**Original Research**

**Canthaxanthin Mitigates Cardiovascular Senescence in Vitro and in Vivo**

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**Abstract**

**Background:** The number of older people in the world is increasing year by year; studies have shown that more than 90% of cardiovascular disease occurs in the older population, indicating that aging is one of the major risks involved in the development of cardiovascular disease. Therefore, retarding the development of cardiac aging is an important strategy to prevent aging-related cardiovascular diseases. **Methods:** In the current study, we examined the anti-cardiovascular aging potential of canthaxanthin in vitro and in vivo experiments. For this, a model of cardiomyocyte senescence induced by D-galactose was established, which was used to investigate the canthaxanthin’s effect on cardiac premature aging. **Results:** We found that canthaxanthin obviously mitigated the cardiomyocyte senescence in vitro. Further mechanistic studies revealed that canthaxanthin seems to alleviate cardiomyocyte senescence by regulating the autophagy process. Furthermore, the effects of canthaxanthin on cardiovascular senescence were further evaluated. We also observed that canthaxanthin mitigated cardiac aging and fibrosis in the aged mice model. **Conclusions:** To sum up, the current work showed that canthaxanthin could obviously alleviate cardiac premature aging, indicating that canthaxanthin could be used as a biologically active molecule for the treatment of cardiac aging and fibrosis.

**Keywords:** cardiovascular aging; canthaxanthin; fibrosis; p16; autophagy

**1. Introduction**

At present, the proportion of older people in the world is increasing year by year, and the proportion of the older population will increase from 12% in 2015 to 22% in 2050 [1]. Therefore, the aging society will face huge economic, social, and health challenges in the next few decades. Studies have found that more than 90% of cardiovascular disease occurs in the older population [2], indicating that aging is one of the major risks involved in the development of cardiovascular disease. Aging is a complex physiological process [3]. Cardiac aging refers to age-dependent degenerative changes, which in turn make the heart more sensitive to stress damage [4]. The risk of age-related cardiovascular disease in older people increases significantly year by year. Therefore, delaying the occurrence and retarding the development of cardiac aging is an important strategy in the prevention of aging-related cardiovascular diseases.

Canthaxanthin (8, 8-carotene-4, 4′-dione) has a molecular formula of C40H38O2 and a relative molecular mass of 564.84 [5]. Canthaxanthin (CX) was first isolated from the edible mushroom Celosia japonica [6]. Canthaxanthin comes from a wide range of sources, including Haematococcus pluvialis and Digitococcus chlorellae [7]. In addition to algae, corn, shrimp, mollusks, and insect larvae also contain the abundant canthaxanthin [7]. Because humans cannot synthesize canthaxanthin, canthaxanthin can only be obtained from the diet. Currently, canthaxanthin is frequently used as a food additive with positive effects on animal health [8]. Similar to carotenoids, CX has good antioxidant and immune regulation properties in vitro and in vivo [9–12]. Furthermore, CX can scavenge free radicals and reactive oxygen species (ROS) and prevent DNA damage [10]. Previous studies have shown that canthaxanthin has various biological activities, such as antioxidant and immune regulation [11]. Dietary CX also significantly increases mitogen-induced lymphocyte proliferation in rats [11]. In addition, it has been reported that canthaxanthin has immunomodulatory activity [12].

Currently, the effect of canthaxanthin on cardiovascular premature aging has not been studied. To this end, in the present study, we established a model of cardiomyocyte senescence induced by D-galactose, which was used to further explore the effect of canthaxanthin on oxidative-stress-induced aging of cardiomyoblasts and arterial endothelial cells. The experimental observations revealed that...
CX might mitigate cardiomyoblast senescence and cardiac premature aging. Further mechanistic studies indicated that canthaxanthin might alleviate cardiomyoblast senescence by regulating the autophagy process. To sum up, this work showed that canthaxanthin obviously mitigates cardiomyoblast senescence and cardiac premature aging, suggesting that canthaxanthin has the potential application value for treating premature aging of the heart.

2. Materials and Methods

2.1 Reagents and Antibodies

Bicinchoninic Acid Assay (BCA) protein concentration assay kit, Reactive oxygen species detection kits, RIPA lysate, and β-galactosidase assay kit were purchased from Beyotime Biotechnology Co., Ltd. (Shanghai, China); 4% paraformaldehyde, trypsin, and D-galactose (D-galactose) were purchased from Sigma Company (Shanghai, China). Cell Counting Kit-8 (CCK-8) kit was purchased from Boster Biological Company (Shanghai, China). The SDS-PAGE gel preparation kit, TBST, PMSF protease inhibitor, SOD detection kits, and MDA detection kits were purchased from Beijing Solar-bio Company (Beijing, China). PVDF membrane was purchased from Millipore Company (Burlington, MA, USA). Anti-Col-1 (1:500 dilution), anti-α-SMA (1:1000 dilution), anti-β-catenin (1:1500 dilution), anti-p21 (1:1000 dilution), anti-p16 (1:1500 dilution), anti-p53 (1:1000 dilution), C62 (1:1500 dilution) and LC3 (1:500 dilution) were purchased from Abcam (Cambridge, UK). Bovine Serum Albumin (BSA), Dulbecco’s Modified Eagle Medium (DMEM), ECL Chemiluminescent, and Western-blot Kits were purchased from Thermo-Fisher Scientific company (Waltham, MA, USA). H9c2 (Cat.no. CRL-1446) and Primary Coronary Artery Endothelial Cells were purchased from ATCC (Manassas, VA, USA).

2.2 Cell Culture

H9c2 (rat embryonic cardiomyoblast-derived cell line) was purchased from the ATCC. HCAES (Human Primary Coronary Artery Endothelial Cells, Cat.no. PCS-100-020) was from ATCC. H9c2 and HCAES cells were cultured in the DMEM medium supplemented with 10% FBS, 100 U/mL penicillin, and 100 µg/mL streptomycin at 37 °C and 5% CO₂. In addition, mycoplasma testing has been performed for the H9c2 and HCAES. H9c2 and HCAES have been authenticated by STR.

2.3 Sa-β-gal Staining

After the cell culture medium was discarded, paraformaldehyde (4%) was added to the cells, and cells were fixed for 0.5 h. After washing, freshly prepared Sa-β-gal staining solution was added to the cells and incubated overnight at 37 °C (the cell culture plate was sealed with plastic wrap to prevent the staining solution from evaporating). The cells were checked under a microscope.

2.4 Western-Blot Analysis

Cells were processed using cell lysate; the cells were then centrifuged to harvest at 12,000 g for 30 min at 4 °C. After centrifugation, the supernatant was collected. A BCA kit was used to determine protein concentration. Proteins were separated by SDS-PAGE and transferred to PVDF membranes. The PVDF membrane was placed in a 5% BSA blocking solution for 2 h at room temperature. PVDF membranes were then placed into the diluted primary antibody solution for 12 h at 4 °C. After washing, PVDF membranes were placed in the secondary antibody solution for 120 min at room temperature (RT). After three washes in TBST (10 min each time), the blots were detected by the enhanced chemiluminescence (ECL) kit. The protein band images were analyzed using image J software (developed by National Institutes of Health, Bethesda, MD, USA).

2.5 CCK-8 Assay for Cell Viability

H9c2/HCAES cells were seeded into 96-well plates at a density of 5000 cells per well and cultured for 24 h, after which the cells were cultured with fresh medium containing D-gal at different concentrations (5, 10, 20, 40, 60 g/L), the cells were cultured for 48 h. After washing, 10 µL of CCK-8 solution was added to each well and incubated for 1.5 h. The absorbance of cell samples was detected using a microplate reader at a wavelength of 450 nm.

2.6 Flow Cytometry

The cells were washed twice with PBS, and 0.25% trypsin was added to digest the cells. When the cells were obviously round, and then the cell suspension was collected. The cells were treated with 5 µL of Annexin V-FITC (for 10 min) and 5 µL of PI solution (for 10 min), respectively. The cell samples were subsequently detected using a Flow cytometer (BD). Data were analyzed using FlowJ software (developed by Flow Cytometry Software Inc, version: 10.8.1, Sunnyvale, CA, USA).

2.7 Mitochondrial Membrane Potential Detection

The cells were digested with trypsin. The cells were then centrifuged at 1000 rpm for 5 min. Cells were washed three times with PBS, 1 mL of complete medium was added, then JC-1 staining solution was added and incubated in the dark for 20 min. The JC-1 staining solution was discarded. After 2 washes with pre-cooled PBS buffer, cells were detected via a Flow cytometer.

2.8 Detection of ROS Content

The medium was discarded, and the cells were washed once with sterile PBS. ROS fluorescent probe (DCFH-DA, 1:1000 dilution) was added. After incubation for 20 min, the medium was discarded, and the cells were washed three times with a serum-free medium. The cells were checked under a fluorescence microscope within 30 min.
Fig. 1. Establishment of H9c2 senescent cell model. (A) The cell viability of H9c2 cells was decreased after D-gal treatment. (B) D-gal induced the senescence of H9c2 cells. (C) Analysis of cell apoptosis by Flow analysis. Cells were treated with different concentrations of D-gal, and then the cell apoptosis was analyzed using an Annexin V-PE/7-AAD double staining kit according to the manufacturer’s instructions. (D) The expression of p16 and p21 was up-regulated in the D-gal treatment group. n = 3 from 3 independent experiments. *p < 0.05, **p < 0.001.

2.9 Detection of Reactive Oxygen Species (ROS) by Flow Cytometry

The reactive oxygen species detection kit (Solarbio, Beijing, China) was used to detect the ROS level. The cells were seeded in 6-well plates. When the cells reached 70–80% confluence, the cells were treated with CX. After washing, the cells have been digested. The cells were collected by centrifugation and rinsed once with a serum-free medium. DCFH-DA was then added and incubated for 20 min. After washing three times with PBS, the cell samples were analyzed by Flow cytometry.

2.10 MDA Detection Kit

Malondialdehyde content was determined using an MDA detection kit. After canthaxanthin administration, the cells were digested and prepared into a single-cell suspension. The cells were then centrifuged for 5 min. The cell samples were centrifuged at 8000 g/min for 10 min. Cell supernatants were assayed by a microplate reader.

2.11 In Vivo Experiments

Experimental animals (C57 mice, weight 28–30 g, 18-month-old) were purchased from Huafu Kang Biotechnology Co (Beijing, China). We treated mice with canthaxanthin (15 mg/kg) by gavage for 6 weeks, and the control group was treated with normal saline by gavage. All animal experiments were approved by the animal ethics committee of Jinan University (approval number: 20200326).

2.12 HE Staining

Paraffin-embedded myocardial tissue was sectioned into a thickness of approximately 4 μm. After the sections were deparaffinized, they were placed in xylene I (20 min), xylene II (20 min), absolute ethanol I (5 min), absolute ethanol II (5 min), and 75% alcohol (5 min), then sections were washed and stained with Hematoxylin and Eosin. After sealing with neutral gum, samples were observed using a microscope. The images were analyzed by ImageJ software.
2.13 Masson Staining
Masson staining was performed to analyze the degree of tissue fibrosis. Cardiac tissue was dehydrated with different concentrations of ethanol and then embedded in paraffin. Subsequent processing of tissue samples was performed according to Masson staining kit instructions.

2.14 Indirect Immunofluorescence Staining
After H9c2 cells were treated with canthaxanthin, PBS was added to rinse the cell slides 3 times (3 min each). Cells were fixed using pre-chilled acetone for 15 min. After the samples were washed 3 times, 0.5% Triton X-100 was added to the cells. After incubation for 20 min, cell samples were washed three times. After blocking with BSA, the cells were placed in the primary antibody solutions for 12 h. After 3 washes with PBS, fluorescently labeled secondary antibody was added and incubated at 37 ℃ for 1 h. After the cell samples were washed three times with PBS, DAPI staining solution was added and incubated in the dark for 10 min. After washing three times with PBS, the samples were treated with 50% glycerol and observed under a confocal microscope (Olympus, FV3000, Tokyo, Japan).

2.15 Statistical Analysis
The data were expressed as mean ± standard error, the comparison between multiple groups was performed by one-way ANOVA, and the comparison between two groups was performed by Student-t test, and p < 0.05 was considered statistically significant.

3. Results
3.1 Establishment of D-gal-Induced H9c2 Cardiomyocyte Senescence Model
D-gal was used to establish a model of cardiomyoblast senescence. For this, we first analyzed the effect of D-gal on the viability of H9c2 cells. In the per-experiment, the cells were cultured for 48 h in a complete medium containing 5 g/L, 10 g/L, 20 g/L, 40 g/L, and 60 g/L D-gal, respec-
Fig. 3. Effect of canthaxanthin on oxidative stress. (A) The reactive oxygen species (ROS) content was significantly decreased by canthaxanthin treatment. (B) The effect of canthaxanthin on the levels of Malondialdehyde (MDA) and Superoxide Dismutase (SOD). n = 3 from 3 independent experiments. **p < 0.01, ***p < 0.001.

3.2 Canthaxanthin Alleviated D-gal-Induced Senescence in H9c2 Cardiomyocytes

In order to study the effect of canthaxanthin on H9c2 senescence, we first determined the concentration of canthaxanthin used in the current work by CCK8 assays. The results showed that 10 µM canthaxanthin showed a good effect on promoting cell proliferation (Fig. 2A). Therefore, we used 10 µM canthaxanthin for the following experiments. To analyze the effect of canthaxanthin on the senescence of H9c2 cells, SA-β-gal staining was performed to detect the cell senescence in each group. As shown in Fig. 2B, compared with the control group, the number of SA-β-gal-positive cells in the D-gal group was significantly increased; compared with the D-gal group, the number of senescent cells was significantly decreased after 48 h of canthaxanthin treatment. Next, we analyzed the effect of canthaxanthin on D-gal-induced mitochondrial membrane potential in H9c2 cells; the results of JC-1 staining showed that compared with the control group, the mitochondrial membrane potential of the D-gal treatment group decreased...
significantly (Fig. 2C). However, canthaxanthin treatment increased the mitochondrial membrane potential. Furthermore, the proportions of cells in S phase also increased (Fig. 2D). The effect of canthaxanthin on D-gal-induced expression of senescence-related markers in H9c2 cells was also investigated. Western blot analyses were performed. As shown in Fig. 2E, compared with the control group, D-gal treatment up-regulated the protein expressions of p16, p21, and p53 in H9c2 cells. In addition, γH2AX (DNA-damage-sensing molecule) also obviously elevated. Canthaxanthin treatment decreased the expression of p16, p21, p53, and γH2AX (Fig. 2F) in the H9c2 cells.

3.3 The Effect of Canthaxanthin on Oxidative Stress in the H9c2 Cell

We investigated the effect of canthaxanthin on D-gal-induced oxidative stress in H9c2 cardiomyoblasts. Flow cytometry analyses were performed to detect ROS content. As shown in Fig. 3A, the mean fluorescence intensity of the D-gal group was enhanced compared with the control group, indicating that the ROS content was significantly increased; compared with the D-gal group, the ROS level in H9c2 cardiomyoblasts was significantly decreased after 48 h of canthaxanthin treatment. Furthermore, the effect of canthaxanthin on the levels of Malondialdehyde (MDA)
Fig. 5. Canthaxanthin alleviated the human primary coronary artery endothelial cells (HCAEC) senescence. (A) D-gal treatment decreased the viability of HCAEC cells compared with the control group. (B) Canthaxanthin reduced the senescence of HCAEC cells. (C) Canthaxanthin increased the proportion of cells in S phase. (D) Canthaxanthin could increase the mitochondrial membrane potential of the senescent cells. (E) Canthaxanthin treatment down-regulated the expression of p53, p21 and p16. (F) γH2AX expression was detected by IFA. The cell membranes were stained using membrane-specific staining reagent DIO (3,3′-dioctadecyloxacarbocyanine perchlorate). n=3 from 3 independent experiments. *p < 0.05, **p < 0.001.

and Superoxide Dismutase (SOD) was also detected. As shown in Fig. 3B, SOD in the D-gal treatment group was significantly decreased, but MDA content was significantly increased. Compared with the D-gal group, the SOD in the canthaxanthin treatment group was increased, and the MDA content was obviously decreased. These results suggest that canthaxanthin could alleviate the oxidative stress of the senescent cardiomyocytes.

3.4 Exploration of the Potential Molecular Mechanism by which Canthaxanthin Could Alleviate Cardiomyocyte Aging

Oxidative stress is closely related to aging, and it has been reported that canthaxanthin has antioxidant effects. Here, we further analyzed whether canthaxanthin could alleviate H9c2 cell senescence through other molecular mechanisms. Previous studies have shown that cardiac aging is closely related to autophagy [13]. In the senescent cells, autophagy levels were significantly reduced [14]. In the current study, we found that in the senescent H2c9 cells, LC3 expression was significantly decreased and p62 expression was increased (Fig. 4A). In contrast, the level of autophagy was significantly increased in the canthaxanthin-treated group by detecting LC3 (Supplementary Fig. 1). These results suggest that canthaxanthin mitigates cardiac premature aging via regulating autophagy. In order to further verify whether canthaxanthin inhibits myocardial aging by regulating autophagy, we conducted a rescue experiment; canthaxanthin was added into the cell medium, at the same time, an inhibitor of autophagy (chloroquine diphosphate, ab142116, 50 µM) was also added. The experimental results showed that the autophagy inhibitor neutralized the anti-aging effects of canthaxanthin (Fig. 4B), which suggests that the anti-aging effect of canthaxanthin may be through the regulation of autophagy (at least in part).

3.5 Canthaxanthin Alleviated the Senescence of HCAEC

We also established the model of D-gal-induced HCAEC senescence. We first analyzed the effect of D-gal on HCAEC viability. Different concentrations of D-gal (5–80 g/L) were used in the pre-experiment, and results showed that 30 g/L D-gal could successfully induce HCAEC senescence. Fig. 5A showed that D-gal treatment decreased the viability of HCAEC compared with the control group. In contrast, canthaxanthin treatment could significantly increase the viability of HCAEC. Furthermore, D-gal treatment led to a significantly higher num-
Fig. 6. Effects of canthaxanthin on oxidative stress in HCAEC. (A) The level of ROS in cells was significantly decreased after canthaxanthin treatment. (B) The SOD in the canthaxanthin treatment group was increased, and the MDA content was decreased. n = 3 from 3 independent experiments. ***p < 0.001.

3.6 Effects of Canthaxanthin on Oxidative Stress in HCAEC

We investigated the effect of canthaxanthin on D-gal-induced oxidative stress in HCAEC. Flow cytometry was used to analyze the ROS content. The results are shown in Fig. 6A; the mean fluorescence intensity of the D-gal treatment group was obviously enhanced. Compared to the D-gal group, the level of ROS was significantly decreased after the H9c2 cells were treated with canthaxanthin for 48 h. Furthermore, the effect of canthaxanthin on the expression levels of MDA and SOD in the HCAEC cells was also analyzed, and the results showed that the SOD in the D-gal group was significantly lower than that of the control group, but the MDA content was significantly higher. Compared to the D-gal group, the SOD in the canthaxanthin treatment group was increased, and the MDA content was decreased (Fig. 6B). These findings indicated that canthaxanthin can relieve the oxidative stress of HCAEC cells.

3.7 Anti-Aging Effects of Canthaxanthin in Vivo

In in vitro experiments, we evaluated the effect of canthaxanthin on heart aging. For an in vivo experiment, we selected 18-month-old mice as research subjects. We treated
the experimental mice with canthaxanthin (15 mg/kg) by gavage and the control mice with normal saline by gavage. Heart tissues were extracted for analysis of aging markers. Canthaxanthin treatment reduced the SA-β-gal staining area of the heart tissue (Fig. 7A). In addition, the expression of both p16 and p21 was also significantly lower than that of the control group (Fig. 7B). The number of Ki67-positive cells was significantly higher in the CX-treated group (Fig. 7C).

Heart aging is closely related to fibrosis; therefore, we also analyzed the effect of CX on myocardial fibrosis in vivo. Masson staining showed that the level of myocardial fibrosis was significantly lower in the CX-treated group (Fig. 8A). Furthermore, CX also inhibited the expression of Col-1 in the cardiac tissue (Fig. 8B), and the expression of α-SMA was also decreased in the heart tissue (Fig. 8C). On this basis, we analyzed a molecular mechanism by which CX might inhibit cardiac fibrosis. It is well known that activation of β-catenin leads to fibroblast activation and fibrogenesis in multiple organ systems. In myocardial tissue, it directly initiates the expression of the Col-1 gene. Therefore, in order to determine whether the effect of CX on myocardial fibrosis is related to β-catenin, Western-blot analyses were conducted to detect the expression level of β-catenin. Compared with the control group, the expression of β-catenin protein in the myocardium of the aged group was higher, and CX treatment reduced the expression of β-catenin protein in the myocardium of the aged mice (Fig. 8D).

Compared with the hearts of young mice, the heart/body-weight ratio (HW/BW) of the 18-month-old mice was abnormal; the ratio was significantly higher in the aged heart (Fig. 9A), suggesting that the myocardium may be hypertrophic. The expression of brain natriuretic peptide (BNP) and β-myosin heavy chain (MHC) (the markers of cardiac hypertrophy in mice) were significantly higher. However, the expressions of BNP and MHC were significantly lower after CX treatment (Fig. 9B). These results suggest that CX may partially alleviate cardiac hypertrophy in aged mice. Additionally, H&E staining showed that
Fig. 8. The effect of CX on myocardial fibrosis in vivo. (A) The level of myocardial fibrosis was down-regulated in the canthaxanthin-treated group. (B) Canthaxanthin inhibited the expression of Col-1 in cardiac tissue. (C) The expression of α-SMA was decreased in the heart. (D) Canthaxanthin treatment reduced the expression of β-catenin in the myocardium of the aged group mice. *p < 0.05.

the cardiomyocytes of young WT mice were arranged compactly in an orderly manner, but the cardiomyocytes of the aged mice were irregularly arranged, with enlarged intercellular spaces. CX treatment significantly alleviated the pathological changes caused by aging (Fig. 9C).

4. Discussion

Population aging has become one of the major social problems in the world [15]. This brings great challenges to limited social medical resources [16]. The aging population has brought unprecedented challenges to medical and health services [17]. Aging is a process of chronic accumulation of biological changes in the body over time. Because of the importance of the heart, cardiac aging is one of the important pathological factors in the aging of the body [18]. Symptoms of cardiac aging include left ventricular wall thickening, myocardial fibrosis, and decreased diastolic function [19]. The expression of aging-related markers p16 and p21 has been found to increase in aging cardiomyocytes of humans and animals. Cardiomyocyte aging is closely related to the occurrence of cardiovascular diseases such as heart failure, atherosclerosis, and arrhythmia [19]. Therefore, there is an urgent need for us to further elucidate the underlying molecular mechanisms of cardiac aging and to find effective treatments or interventions to delay or retard cardiac aging. In the present study, we found that CX has a promising anti-aging effect on the heart.

In the present study, H9c2 was used as a cell model to establish a senescent cardiomyoblast model. In in vitro experiments, we found that CX could alleviate the senescence of H9c2. The proportion of Sa-β-gal-positive cells was significantly reduced in the CX-treated group. Furthermore, aging-related markers were also reduced. Additionally, in vivo, we found that CX exhibited anti-aging effects. Furthermore, the expression of p16 and p21 was reduced after CX treatment, and the rate of Ki67-positive cells was increased. Heart aging is closely related to fibrosis. We found that the level of myocardial fibrosis was reduced after CX treatment. In addition, in recent years, CX has been reported to exhibit a number of other important biological activities [20]. For example, (a) CX inhibited the proliferation of cancer cell lines [21]; (b) CX enhanced the productive and reproductive performance of broiler breeders [22]; and (c) CX has antioxidant properties [10,23].
Autophagy is closely related to cardiomyocyte aging, which is responsible for degrading the damaged organelles and macromolecules [24]. Autophagy can be activated under stress. However, studies have shown that autophagy gradually decreases with aging. Therefore, the reduction of autophagy is one of the important reasons for cardiac aging [25]. In the current study, we showed that CX can increase autophagy in the senescent cardiomyoblast, which may be one of the potential mechanisms by which CX exhibits an anti-aging effect.

The incidence of age-related cardiovascular diseases increases exponentially with the global aging trend [26,27]. Studies have shown that vascular aging plays a key role in the occurrence and development of aging-related diseases [26,27]. Senescent endothelial cells accumulate continuously in the vessel wall, leading to the decline of endothelial function [28]. Previous studies have shown that senescent endothelial cells accumulate in atherosclerotic plaques, leading to vascular dysfunction [29]. Therefore, endothelial cells can be used as a tool for evaluating vascular aging and function. In the present study, we found that CX could significantly alleviate the senescence of vascular endothelial cells. These results suggest that CX can relieve vascular endothelial cell senescence.

Cardiac aging is closely related to oxidative stress [30]. Oxidative stress occurs when ROS exceeds the buffering capacity of the antioxidant defense system. ROS-mediated oxidative damage plays an important role in vascular aging. In endothelial cells, nitric oxide (NO) is considered to be the main substance maintaining endothelial-cell function. In the present study, we found that CX can significantly increase vascular NO content.

CX has many advantages over the anti-aging substances that have been mentioned in previous reports. CX can be eaten as food or applied as a medication (because anti-aging is a long-term process). An interesting question is, how much CX does the body get from food every day? As of now, there is no data on how much CX can be obtained from food every day because inter-individual variability is very large. However, studies have shown that CX can be used in a very wide range of safe concentrations.

Of course, current research also has certain limitations. H9c2 cells are cardiomyoblast-derived immortal cells that are fundamentally different from terminally differentiated cardiomyocytes with respect to cell cycle progression. Since cell cycle arrest is a major determinant of cellular senescence, H9c2 cells are not the best model for cardiomyocyte senescence. Theoretically, primary car-
diomyocytes should be the best cellular model, but under our experimental system, primary cardiomcyocytes cannot meet the current experimental requirements. Therefore, we finally chose H9c2 as a model to evaluate the anti-aging potential of canthaxanthin.

5. Conclusions

In the current study, we showed that CX can alleviate cardiomyoblast senescence via regulating the autophagy process, indicating that canthaxanthin may be an important bioactive molecule that has shown potential for alleviating cardiac aging and fibrosis.

Abbreviations

CX, Canthaxanthin; MHC, β-myosin heavy chain; HCAES, human primary coronary artery endothelial cells; DMEM, dulbecco’s modification of eagle’s medium; ROS, reactive oxygen species.

Availability of Data and Materials

The data used during the current study are available from the corresponding author on reasonable request.

Author Contributions

ZW, KZ and SD designed the research; WL, MZ, JY performed the research; JF and KZ provided help and advice on the conclusions; MZ and JF analyzed the data; ZW wrote the manuscript. All authors contributed to editorial changes in the manuscript. All authors read and approved the final manuscript. All authors have participated sufficiently in the work and agreed to be accountable for all aspects of the work.

Ethics Approval and Consent to Participate

All animal experiments were approved by the animal ethics committee of Jinan University (approval number: 20200326).

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Conflict of Interest

The authors declare no conflict of interest.

Supplementary Material

Supplementary material associated with this article can be found, in the online version, at https://doi.org/10.31083/j.fbl2902070.

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