Abstract. Mice that lacked manganese-superoxide dismutase (Mn-SOD) activity exhibited the typical pathology of dilated cardiomyopathy (DCM). The aim of the present study was to investigate the effect of supplementation with omega-3 polyunsaturated fatty acids (n-3 PUFA) on heart function and oxidative stress biomarkers in mice with DCM. In the present study, heart/muscle-specific Mn-SOD-deficient mice (H/M-Sod2-/-) were treated with n-3 PUFA (30 mg/kg/day) for 10 weeks, and the reactive oxygen species (ROS) production in their heart mitochondria and cardiac function was subsequently assessed. n-3 PUFA treatment diminished ROS production and suppressed the progression of cardiac dysfunction. Furthermore, n-3 PUFA treatment effectively reversed the cardiac dysfunction and dilatation observed in symptomatic H/M-Sod2-/- mice. Notably, n-3 PUFA treatment ameliorated a molecular defect in connexin 43. Hematoxylin-eosin staining indicated that the phenotype of DCM was also ameliorated following n-3 PUFA treatment. Furthermore, echocardiography demonstrated that cardiac function was significantly improved in the mice treated with n-3 PUFA (P<0.05). Meanwhile, pre-treatment with n-3 PUFA significantly decreased cardiomyocyte apoptosis (P<0.001). In conclusion, n-3 PUFA treatment is able to prevent murine DCM, primarily by reducing ROS production and improving myocardial apoptosis. Therefore, the impairment of ROS production is proposed as a potential therapy for DCM.

Introduction

As a progressive heart muscle disease, dilated cardiomyopathy (DCM) is characterized by a dilated ventricular chamber, including impaired contraction of the left or both ventricles (1,2). DCM typically leads to congestive heart failure, thereby enhancing morbidity and mortality among patients (3,4). Several clinical studies have indicated that reactive oxygen species (ROS) are important in the pathogenesis of cardiovascular diseases, including DCM (5-7). Furthermore, it has been suggested that an imbalance between ROS production and antioxidant defenses may result in oxidative stress-related disorders, including DCM (8,9). Epidemiological studies show that patients with cardiovascular diseases can be prevented from ROS injury through antioxidant reagents (10). Additionally, several potential sources of ROS have been proposed, including mitochondrial respiratory chain enzymes, xanthine oxidase, nicotinamide adenine dinucleotide phosphate (NADPH) oxidase and nitric oxide synthase (11). Abnormal ROS production may further result in aberrant morphological or functional changes, which are demonstrated as cardiac dysfunction (12).

Omega-3 polyunsaturated fatty acid (n-3 PUFA), a free radical scavenger, has long been suggested to be effective for combating oxidative stress (13-15). However, the specific mechanism through which n-3 PUFA protects the heart from DCM remains to be elucidated. Therefore, the present study aimed to determine the potential role of n-3 PUFA on DCM.

It has previously been indicated that the typical pathology of DCM is exhibited by mice lacking manganese-superoxide dismutase (Mn-SOD) activity (16,17). Therefore, the present study used such mice as a DCM model to explore the effect of n-3 PUFA on the parameters of heart function and oxidative stress biomarkers in DCM mice.

Materials and methods

In vivo study protocols. A total of 8 male heart/muscle-specific Mn-SOD-deficient (H/M-Sod2-/-) mice (age, 8 weeks; weight, 22.5±3.1 g) were purchased from The Fourth Affiliated Hospital of Harbin Medical University (Harbin, China). The mice were...
kept in a temperature-(20-24°C) and humidity-controlled (45-55%) environment, under a 12-h light/dark cycle, with free access to food and water. H/M-Sod2+/− mice were fed with 3 mg/kg/day n-3 PUFA (n=4) (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) or vehicle (n=4) (lauric acid, Sigma-Aldrich; Merck KGaA) (18) for 10 weeks, and cardiac function was evaluated by echocardiography. Mice were subsequently sacrificed, hearts were harvested and the proteins extracted. The present study was approved by the Zhejiang Province Hospital of Integrated Traditional Chinese and Western Medicine (Hangzhou, China).

Echocardiography. Two-dimensional (2D) echocardiography was performed with certain modifications according to previously reported methods (19). On the day of evaluation, sodium pentobarbital (50 mg/kg; P3761, Sigma-Aldrich; Merck KGaA) was administered by intraperitoneal injection to anesthetize the mice. After shaving the chest, 2D echocardiography was conducted with an echocardiographic system (model SSD-900; Hitachi-Aloka Medical, Ltd., Tokyo, Japan) and 7.5 MHz probe (UST-987-7.5; Hitachi-Aloka Medical, Ltd.). Furthermore, the fraction shortening (FS), ejection fraction (EF), left ventricular internal diameter (LVID) during systole and diastole, and endystolic and end-diastolic volumes were calculated using Vevo Vasc Analysis software (version 2.2.3; Fujifilm VisualSonics, Inc., Toronto, Canada) as previously described (20).

Primary cardiomyocyte culture. Primary cardiomyocytes from mouse neonatal hearts were isolated as previously described (21). In brief, hearts were isolated and digested with collagenase type II solution (Worthington Biochemical Corporation, Lakewood, NJ, USA). Following digestion, the cells were plated in a 25-cm² culture flask for 2 h at 37°C in Dulbecco's modified Eagle's medium (DMEM; GE Healthcare Life Sciences, Logan, UT, USA) supplemented with 1% (v/v) fetal bovine serum (Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA) to collect cardiomyocytes. Attached cells were considered as non-myocytes and discarded, and the unattached cells were primarily cardiomyocytes.

Histological studies. For the histological analysis, heart tissues were fixed at room temperature for 10 min in 10% buffered formalin. Fixed tissues were subsequently dehydrated, embedded in paraffin, sectioned into 4-µm slices and stained with hematoxylin and eosin (H&E). Furthermore, Masson's trichrome was used to stain myocardial sections in order to evaluate the cardiomyocyte diameter and degree of fibrosis. Images were captured with a Pixera Pro600EX camera and a VANOX-S microscope (Olympus Corporation, Tokyo, Japan). Additionally, the fibrotic area and cardiomyocyte diameter (>30 cells) were quantified with Qwin Plus V3 (Leica Microsystems, Inc., Buffalo Grove, IL, USA), and the collagen volume percentage was calculated as the mean of 5 fields for each animal.

Hoechst 33258 staining. Primary cardiomyocytes were cultured in 6-well tissue culture plates (1x10⁶ cells per well). The cells were incubated at 37°C in serum-free DMEM for 16 h at 70-80% confluence. The cells were then washed three times with cold PBS and fixed with 4% formaldehyde (Zhongshan Technology Co., Ltd., Zhongshan, China) in PBS for 20 min at room temperature. Next, the cells were washed three times with cold PBS and stained with Hoechst 33258 (10 µg/ml; 50 µl/slide; Sigma-Aldrich; Merck KGaA) at room temperature for 5 min. Following staining, cold PBS was used to further rinse the cells, and were then examined under a fluorescence microscope.

Apoptosis assay. In order to detect the effects of n-3 PUFA on cell apoptosis, primary cardiomyocytes isolated from mice were washed with cold PBS three times. Next, flow cytometry was used to determine cell apoptosis with an Annexin-V fluorescein isothiocyanate-propidium iodide (FITC-PI) apoptosis kit (Invitrogen, Carlsbad, CA, USA). In summary, the cells (1x10⁶) were washed with 1X PBS three times and suspended at 2-3x10⁶ cells/ml in X1 Annexin-V binding buffer [10 mM HEPES/NaOH, (pH 7.4), 140 mM NaCl, 2.5 mM CaCl₂]. Annexin-V FITC and PI buffer were then added to the cells, which were then incubated at room temperature for 15 min in the dark. The cells that did not undergo any treatment were used as an internal control. Following incubation, the cells were filtered using a filter screen and analyzed using a flow cytometer within 1 h of staining. Cell apoptosis was analyzed using BD CellQuest Pro™ Analysis Software (BD Biosciences, San Jose, CA, USA).

Determination and quantification of ROS. Primary cardiomyocytes isolated from the mice were cultured on slides in a 6-well chamber at 60% confluence at 37°C. Two days later, the slides were washed with cold PBS three times. Additionally, the slides were treated with 5 µM dihydroethidium (DHE; Vigorous Biotechnology Beijing Co., Ltd., Beijing, China) in serum-free DMEM F-12 medium (GE Healthcare Life Sciences) for 30 min at 37°C in darkness. Furthermore, the cells were fixed in 4% paraformaldehyde for 30 min at room temperature. The slides were then washed with cold PBS three times and mounted. Finally, immunofluorescence images were captured by fluorescence microscopy, and in order to quantify the intracellular ROS, the relative fluorescence intensities were analyzed using flow cytometry in the primary cardiomyocytes. Briefly, 1x10⁶ cells were centrifuged (200 x g) for 10 min at room temperature and the supernatant was discarded. The cell pellet was resuspended in 1 ml of PBS at room temperature. A 2 mM solution of H₂DCFDA (Invitrogen; Thermo Fisher Scientific, Inc.) was freshly prepared in ethanol and 5 µl was added to the cell suspension (final concentration 10 µM) and incubated at 37°C for 20 min. Cells were centrifuged at 200 x g for 5 min at room temperature, the supernatant was discarded, and the cell pellet was resuspended in 500 µl of PBS. Flow cytometric analysis was performed in duplicate using a flow cytometer and acquired data were analyzed using WinMDI v2.8 software (BD Biosciences).

Protein extraction, western blotting and antibodies. Cellular proteins were extracted using RIPA buffer [50 mM Tris/HCl, (pH 7.4), 150 mM NaCl 1% (v/v) NP-40, 0.1% (w/v) SDS; Beijing Solarbio Science & Technology Co., Ltd., Beijing, China] with 0.3% (v/v) protease inhibitor (Sigma-Aldrich; Merck KGaA), 1% (v/v) phenylmethane sulfonyl fluoride (Beijing Solarbio
Science & Technology Co., Ltd.) and 0.1% (w/v) phosphorylated proteinase inhibitor (Sigma-Aldrich, Inc.; Merck KGaA) for 15 min at room temperature. The lysates were collected for the total protein following centrifugation at 10,000 x g at 4˚C for 15 min. Furthermore, the bicinechonic acid protein assay kit (Pierce; Thermo Fisher Scientific, Inc.) was used to determine the protein concentration. A total of 15 µg protein per lane was separated by 10% SDS-PAGE and transferred onto a polyvinylidene difluoride membrane. After blocking with 8% milk in PBST for 2 h at room temperature, the membranes were then incubated with primary antibodies against GAPDH (cat. no. 2118; 1:5,000), cleaved-caspase 3 (cat. no. 9664; 1:1,000), p-JNK (cat. no. 4668; 1:1,000), JNK (cat. no. 9252; 1:1,000), p-NF-κB (cat. no. 3033; 1:1,000), NF-κB (cat. no. 8801; 1:1,000) (all Cell Signaling Technology, Inc., Danvers, MA, USA) and NOX4 (cat. no. ab109225; 1:1,000; Abcam, Cambridge, UK) overnight at 4˚C. Next, the membranes were washed with TBST three times. Additionally, they were incubated with horseradish peroxidase (HRP)-conjugated goat anti-rabbit or HRP-conjugated mouse anti-goat IgG (both 1:5,000; ZB-2301, Zhongshan Golden Bridge Biotechnology Co., Ltd., Beijing, China) for 2 h at room temperature and then washed three times with TBST. Enhanced chemiluminescence (EMD Millipore, Billerica, MA, USA) was used to determine the protein concentrations according to the manufacturer's recommendations. The relative contents of protein were normalized against GAPDH. All experiments were repeated three times. ImageJ 1.43b software (National Institutes of Health, Bethesda, MD, USA) was used for densitometry analysis.

**Determination of protein carbonylation and ATP content.**

The nuclear and mitochondrial fractions of the heart were isolated in order to quantify cardiac protein carbonylation. In brief, the crude nuclear fractions were isolated from tissue homogenates at 1,000 x g for 5 min at 4˚C and washed with PBS at 4˚C. The mitochondrial fractions were separated as previously described (22). The carbonylation of mitochondrial and nuclear protein was evaluated using an Oxyblot protein oxidation detection kit (EMD Millipore) according to the manufacturer's instructions. Immunoreactive spots were visualized with enhanced chemiluminescence (GE Healthcare, Buckinghamshire, UK) and quantified with ImageJ 1.43b (National Institutes of Health, Bethesda, MD, USA). Furthermore, the ATP content was determined using an ATP assay kit (colorimetric/fluorometric) (cat. no. ab83355; Abcam), according to the manufacturer's instructions.

**Statistical analysis.** Data are presented as the mean ± standard error from three independent experiments. Statistical analysis was performed with Student's t-test. *P*<0.05 was considered to indicate a statistically significant difference.

**Results**

Cardiac dilation in hearts from H/M-Sod2−/− mice. Compared with wild-type (WT) mice, hearts from H/M-Sod2−/− mice yielded increased parasternal long-axis M-mode echocardiographic images (Fig. 1A). As shown in Fig. 1B, H&E staining revealed an evident dilation of DCM hearts from H/M-Sod2−/− mice compared with those from WT mice. Furthermore, the myocardial sections from H/M-Sod2−/− mice indicated eccentric hypertrophy and myocardial disarray compared with hearts from WT mice (Fig. 1C). These data showed the cardiac dilation phenotype in hearts from H/M-Sod2−/− mice.

**n-3 PUFA ameliorates cardiac enlargement in H/M-Sod2−/− mice.** To identify whether n-3 PUFA prevents cardiac enlargement in H/M-Sod2−/− mice, Masson's trichrome staining demonstrated that interstitial fibrosis was markedly improved after n-3 PUFA treatment, suggesting that n-3 PUFA was able to partially reverse cardiac fibrosis in the H/M-Sod2−/− mice (Fig. 2A). Furthermore, the cardiomyocyte diameter was significantly decreased in H/M-Sod2−/− mice treated with n-3 PUFA, compared with untreated mice, indicating that cardiac hypertrophy was ameliorated by n-3 PUFA treatment (Fig. 2B). Following n-3 PUFA treatment, the percentage of FS was significantly enhanced, compared with untreated mice (Fig. 2C). These data suggested that
Figure 2. n-3 PUFA prevents cardiac enlargement in H/M-Sod2−/− mice. (A) Masson's trichrome staining demonstrated that interstitial fibrosis was markedly ameliorated following n-3 PUFA treatment. (B) Cardiomyocyte diameter was significantly decreased in the H/M-Sod2−/− mice treated with n-3 PUFA, compared with untreated mice. (C) Following n-3 PUFA treatment, FS was significantly increased. n=5; *P<0.05; **P<0.01. n-3 PUFA, omega-3 polyunsaturated fatty acids; H/M-Sod2−/−, heart/muscle-specific manganese-superoxide dismutase-deficient mice; FS, fraction shortening; WT, wild-type.

Figure 3. n-3 PUFA decreases ROS production in H/M-Sod2−/− mice. (A) Dihydroethidium staining demonstrated that n-3 PUFA treatment markedly decreased ROS production in primary cardiomyocytes isolated from H/M-Sod2−/−. Protein carbonylation was significantly decreased in (B) the nuclei and (C) mitochondria of primary cardiomyocytes treated by n-3 PUFA of H/M-Sod2−/− mice. (D) n-3 PUFA treatment recovered 21% more ATP content compared with the control H/M-Sod2−/− mice. n=5; *P<0.05; **P<0.01 vs. WT. n-3 PUFA, omega-3 polyunsaturated fatty acids; ROS, reactive oxygen species; H/M-Sod2−/−, heart/muscle-specific manganese-superoxide dismutase-deficient mice; WT, wild-type.
n-3 PUFA treatment ameliorated cardiac dysfunction in H/M-Sod2−/− mice.

n-3 PUFA decreases ROS production in H/M-Sod2−/− mice. The ROS production in primary cardiomyocytes isolated from H/M-Sod2−/− and WT mice was identified. As shown in Fig. 3A, DHE staining revealed that n-3 PUFA treatment markedly decreased the ROS production in primary cardiomyocytes isolated from H/M-Sod2−/−. Subsequently, oxidative protein damage was investigated in nuclei and mitochondria. Protein carbonylation was significantly decreased in the nuclei and mitochondria of primary cardiomyocytes of H/M-Sod2−/− mice treated with n-3 PUFA (Fig. 3B and C).

Furthermore, n-3 PUFA treatment recovered 21% more ATP content compared with the untreated H/M-Sod2−/− mice (Fig. 3D). These data indicated that n-3 PUFA decreases ROS generation, protein damage and the ATP content in H/M-Sod2−/− mice.

n-3 PUFA reduces primary cardiomyocyte apoptosis. In order to investigate the potential role of n-3 PUFA on cell apoptosis, Hoechst staining and flow cytometry were applied to determine apoptosis of primary cardiomyocytes isolated from H/M-Sod2−/− mice. Following n-3 PUFA treatment, primary cardiomyocytes isolated from H/M-Sod2−/− mice after apoptosis were evidently decreased compared with the
untreated group (Fig. 4A and B). Notably, the phosphorylation levels of c-Jun N-terminal kinases (JNK), nuclear factor (NF)-κB and cleaved-caspase 3 were significantly decreased with n-3 PUFA treatment, whereas the expression of NADPH oxidase 4 (NOX4) was significantly increased (Fig. 4C).

Discussion

The present study identified that n-3 PUFA treatment was able to partially abolish cardiac enlargement and dysfunction in H/M-Sod2-/- mice and that the protective effects of n-3 PUFA predominantly originated from reduced ROS production and cardiomyocyte apoptosis. Together, these data indicate that decreased oxidative damage contributes to the reduction of cardiac enlargement in H/M-Sod2-/- mice. Knockout of the Mn-SOD gene in the heart and muscle may lead to cardiac oxidative stress, which leads to contractile dysfunction, fibrosis and myocyte damage (23). According to echocardiographic analysis, hearts from H/M-Sod2-/- mice were significantly enlarged. Furthermore, EF and FS were also found to be reduced in the hearts of H/M-Sod2-/- mice, indicating severe DCM and cardiac dysfunction of these mice. Notably, n-3 PUFA treatment was demonstrated to improve histological abnormalities in DCM hearts of H/M-Sod2-/- mice, such as fibrosis, compared with hearts from the control group. This observation is in accordance with the Masson's trichrome-stained section analysis and suggests that fibrosis is improved in DCM hearts.

ROS are widely associated with age-related diseases, such as Alzheimer's and Parkinson's disease, and heart failure (24,25). During mitochondrial respiration, small amounts of mitochondrial ROS production may be cleared by scavenging systems. In the presence of NOX4, superoxide anions were significantly decreased in dilated cardiomyopathy, suggesting the protective role of NOX4 on ROS production in HF (12). It is widely reported that abnormal ROS production and apoptosis are important in the pathology of DCM (28,29). Therefore, to prevent diabetic cardiomyopathy, it is important to simultaneously inhibit oxidative stress and apoptosis. The present study explored the effects of n-3 PUFA on cardiomyocyte apoptosis. It was observed that n-3 PUFA treatment led to a significant reduction of cell apoptosis in primary cardiomyocytes isolated from H/M-Sod2-/- mice compared with the untreated group. Accordingly, the protein level of cleaved-caspase 3 was significantly decreased following n-3 PUFA treatment. Furthermore, it was also shown that n-3 PUFA demonstrated anti-inflammatory effects since the phosphorylation levels of JNK and NF-κB were significantly decreased.

In conclusion, oxidative stress was shown to increase in the DCM model of H/M-Sod2-/- mice. Notably, the results indicate that n-3 PUFA may be used as an antioxidant to protect hearts from DCM, primarily by reducing ROS production and cardiomyocyte apoptosis. Finally, the present study may assist in the development of a novel therapy and prevention for DCM in human patients.

References


