Omentin-1 alleviate interleukin-1β(IL-1β)-induced nucleus pulposus cells senescence

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ABSTRACT
One of the main causes of low back pain (LBP) and degenerative musculoskeletal disorders is intervertebral disc degeneration (IVDD). Inflammation-associated senescence of Human nucleus pulposus cells (HNPCs) plays an essential function in the disease progression of IVDD. Omentin-1 is an adipokine that has been recently reported to have anti-inflammatory potential. In our research, IL-1β was used to simulate the inflammatory environment in the IVDD. We investigated in vitro the effects of Omentin-1 on HNPCs, including the components of senescence, cell cycle and extracellular matrix (ECM) synthesis. The results showed that the addition of Omentin-1 improved IL-1β-induced senescence in HNPCs. G1 phase cell cycle arrest and reduced ECM synthesis in HNPCs. Furthermore, we demonstrated that the effect of Omentin-1 in reducing senescence of HNPCs is dependent on SIRT1. These findings suggest that Omentin-1 plays an important function in protecting HNPCs against senescence and has the potential for IVDD gene target therapy.

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Highlights
• Our study demonstrated the addition of Omentin-1 improved IL-1β-induced senescence in HNPCs for the first time.
• In the present study, we found that the effect of Omentin-1 in reducing senescence of HNPCs is dependent on SIRT1.
• These findings suggest that Omentin-1 plays an important function in protecting HNPCs against senescence and has the potential for IVDD gene target therapy.

1. Introduction
The leading cause of low back pain is intervertebral disc degeneration (IVDD), which places a tremendous burden on global health care systems and socioeconomics [1–3]. Since the specific pathological mechanism of IVDD is not clear, symptomatic treatment by surgical removal of the nucleus pulposus is currently one of the most effective treatments, but many patients still suffer from postoperative low back pain and even reduced quality of life. Therefore, research on the pathogenesis of IVDD is urgent.
It is now widely believed that IVDD is mainly caused by the following factors, including aging, genetics, gender, susceptibility to injury, and environment [4,5]. Age-related factors are most associated with the pathological processes, therefore, as IVDD disease progresses, more and more senescent cells accumulate in the intervertebral disc (IVD) tissue [6,7]. The nucleus pulposus cells (NPCs) in IVD are mainly located in the middle part of IVD and their main function is to produce and stabilize the extracellular matrix (ECM) [8]. It is considered that the progression of IVDD begins with degenerative changes in NPCs, accelerated by the degradation of the ECM [9,10]. During IVDD pathology, NPCs produce excess inflammatory factors that trigger the subsequent pathogenic process of aging, of which the most widely studied inflammatory factors are IL-1β and TNF-α [11,12]. Based on the relevant literature, we selected IL-1β for the induction of senescence in NPCs, simulating the microenvironment of IVDD in vitro [12,13].

Adipokines are active factors derived from white adipose tissue [14]. To date they usually include the following: leptin, adiponectin, vaspin and omentin. According to available literature, they functionally regulate various biological processes such as energy balance, inflammation and bone metabolism [14,15]. Omentin-1 (also known as intelectin-1 because it binds lectin) is an adipokine that was first discovered and named in 2006, so designated because it is found in retinal adipose tissue [16]. Previous studies have shown that Omentin-1 can have anti-inflammatory potential by modulating the immune response of the body [17,18]. In studies of osteoarthritis (OA), it has been well established that Omentin-1 may have potential chondroprotective therapeutic capabilities [19,20]. However, studies on the role of Omentin-1 in IVDD have not been reported.

In the present study, our work focused on exploring the potential of Omentin-1 in protecting HNPCs from senescence and, we preliminarily explored the underlying mechanisms of this protective effect.

2. Materials and methods

2.1 Cell culture and treatment experiment

Human nucleus pulposus cells (HNPCs) and related culture medium (Nucleus Pulposus Cell Medium) were purchased from the Sciencell company (Sciencell, CA, USA) as described previously. HNPCs were incubated at 37°C in a humidified atmosphere of 5% (v/v) CO2. Only HNPCs from 2–3 generations were used in the following experiments. The fully-grown confluent HNPCs were then stimulated with IL-1β (10 ng/mL) for 24 h [21,22]. For omentin-1 co-treatment, 150 and 300 ng/mL concentrations of human recombinant omentin-1(R&D Systems, MN, USA) were added to the cell media for the same period.

2.2 Senescence-associated beta-galactosidase (SA-β-Gal) staining

HNPCs were seeded in 6-well plates, subsequently, different experimental groups were set up and given different treatments. Then, different groups of cells were stained with Sa-β-gal staining kit (Beyotime, shanghai, China), and the proportion of Sa-β-gal staining-positive HNPCs was observed under a light microscope. The positively stained cells were then counted and normalized to the total cell count [19].

2.3 Cell counting kit-8 (CCK8) assay

The Cell Counting Kit 8 (CCK8) assay (Beyotime) was used to detect cell viability according to the manufacturer’s instruction. Briefly, HNPCs were seeded in 96-well plates at 2000 cells per well and then given different treatments. Subsequently, CCK-8 reagent was added to each well at 0, 24, and 48 h timepoints and incubated for 2 h at 37°C. The optical density (OD) at 450 nm wavelength was measured using a microplate reader (ELx800, BioTek, USA).

2.4 Quantitative real-time PCR (qRT-PCR)

Total cellular RNA was isolated with Trizol reagent (Invitrogen, Thermo Fisher). Purified RNA was reverse transcribed using PrimeScript RT Master
Mix (Perfect Real Time, Takara, California, USA). The primer sequences are presented in Additional file 1: Table S1. The complementary DNA (cDNA) produced by the reverse transcription was amplified by using SYBR Premix Ex Taq (Takara) following the manufacturer’s instructions through a 7500 Real-Time PCR system (Applied Biosystems, Foster City, California, USA). GAPDH was used as an internal standard to calculate gene expression using the 2^\(-\Delta\Delta C_t\) method [23].

### 2.5 Western blot analysis

After the indicated treatment, cell samples were lysed using cell lysis buffer (Cell Signaling, USA) containing protease inhibitor cocktail (Roche, USA). Protein concentrations were determined using the bicinchoninic acid (BCA) method. Equal amount of the extracted proteins was separated by 8%–12% sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) and transferred to polyvinylidene fluoride (PVDF) membranes via electroblotting for 2 h. Blots were blocked in 5% nonfat dry milk for 1 h at room temperature and then incubated at 4°C overnight with primary antibodies against anti-Omentin-1 (diluted 1:1000, Abcam, catalog number: ab252927), anti-P16 (diluted 1:1000, Abcam, catalog number: ab32441), anti-MMP13 (diluted 1:250, Abcam, catalog number: ab41037), anti-P16 (diluted 1:2000; Abcam, catalog number: ab39012), anti-SIRT1 antibody (diluted 1:2000; Abcam, catalog number: ab34712), anti-Aggrecan antibody (diluted 1:1000, Abcam, catalog number: ab36861), anti-ADAMTS-5 antibody (diluted 1:250, Abcam, catalog number: ab36861), anti-MMP13 antibody (diluted 1:250, Abcam, catalog number: ab36861), anti-SIRT1 antibody (diluted 1:2000; Abcam, catalog number: ab36861), anti-SIRT1 antibody (diluted 1:1000, Abcam, catalog number: ab36861), anti-SIRT1 antibody (diluted 1:500; Abcam, catalog number: ab36861), anti-SIRT1 antibody (diluted 1:250; Abcam, catalog number: ab36861), anti-SIRT1 antibody (diluted 1:1000; Abcam, catalog number: ab36861), anti-SIRT1 antibody (diluted 1:200; Abcam, catalog number: ab36861), and anti-SIRT1 antibody (diluted 1:1000, Cell Signaling Technology, catalog number: #9282), anti-Col II antibody (diluted 1:1000, Abcam, catalog number: ab36861). Membranes were then washed 3 times with TBS containing 0.05% Tween-20 (TBS-T buffer) and then incubated with horseradish peroxidase-conjugated secondary antibodies at 37°C for 1 h. Finally, the blots were developed with an enhanced chemiluminescence (ECL) kit (Pierce Biotechnology, USA).

### 2.6 Cell cycle analysis

HNPCs were seeded in six-well plates and grown to 70% to 80% confluence. After serum starvation for eight hours, HNPCs were incubated with different treatments. Subsequently, HNPCs were digested with trypsin (0.25% without EDTA; Gibco, Thermo Fisher) and centrifuged to collect the cell pellets. After fixation with 70% ethanol and staining with propidium iodide dye (50 μg/ml, Beyotime) for 30 minutes, HNPCs were subjected to flow cytometry analysis.

### 2.7 Transfection

We performed a siRNA-based method to knock down the expression of SIRT-1 in HNPCs. The specific siRNA sequences against SIRT1 were designed and synthesized by GenePharma (Shanghai China) with the forward sequence: GCAAUAGGCCCUCUUAAAUATT and the reverse sequence: UAAUAAGGCCUAUUGCTT. To transfect the cells, we administered a mixture of Lipofectamine 2000 Reagent (Invitrogen) and SIRT-1 oligomers to 50% confluent HNPCs. Then the transfected cells were grown for another 48–72 hours.

### 2.8 Statistical analysis

All the experimental data in this study are representative of three independent experiments performed and are presented as mean ± standard deviation (SD). Student t-test was used for comparisons between two groups and one-way ANOVA test was used between more than two groups. A p-value <0.05 was considered statistically significant.

### 3. Results

We hypothesized that Omentin-1 protects HNPCs from aging by targeting and regulating SIRT1. In our research, IL-1β was used to simulate the inflammatory environment in the IVDD. We investigated in vitro the effects of Omentin-1 on HNPCs, including the components of senescence, cell cycle and extracellular matrix (ECM) synthesis. Furthermore, we demonstrated that the effect of Omentin-1 in reducing senescence of HNPCs is dependent on SIRT1.
3.1 Omentin-1 expression was upregulated in senescent HNPCs induced by IL-1β

We isolated and cultured HNPCs as before and took the second and third generation cells for subsequent experiments. To evaluate the expression of Omentin-1 in HNPCs treated with IL-1β, we treated HNPCs with different doses of IL-1β (0 ng/ml, 10 ng/ml, 20 ng/ml, and 50 ng/ml) at different time points. As shown in Figure 1(a), the cell viability of IL-1β-treated HNPCs was lower than that in the control group at 24 and 48 hours. Consequently, we chose the time point for IL-1β stimulation of HNPCs to be 24 hours. Moreover, there was a quantitative relationship between the expression level of Omentin-1 and the concentration of IL-1β in HNPCs (Figure 1(b-d)). Meanwhile, the literature has demonstrated that IL-1β can induce premature senescence in HNPCs [12,13]. Taken together, we used IL-1β (10 ng/mL for 24 h) in follow-up experiments to induce senescence in HNPCs.

![Figure 1](image)

Figure 1. Omentin-1 expression was upregulated in senescent HNPCs induced by IL-1β. a. Cell viability of IL-1β-induced HNPCs was assessed by CCK8 assay (n = 3/group). b. Omentin-1 mRNA expression in HNPCs stimulated by different concentrations of IL-1β was assessed by qRT-PCR (n = 3/group). c, d. Omentin-1 protein expression in HNPCs stimulated by different concentrations of IL-1β was assessed by Western blot analysis (n = 3/group). *P < 0.05 between 0ng/ml and 10 ng/ml groups, **P < 0.05 between 0 ng/ml and 20 ng/ml groups, †P < 0.05 between 0 ng/ml and 50 ng/ml groups.

3.2 Omentin-1 alleviated IL-1β-stimulated senescence in HNPCs

To investigate whether Omentin-1 plays a biological role in affecting IL-1β-induced senescence in HNPCs. We performed this part of the experiment. In the first step, Sa-β-gal staining was conducted, as shown in Figure 2 (a, b), the positive rate of Sa-β-gal staining showed an increase in the IL-1β group compared to the control group. However, the addition of Omentin-1 diminished the positive rate of Sa-β-gal staining increased by IL-1β, and the higher dose of Omentin-1 significantly diminished the positive rate of Sa-β-gal staining increased by IL-1β.

Subsequently, we determined the mRNA and protein levels of P16 and P53 (cellular senescence markers) in HNPCs of each experimental group. As shown in Figure 2(c, d), the mRNA levels of p16 and p53 were upregulated in the IL-1β group compared to the control group. However, the addition of Omentin-1 diminished
the mRNA levels of p16 and p53 increased by IL-1β, and the higher dose of Omentin-1 significantly diminished the mRNA levels of p16 and p53 increased by IL-1β. Meanwhile, we verified the Omentin-1 inhibitory function on P16 and P53 at the protein level. (Figure 2(e-g)).
In summary, Omentin-1 protected against senescence induced by IL-1β stimulation of HNPCs.

3.3 Omentin-1 promoted extracellular matrix synthesis in HNPCs

To explore the in-depth potential mechanism by which Omentin-1 can alleviate the senescence of HNPCs in vitro. We performed this part of the experiment. Firstly, we did cell cycle and cell viability assays, as shown in Figure 3(a,b). Compared with the control group, the G1 phase cell cycle was prolonged along with reduced cell viability after 24 hours of intervention with IL-1β. However, the addition of Omentin-1 shortened the cell cycle in the G1 phase that was prolonged by IL-1β and improved the cell viability that was reduced by IL-1β. At the same time, with higher dose of Omentin-1 addition, this effect of improvement was more obvious.

Subsequently, we determined the mRNA and protein levels of Colleagen II, Aggrecan (synthesis-related markers of ECM); MMP 13 and ADAMT5 (degradation-related markers of ECM) in HNPCs of each experimental group, respectively. In this experiment, it was found that IL-1β could down-regulate the expression of synthesis-related markers at the mRNA levels compared to the control group. However, the addition of Omentin-1 up-regulated the mRNA levels of synthesis-related markers attenuated by IL-1β, and the higher dose of Omentin-1 significantly up-regulated the mRNA levels of synthesis-related markers attenuated by IL-1β. The results of Western blot analysis further confirmed the results of qRT-PCR at the protein level (Figure 3(d,e)).

Taken together, Omentin-1 promoted the synthesis of extracellular matrix in HNPCs.

3.4 Omentin-1 protects against IL-1β-induced senescence in HNPCs by targeting SIRT1

SIRT1 is one of the most essential senescence genes, so we investigated in HNPCs whether Omentin-1 would influence the expression of SIRT1. Our experiments show that IL-1β mediated a decrease in mRNA and protein levels of SIRT1 in HNPCs and that the addition of Omentin-1 effectively restored this alteration. As shown in Figure 4(a), the 24-h IL-1β intervention reduced SIRT1 mRNA level compared with the control group. However, the addition of Omentin-1 up-regulated SIRT1 mRNA levels. Also, the effect of this rescuing became more pronounced as the dose of Omentin-1 addition was increased. We subsequently obtained the same validation at the protein level (Figure 4(b, c)).

To investigate whether variation in SIRT1 expression affects the role of Omentin-1 in protecting against senescence in HNPCs. We performed this part of the experiment. Firstly, we transfected SIRT1-specific RNAi oligonucleotides into HNPCs and verified the successful construction of knockdown cell models by qRT-PCR and Western blot (Additional file 2: Fig. S1). By examining the mRNA and protein levels of P16 and P53, we confirmed that with the knockdown of SIRT1 in HNPCs, the inhibitory effect of Omentin-1 on these two proteins was attenuated (Figure 4(d-f)). As shown in Figure 4(g, h), the effect of Omentin-1 in reducing the proportion of positive IL-1β-induced SA-β-Gal staining was attenuated with the knockdown of SIRT1 in HNPCs.

In summary, Omentin-1 plays a role in protecting HNPCs from IL-1β-induced senescence by targeting SIRT1.

4. Discussion

Omentin-1 is a novel adipokine derived mainly from visceral adipose tissue [24]. Omentin-1 was found to be a secreted glycoprotein and a lactoferrin-binding protein that binds to galactofuranosyl residues on microorganisms [25]. Most studies point to the anti-inflammatory ability of Omentin-1 to counteract chronic inflammatory processes. For example, Omentin-1 exhibited anti-inflammatory and anti-atherosclerotic properties in obese individuals [26]. In addition, this adipokine was negatively associated with the progression of two diseases, inflammatory bowel disease and metabolic syndrome [27,28]. There are also studies on Omentin-1 in the field of OA in degenerative skeletal disorders. Li et al [20]. found that Omentin-1 was able to suppress IL-1β-induced degradation of type II collagen and proteoglycan in human chondrocytes, demonstrating the inhibitory role of Omentin-1 in inflammation. Chai et al [19] identified that the adipokine Omentin-1
can play a role in protecting chondrocytes against senescence. These findings all suggest that Omentin-1 may have chondroprotective therapeutic potential.

In recent years, there have also been a few studies related to adipokines in IVDD. Regarding leptin, current studies reveal its important role in the pathological development of IVDD, which
enhances intervertebral disc cells proliferation [29], cytoskeletal remodeling [30–32], and production of pro-inflammatory cytokines [33]. In addition, it has been found that adiponectin may protect against IVDD by inhibiting the expression of pro-inflammatory mediators [34,35]. In our study, we found that the adipokine Omentin-1 plays an important role in protecting HNPCs from IL-1β-induced senescence and could be a genetic target for the treatment of IVDD.
The etiology of IVDD is associated with many factors, among which the accumulation of senescent NPCs provides new insights into the pathogenesis of IVDD [13,36]. On the one hand, the senescence of NPCs is followed by a lack of new cell production and a gradual decrease in the number of functional cells. On the other hand, as senescence progresses in NPCs, the microenvironment of the disc is altered. The final outcome develops as a cellular senescence-associated phenotype in which pro-inflammatory factors are overexpressed, ECM synthesis is reduced, and growth factors and chemokines are downregulated [6,37,38]. The main pathological features of IVDD are increased production of degradative enzymes and decreased synthesis of ECM [39]. Therefore, we examined the synthesis-related markers Colleagen II, Aggrecan, and degradation-related markers MMP 13 and ADAMT5 in our experiments [40].

Silent mating-type information regulator 2 homolog-1 (SIRT1), a nicotinamide-dependent class 3 histone deacetylase, is essential for cell viability and extends the lifespan of species ranging from yeast to mammals [41]. Increasingly, studies on the important role of SIRT1 in diseases associated with aging are gradually [42–46]. It has been reported in the literature that SIRT1 is associated with the modulation of several cellular processes, namely cellular senescence, cell cycle and metabolism [41,47,48]. The significant role of SIRT1 in the pathogenesis of two skeletal degenerative diseases, OA and IVDD, has been demonstrated in the literature [49,50]. Therefore, in the present experiments, we hypothesized that Omentin-1 exerts a protective effect against IL-1β-induced senescence in NPCs through the targeted regulation of SIRT1. Literature studies have verified that SIRT1 can arrest NPC senescence and death through NF-κB or PI3K/Akt signaling pathways [51,52]. Meanwhile, a recent study reported that after IL-1β exposure, deoxycine exerts antioxidant stress effects on HNPCs by affecting NF-κB, P65 [53]. Therefore, we can continue to conduct in-depth research in related directions.

5. Conclusion

In conclusion, our study identified for the first time the important function of Omentin-1 in protecting HNPCs from IL-1β-induced aging, while our study explored the underlying molecular mechanisms of Omentin-1 in inflammation-associated senescence. In the future, Omentin-1 may have potential for IVDD gene targeting therapy.

Data availability statement

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

Disclosure statement

No potential conflict of interest was reported by the author(s).

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Author contributions

Xin Huang designed the research and wrote the manuscript. Changhong Chen contributed to the literature search. Yaofei Chen and Jun Xu performed the experiments. Lin Liu reviewed and edited the article. All authors read and approved the article.

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