activity and muscle fibre mass, cause marked changes in muscle fibre type, lead to ultrastructural damage and increase apoptosis in the genioglossus. Most impressively, this study identifies that genioglossus injury might be mainly mediated by mitochondrial disturbance and the accompanying oxidative stress in HFD-induced obese mice, which may provide potential therapeutic targets for OSA co-occurrence with obesity.

## Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation,

or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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## Disclosure

The authors report no conflicts of interest in this work.

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