Sulforaphane protects against oxidative stress-induced apoptosis via activating SIRT1 in mouse osteoarthritis

MANGMANG CHEN¹², LIUPENG HUANG², YANGXUN LV², LIUBING LI¹ and QIRONG DONG¹

¹Department of Orthopedics, The Second Affiliated Hospital of Soochow University, Suzhou, Jiangsu 215004; ²Department of Orthopedics Surgery, The Dingli Clinical Institute of Wenzhou Medical University, Wenzhou Central Hospital, Wenzhou, Zhejiang 325000, P.R. China

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Abstract. Osteoarthritis (OA), the most common form of human joint disease, is characterized by progressive degeneration of the articular cartilage, synovitis and subchondral osteoporosis. Chondrocyte apoptosis is the primary pathogenic mechanism of OA and is considered to be a potential therapeutic target. Sulforaphane (SFN), a dietary isothiocyanate obtained from cruciferous vegetables, has been reported to exert an anti-apoptotic effect by activating sirtuin 1 (SIRT1). To the best of our knowledge, however, the effects of SFN on apoptotic responses in OA have not been reported. In the present study, SFN was shown to significantly inhibit chondrocyte apoptosis while enhancing expression levels of SIRT1 in a H2O2-induced OA mouse model. The anti-apoptotic effect of SFN was reversed by SIRT1 small interfering RNA, implying that SIRT1 exerted a protective role against the effect of SFN on chondrocytes. The expression levels of C/EBP homologous protein, 78-kDa glucose regulated protein, Bax, Bcl-2 and cleaved caspase 3 were found to be downregulated in SFN-treated mice. Furthermore, SFN ameliorated cartilage degradation in the OA mouse model. These findings indicate that SFN exerted an anti-apoptotic effect on chondrocytes and ameliorated OA in vivo by activating the SIRT1 signaling pathway.

Introduction

Osteoarthritis (OA), a degenerative disease of joints, is the most common cause of musculoskeletal disability (1,2). Various factors, such as age, genetics, adiposis and sex, are involved in the development of osteoarthritis (3). Destroyed cartilage structure and loss of function are the central hallmarks of OA (4). Although studies have revealed potential mechanisms involved in the progression of OA, its pathological etiology is unknown. Moreover, there are no effective therapeutic options that can prevent disease progression (5,6). OA is associated with degradation of matrix cartilage and chondrocytes, which are the only type of cell in articular cartilage (7). Extracellular matrix (ECM) molecules are involved in maintaining the structure and function of regular cartilage, whose components include proteoglycans and collagen, produced by chondrocytes (8). Articular cartilage degradation is initiated by several inflammatory cytokines (IL-6, TNF-α, prostaglandin 2) and reactive oxygen species (ROS) that induce oxidative stress (9). Excessive chondrocyte apoptosis and senescence occur due to ROS accumulation (10). Therefore, inhibiting chondrocyte apoptosis and senescence is a potential strategy for preventing articular cartilage degradation in OA.

Endoplasmic reticulum (ER), which is the first site in the secretory pathway, synthesize and fold proteins (11). ER regulate the balance between pathological and physiological conditions (12-14). The ER stress (ERS) response enhances cell survival by decreasing the load of unfolded proteins. However, cells undergo apoptosis when ERS is overwhelming (15). ERS-associated apoptosis is an important signaling pathway during apoptosis (16). It has been demonstrated that ERS signaling in this pathway promotes repair of the unfolded protein response (UPR) of the ER (17). UPR serves a critical role in cell growth regulation, differentiation and apoptosis (18). ERS has been shown to stimulate the C/EBP-homologous protein (CHOP) pathway while activating glucose-regulated protein 78 (GRP78) (19). The Bcl-2 protein family acts against apoptosis by inhibiting apoptotic mediators, such as Bax and Cytosome c (20). Studies have reported that ER and mitochondrial pathways regulate apoptosis during rat intervertebral disc degeneration (21,22). These findings show the importance of ER and mitochondria in chondrocytes.

There are seven known members of mammalian Sirtuins (SIRTs). Among them, SIRT1 regulates multiple functions in metabolic syndrome, oxidative stress, inflammation and aging (23-25). Li et al (26) reported that sulforaphane (SFN) can downregulate ER-induced apoptosis protein expression levels in a rat model of hypoxia/reoxygenation injury.
by activating SIRT1. Activated SIRT1 inhibits oxidative damage and inflammation (26) and is therefore a therapeutic target for OA via regulating mitochondrial biogenesis (27). Feng et al (28) reported that curcumin exerts a protective effect against OA by activating SIRT1.

SFN (a type of dietary isothiocyanate) is primarily found in cruciferous vegetables (29). It exhibits various pharmacological properties, including anti-cancer, anti-inflammatory and anti-oxidative stress (30). SFN exerts a protective effect during post-ischemic cardiac injury (31). Moreover, it exerts cardioprotective effects by activating SIRT1 and suppressing ER stress (32). Davidson et al (33) reported that SFN inhibits inflammation via the NF-κB signaling pathway in OA mice. Moreover, they found that SFN suppresses IL-1/NF-κB and Wnt3a/T cell factor/lymphoid enhancer factor signaling while enhancing TGF-β/SMAD2/3 and bone morphogenetic protein 6/SMAD3/5/8 signaling in IL-1/oncostatin M-induced chondrocytes (34). However, its anti-apoptotic effects during OA have not been established. The aim of the present study was to elucidate the role of SFN in apoptosis, including its specific mechanisms of action, in H2O2-treated chondrocytes. The effect of SFN on cartilage degeneration was investigated in a mouse model of OA.

Materials and methods

Reagents and antibodies. SFN (purity ≥98%), Safranin O and toluidine blue stain were obtained from Beyotime Institute of Biotechnology; collagenase type II, H2O2 and DMSO were bought from Sigma-Aldrich (Merck KGaA). BCA protein assay kit was obtained from Beyotime Institute of Biotechnology; primary antibodies against GRP78, CHOP and GAPDH were obtained from Wuhan Sanying Biotechnology; primary antibodies against Bax, Bcl-2, Cleaved caspase-3 and SIRT1 were purchased from Cell Signaling Technology, Inc.; In Situ Cell Death Detection kit was obtained from Roche Diagnostics; secondary antibodies (cat. no. SA00001-14; Wuhan Sanying Biotechnology), including Goat Anti-Rabbit IgG (cat. no. B900210; Wuhan Sanying Biotechnology) and Alexa Fluor® 488-labeled goat anti-rabbit IgG (H+L) secondary antibody (cat. no. SA00009-3; Wuhan Sanying Biotechnology), were obtained from Jackson ImmunoResearch Laboratories, Inc.; Cell-Counting Kit-8 (CCK-8) was purchased from Dojindo Molecular Technologies, Inc. DMEM-F12 was purchased from Gibco (Thermo Fisher Scientific, Inc.).

Primary chondrocyte isolation and culture. A total of 10 C57BL/6 mice (5 males and 5 females; age, 10 days; weight, 4 g; the Animal Center of the Chinese Academy of Sciences) were euthanized by 100 mg/kg sodium pentobarbital. All animal procedures were performed in accordance with the Guidelines for Care and Use of Laboratory Animals of Wenzhou Medical University (35) and experiments were approved by the Animal Ethics Committee of Wenzhou Medical University. The knee cartilage of mice were collected carefully under aseptic conditions using a dissecting microscope, and tissue was treated with 2 mg/ml (0.1%) collagenase II for 4 h at 37°C. The digested tissue was centrifuged (800 x g) for 5 min at 37°C. Then, chondrocytes (5x10^6 cells/cm^2) were seeded into culture flasks. Cells were cultured in DMEM/F12 supplemented with 10% fetal bovine serum and 1% antibiotics (penicillin/streptomycin) and incubated in an atmosphere of 5% CO2 at 37°C. The culture medium was replaced every 2-3 days. When 80-90% confluence was attained, cells were fused and 0.25% trypsin-EDTA solution was used to subculture the chondrocytes. In order to avoid phenotypic loss, chondrocytes from the first and second channels were used.

Cell viability assay. CCK-8 assay was used to evaluate chondrocyte viability according to the manufacturer's instructions. First, chondrocytes were incubated in 96-well plates (8,000 cells/well) for 24 h. Then, they were treated with a concentration gradient (0.0, 12.5, 25.0, 50.0, 100.0 and 200.0 µM) of SFN for 24 h and 48 h. Finally, 10 mol/l CCK-8 solution was added to each well for 2 h at 37°C, after which optical density was spectrophotometrically measured at 450 nm (Thermo Fisher Scientific, Inc.).

Intracellular ROS production assay. Intracellular ROS levels were detected using the HDCFDA Probe Assay kit (Beyotime Institute of Biotechnology), according to the manufacturer's instructions. Intracellular ROS generation was measured at 485 nm (excitation) and 535 nm (emission) using a micro-plate reader (Bio-Rad Laboratories, Inc.).

Western blot analysis. In order to obtain total proteins from chondrocytes, RIPA lysis buffer (Beyotime Institute of Biotechnology) was added to 1 mM PMSF and kept on ice for 10 min, after which the mixture was centrifuged at 1,000 x g for 15 min at 4°C. BCA protein detection kit was used to estimate protein concentration. Then, 40 ng protein was separated by SDS-PAGE (10%) and transferred to polyvinylidene fluoride membrane. Membranes were blocked using 5% skimmed milk for 2 h at 37°C and detected overnight at 4°C with the following antibodies: CHOP (1:1,000; cat. no. 15204-1-AP; Wuhan Sanying Biotechnology), GRP78 (1:1,000; cat. no. 115871-1-AP; Wuhan Sanying Biotechnology), SIRT1 (1:1,000; cat. no. 15071; Cell Signaling Technology, Inc.), Bax (1:1,000; cat. no. 15023; Cell Signaling Technology, Inc.), Cleaved caspase-3 (1:1,000; cat. no. 15204-1-AP; Wuhan Sanying Biotechnology), and GAPDH (1:1,000; cat. no. 10494-1-AP; Wuhan Sanying Biotechnology).

The membrane was washed using TBS-Tween-20 (TBST; 5% Tween-20) and incubated with the corresponding secondary antibodies (1:1,000; cat. no. SA00001-14; Wuhan Sanying Biotechnology) for 2 h at room temperature. After being washed three times using TBST, the blots were visualized using electrochemiluminescence plus reagent (Invitrogen; Thermo Fisher Scientific, Inc.). Image Lab 3.0 software (Bio-Rad Laboratories, Inc.) was used to measure the intensity of each band.

TUNEL assay. Apoptotic chondrocytes were measured by TUNEL staining using an In Situ Cell Death Detection kit, according to the manufacturer's instructions. The chondrocytes (1x10^5) were seeded in a 6-well plate and treated with SFN (50 µM) or H2O2 (50 µM) for 24 h at 37°C, fixed in 4% paraformaldehyde for 45 min at 37°C, incubated with 0.5% Triton X-100 for 15 min at 37°C and washed using PBS.
for 5 min. Finally, cells were stained using the In Situ Cell Death Detection kit for 60 min at 65°C, after which the nuclei were stained with DAPI for 1 min at 37°C (Beijing Solarbio Science & Technology Co., Ltd.). In total, 25 fields of each slide were randomly selected and images were observed using a fluorescence microscope (magnification, x100; scale bar, 50 µm; Olympus Corporation).

Small interfering (si)RNA transfection for 24 h. Double-stranded siRNA for mouse SIRT1 gene silencing was purchased from Abcam. The SIRT1 siRNA sequence was as follows: 5′-UGG GAACUAGCUAGGACUG-3′. Cells (1×10⁴) were seeded into a 6-well plate, incubated for 24 h at 37°C and transfected with negative control [non-specific non-targeting siRNA (scramble); Abcam] or SIRT1 siRNA duplexes for 36 h at 50 nM using Lipofectamine® 2000 siRNA transfection reagent (Thermo Fisherr Scientific, Inc.), according to the manufacturer's instructions. The chondrocytes were serum-starved overnight followed by incubation at 37°C with SFN (50 µM) for 24 h. After 24 h, the interference efficiency of the siRNA was evaluated by reverse transcription-quantitative PCR (RT-qPCR).

RT-qPCR. Total RNA was extracted from cells using TRIzol® reagent (Invitrogen; Thermo Fisherr Scientific, Inc.). The concentration and purity of RNA was evaluated by spectrometry at 260 and 280 nm. The synthesis of cDNA was performed using the PrimeScript™ RT reagent kit [cat. no. 2312, Hangzhou Multisciences (Lianke) Biotech Co., Ltd.]. qPCR was performed using SYBR Green Real-Time PCR Master mix (Thermo Fisherr Scientific, Inc.) according to the manufacturer's protocol. The amplification conditions were as follows: SIRT1, 45 sec at 94°C followed by 30 cycles of 30 sec at 94°C, 30 sec at 52°C and 60 sec at 72°C; GAPDH, 45 sec at 94°C followed by 30 cycles of 30 sec at 94°C, 30 sec at 58°C and 60 sec at 72°C. Primers sequences, designed by Primer-Express V3.0 (Thermo Fisherr Scientific, Inc.), were as follows: SIRT1 forward, 5′-CTCTGGAATGTAGCAGACGG TAGC-3′ and reverse, 5′-TGATAGTGGACAAAGTTC-3′ (product, 213 bp); and GAPDH forward, 5′-TTTGCTCA GGTGTCCCTG-3′ and reverse, 5′-TTTGTGCAAGCTCAT TCTTG-3′ (product, 457 bp). Relative gene expression was analyzed using the 2−ΔΔCT method (36).

Immunofluorescence. Cells were rinsed using PBS and fixed in 4% paraformaldehyde for 15 min at 37°C after which they were washed three times using PBS. Then, they were treated with 0.1% Triton X-100 diluted in PBS for 15 min at room temperature. Next, chondrocytes were blocked using 10% goat serum (cat. no. SL038; Beijing Solarbio Science & Technology Co., Ltd.; dissolved in PBS) for 4 h at 37°C and incubated at 4°C overnight in the presence of primary antibodies against collagen II (1:300; cat. no. 28459-1-AP; Wuhan Sanying Biotechnology). Following incubation, cells were exposed to Alexa Fluor® 594-labelled conjugated secondary antibodies (1:400; cat. no. IC1051T; R&D Systems, Inc.) for 1.5 h. Finally, they were exposed to DAPI (10 µg/ml; 37°C) (Beyotime Institute of Biotechnology) for 1 min and observed using a fluorescence microscope (magnification, x200; scale bar, 20 µm; Olympus Corporation). Fluorescence intensity was assessed using ImageJ software 6.0 (National Institutes of Health).

Immunohistochemical assay. The paraffin-embedded sections (6 µm) were deparaffinized in xylene for 15 min at 37°C and rehydrated by incubation in series of graded ethanol (100, 95, 85, 75 and 0%) for 5 min at 37°C and endogenous peroxidase. Then, sections were treated with 3% (v/v) hydrogen peroxide for 10 min at 37°C. Antigen retrieval of the sections was performed using 0.4% pepsin (Sangon Biotech Co., Ltd.) in 5 mM HCl at 37°C for 20 min. Then, sections were incubated in the presence of 10% BSA (cat. no. ST025; Beyotime Institute of Biotechnology) for 45 min at room temperature, after which they were treated with primary antibodies against SIRT1 (1:500; cat. no. 2493; Cell Signalining Technology, Inc.) overnight at 4°C. Finally, they were treated with horseradish peroxidase-conjugated secondary antibodies (1:1,000; cat. no. SA00001-14; Wuhan Sanying Biotechnology) for 3 h at 4°C. Following incubation at room temperature for 1 h, the slices were rinsed (5 min, 37°C), stained using a Metal Enhanced DAB Substrate kit (2 min, 37°C; cat. no. DA1015; Beijing Solarbio Science & Technology Co., Ltd.), dehydrated (5 min, 37°C), mounted and examined under a microscope (magnification, x40; CX4i; Olympus Corporation). Quantitative analysis was performed using Image-Pro Plus 6.0 software (Media Cybernetics, Inc.).

Animals. A total of 45 C57BL/6 male wild-type mice (age, 10 weeks; weight, 40 g) were obtained from the Animal Center of the Chinese Academy of Sciences (Shanghai, China). The protocol for animal care and use conformed to The Guide for the Care and Use of Laboratory Animals of the National Institutes of Health published by the National Institutes of Health (NIH Publication No. 85-23, revised 1996) (37) and was approved by the Animal Care and Use Committee of Wenzhou Medical University. Experimental OA mice were established by destabilization of the medial meniscus (DMM) as previously described (38). Mice were housed under 12-h light/dark cycles and constant temperature (21-23°C) and humidity (50-60%). Water and food were provided ad libitum. Anesthesia was performed via intraperitoneal injection of 2% (w/v) pentobarbital (40 mg/kg). Faint breathing, myasthenia, lack of independent reaction, cyanosis or coma were considered to indicate that mice were close to death; mice were euthanized by cervical dislocation. The joint capsule was carved followed by transection of the medial meniscotibial ligament of the right knee using microsurgical scissors. During surgery, the lateral meniscotibial ligament was always protected. As a control, the medial meniscotibial ligament was not transected while arthrotomy was performed in the left knee. Mice were randomly separated into three groups (n=15/group): Control (sham-operated), DMM and DMM + SFN [mice were intraperitoneally administered with SFN at 20 mg/kg body weight (39,40)]. Parameters indicating the condition of mice were observed daily, including fur brightness, food and water intake, defecation and behavior. Furthermore, body weight was measured each week.

Histopathologic analysis. The extent of cartilage degeneration and synovitis in stained sections was assessed
using light microscopy with the Osteoarthritis Research Society International (OARSI) scoring system, as previously described (41).

**Statistical analysis.** The data are presented as mean ± SD (n=3). Statistical analysis was performed using GraphPad Prism version 5.0 software (GraphPad Software, Inc.). Comparisons between groups were performed using one-way ANOVA followed by Tukey’s post hoc test. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**Identification of mouse chondrocytes.** Safranin O and toluidine blue staining were used to characterize mouse chondrocytes. Chondrocytes were stained red by Safranin O and the cytoplasm was stained purple by toluidine blue (Fig. 1a and B). In addition, collagen II in chondrocyte cytoplasm was stained red by immunofluorescence with no positive staining observed in the nucleus (Fig. 1c). These results imply that the cells isolated from mouse articular cartilage were chondrocytes.

**Cytotoxicity of SFN on mouse chondrocytes.** The chemical structure of SFN is shown in Fig. 1d. In order to determine the cytotoxic effects of SFN, CCK-8 assay was performed. Cells were incubated with increasing concentrations (0.0, 12.5, 25.0, 50.0, 100.0 and 200.0 µM) of SFN for 24 and 48 h. SFN exhibited cytotoxic effects at ≥100 µM after 24 or 48 h but not <100 µM (Fig. 1E and F). Therefore, 50 µM was selected for use in subsequent experiments.

**Effects of SFN on apoptosis in H₂O₂-treated chondrocytes.** In order to determine the anti-apoptotic effect of SFN on H₂O₂-induced chondrocytes, the expression levels of Bax, Cleaved caspase-3 and Bcl-2 were evaluated by western blot analysis. H₂O₂ elevated levels of Bax and Cleaved caspase-3, whereas Bcl-2 levels were downregulated (Fig. 2A-D). SFN downregulated levels of Bax and Cleaved caspase-3. Moreover, SFN significantly activated SIRT1 (Fig. 2A and E). These data indicate that SFN exerted an anti-apoptotic effect in mouse chondrocytes.

**SFN alleviates ERS and apoptosis of H₂O₂-treated chondrocytes.** In order to determine whether ERS in chondrocytes was associated with the cytoprotective effects of SFN on cartilage, cells were treated with H₂O₂, a typical ERS inducer. Expression levels of ERS-associated factors GRP78 and CHOP were measured. Western blotting assay exhibited increased expression levels of these proteins in response to H₂O₂; these effects were stopped by SFN (Fig. 3A-C). Moreover, apoptosis of cells in the H₂O₂ group was high compared with the control group. SFN treatment decreased cell apoptosis (Fig. 3D and E). Furthermore, H₂O₂ induced ROS production, which was inhibited by SFN (Fig. 3F).

**SFN pretreatment enhances SIRT1 activity in H₂O₂-treated chondrocytes.** In order to confirm the aforementioned results, SIRT1 siRNA was used to detect the potential mechanism of SFN on H₂O₂-stimulated chondrocytes. The mRNA expression levels of SIRT1 were significantly lower in SIRT1-silenced cells than in control cells (Fig. 4F). SIRT1 levels were elevated in the H₂O₂ + SFN group but suppressed in the SIRT1 siRNA transfection group (Fig. 4A and E). Moreover, Bel-2 levels were suppressed in the SIRT1 siRNA transfection group whereas the expression levels of Bax and Cleaved caspase-3 increased significantly (Fig. 4A-D). These results demonstrated the anti-apoptotic effect of SFN, which was mediated by activation of SIRT1.

**SFN ameliorates OA development in a DMM mouse model.** The chondroprotective effect of SFN on OA was investigated in vivo. A DMM mouse model was established, followed by intraperitoneal injection of 20 mg/kg SFN dissolved in DMSO every 2 days for 8 weeks. Safranin O staining was...
used to evaluate the effect of SFn on the development of OA. Compared with the sham group, the surface of the cartilage was notably destroyed in the DMM group (Fig. 5A). The degradation level of the cartilage matrix in the SFn group was lower than the OA but higher than the sham group. The oarSi scores in the SFn group were lower than those of the DMM group (Fig. 5B). Moreover, synovitis score in the DMM group was higher than in the sham group; SFN reversed this effect (Fig. 5a and c). In order to investigate the effect of SFn on SirT1 expression in vivo, immunohistochemical staining for SirT1 was performed (Fig. 5 d and e). Compared with the DMM group, significantly more SIRT1-positive chondrocytes were detected in the DMM + SFn group. These results imply that SFn exhibited chondrocyte-protective effects during the progression of OA.

Discussion

OA is a chronic joint degradation disease that is characterized by long-term pain and joint limitation (42). Oral or topical non-steroidal anti-inflammatory drugs (such as ibuprofen and diclofenac sodium) are the primary therapeutic options for OA but only relieve clinical symptoms (3). These drugs are also associated with side effects, such as heart attack and stroke (43). Nonsteroidal anti-inflammatory drugs can only delay the progression of OA and surgery is the recommended option (44). Therefore, an agent that prevents OA progression, accompanied by fewer side effects, would be a potential therapeutic option for OA. Previous studies aimed to evaluate the efficacy of anti-apoptotic compounds, which exhibit few side effects (45,46). Biochemical and biomechanical factors (such as ROS) induce chondrocyte apoptosis, as well as an imbalance between catabolism and anabolism of the ECM (47). Pharmacological administration of compounds (such as vitamin C and Glutathione) with the ability to regulate genetic mechanisms involved in apoptosis exhibit beneficial effects against OA development in vitro and in vivo (48). Therefore, the mechanism involved in chondrocyte apoptosis may be a potential therapeutic target for OA.

SFN, an isothiocyanate found in cruciferous vegetables, has been shown to exhibit anti-anxiety, sedative-hypnotic and anti-depression effects (49,50). It is effective against oxidative-induced cardiomyocyte damage (51). Moreover, SFN has
been shown to be effective against ERS in different models of cell injury, including chick yolk sac membrane and chorioallantoic membrane models (52). It inhibits ERS by suppressing the apoptosis of cells in type 1 diabetes mellitus (53). The anti-apoptotic effect of SFN was also found to be present in human hepatocytes (54). Wang et al (52) reported that SFN ameliorates ethanol-suppressed embryonic angiogenesis by alleviating excessive ROS production and ERS. Moreover, Li et al (26) documented that SFN suppresses expression levels of ERS-associated apoptosis proteins by activating SIRT1. The present study determined whether SFN ameliorates chondrocyte apoptosis and delays OA progression. H2O2, a key pro-apoptosis factor, was used to induce mitochondrial dysfunction, and upregulated the expression levels of caspase family proteins (55). The effect of SFN on ERS was evaluated in OA mouse chondrocytes. ER are found in the cytoplasm of all eukaryotic cells (56). ERS is involved in the development of OA (57). In the present study, the levels of ERS-associated apoptosis proteins (GRP78 and CHOP), Bax and Cleaved caspase-3 were elevated in chondrocytes following H2O2 treatment. However, pretreatment with SFN downregulated GRP78, CHOP, Bax and Cleaved caspase-3 levels and elevated Bcl-2 levels compared with the H2O2 group. These findings imply that SFN exerted chondroprotective effects against H2O2-induced apoptosis by decreasing ERS-dependent apoptosis.
SirT1 exerts anti-apoptotic effects in OA (58). Certain drugs, such as melatonin, curcumin and sildenafil, have been shown to exhibit chondroprotective effects in OA by modulating the SirT1 signaling pathway (59,60). The present study evaluated the effect of SirT1 pathway signaling in SFN-induced chondroprotection. Compared with the H2O2 group, SirT1 levels were significantly elevated in the SFN-pretreatment group. SirT1 siRNA was used to detect the potential mechanisms of SFN in H2O2-stimulated apoptotic chondrocytes. Consistent with previous studies, RNA interference against SirT1 suppressed SFN-mediated inhibition of the anti-apoptosis effect (61,62). Expression levels of SirT1 were suppressed whereas levels of Cleaved caspase-3 were significantly elevated in the SirT1 siRNA transfection group. These findings suggested that SirT1 activation and ERS are associated with H2O2-induced apoptosis. A DMM mouse model was used to investigate the chondroprotective effects of SFN in vivo. Histological staining showed that SFN significantly alleviated disease progression in mice. This study has certain limitations; for example, flow cytometry analysis revealed the anti-apoptosis effect of SFN on OA, which needs to be further studied in future.

SFN protected chondrocytes against H2O2-induced apoptosis via the SirT1 signaling pathway. Furthermore, SFN was shown to significantly relieve disease progression in a DMM mouse model.
Figure 5. SFN inhibits OA development in a DMM mouse model. (A) Safranin O staining was used to evaluate histomorphometric differences between the sham, DMM and DMM + SFN groups. Magnification, x20 and x40. (B) OARSI and (C) synovitis scores for each cartilage sample. (D) Immunohistochemistry of SIRT1 in a DMM mouse model. (E) Quantification of SIRT1-positive cells in cartilage samples. Data are presented as the mean ± SD (n=15). *P<0.01 vs. sham, **P<0.01 vs. DMM group. SFN, sulforaphane; OA, osteoarthritis; DMM, destabilization of the medial meniscus; OARSI, Osteoarthritis Research Society International; SIRT1, sirtuin 1.
mouse model. Therefore, SFN has a potential value in the prevention and treatment of OA.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors’ contributions

MC and LH wrote the manuscript. YL and LL designed and supervised the study. MC, LH, YL and QD performed the experiments. MC, YL and LL analyzed and interpreted the experimental data. All the authors discussed the results and commented on the manuscript. MC LH, YL and QD confirm the authenticity of all the raw data. All authors read and approved the final manuscript.

Ethics approval and consent to participate

The protocol for animal care and use conformed to The Guide for the Care and Use of Laboratory Animals of the National Institutes of Health and was approved by the Animal Care and Use Committee of Wenzhou Medical University (ethics approval no. wydw2019-0377).

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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