

Isovitexin Depresses Osteoarthritis Progression via the Nrf2/NF- κ B Pathway: An in vitro Study

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Purpose: Osteoarthritis (OA) is a multifactorial joint disease and inflammatory processes contribute to joint destruction. Isovitexin (IVX) is a flavone component found in passion flower, Cannabis and, and the palm that is known for its anti-inflammatory properties.

Materials and Methods: This study investigated in vitro the role and underlying mechanism used by IVX in its regulation of OA development. Effects of IVX on the viability of chondrocytes were measured by CCK-8 assays. The phenotypes of extracellular matrix (ECM) degeneration and inflammation were measured by qPCR, Western blot, and ELISA; and NF- κ B pathway was detected by immunofluorescence and Western blot. Molecular docking was applied to predict the interacting protein of IVX, while Nrf2 was knocked down by siRNAs to confirm its role.

Results: We demonstrated that IVX suppressed ECM degeneration and suppressed pro-inflammatory factors in IL-1 β -treated chondrocytes. Additionally, IVX impact on NF- κ B signaling in IL-1 β -exposed chondrocytic cells; Mechanistically, it was also demonstrated in molecular docking and knock down studies that IVX might bind to Nrf2 to suppress NF- κ B pathway.

Conclusion: Our data suggest that IVX halts OA disease advancement through the Nrf2/NF- κ B axis, suggesting a possibility of IVX as a target for OA therapy.

Keywords: osteoarthritis, isovitexin, extracellular matrix, NF- κ B, Nrf2

Introduction

In general, Osteoarthritis (OA), is known as a chronic, incurable, as well as degenerating illness, which is marked with tremendous pain, limited mobility and disability.^{1,2} With a large aging population, increased incidences of obesity, and occasional injury, OA has become prevalent around the world. Thus far, around 27 million Americans and over 250 million people globally are affected by this disease.^{3,4} Advances in biological tissue engineering have attempted to ease treatment and rehabilitation of OA.⁵ But more work needs to be done to achieve success. Stem cell therapy also is lagging behind in its ability to relieve OA.^{6,7} Additionally, joint replacement surgery is costly.⁸ Given these complications, it is urgent and necessary to develop new drugs with high efficacy to relieve OA symptoms. This study, therefore, examined the role and efficacy of IVX in managing OA.

Inflammatory factors are known to control OA development.⁹ Among the chief cytokines, interleukin-1 β (IL-1 β) was reported to be highly expressed in OA animals and humans, and the level of IL-1 β positively correlated with the progression of disease.^{10,11} As expected, high IL-1 β levels increased the production of tumor necrosis factor- α (TNF- α), interleukin-6 (IL-6), cytochrome C oxidase

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subunit 2 (COX-2), and inducible nitric oxide synthase 2 (iNOS), which severely impaired the metabolic turnover of extracellular matrix (ECM) inside knee joints.^{12,13} Conversely, IL-1 β or IL-1 β receptor inhibition were shown to have a therapeutic effect on OA.^{14,15}

Moreover, nuclear factor kappa-light-chain-enhancer or NF- κ B signaling modulates inflammation and IL-1 β -mediated catabolism in chondrocytes.¹⁶ Under conditions of IL-1 β stimulation, a phosphate group is added to the NF- κ B inhibitor alpha (I κ B α) which becomes marked for proteasomal degradation. This allows the p65 protein to be liberated from its inhibitory complex and travel to the nucleus,^{16,17} where it activates target-specific pro-inflammatory or catabolic downstream genes.^{18,19}

Nrf2, an essential transcription factor, is primarily involved in cell homeostasis. In fact, Nrf2 was shown to activate antioxidant, anti-apoptotic, and anti-inflammatory pathways to maintain homeostasis.²⁰ Multiple studies have also demonstrated an important interaction involving NF- κ B and Nrf2 needed to maintain cell homeostasis and improve inflammation.^{21,22} Interestingly, Nrf2 stimulation was also shown to depress the IL-1 β -mediated stimulation of NF- κ B in human chondrocytic cells.²³ Furthermore, an imbalance in the levels of Nrf2 and NF- κ B was reported to cause numerous diseases, including neurodegeneration.²⁴ Based on these reports, the Nrf2/NF- κ B pathway is deemed crucial for investigations into promising targets for OA therapy. Heme oxygenase 1 (HO-1) is considered one of the main effectors of Nrf2-dependent cell responses.²⁵ Moreover, Nrf2/HO-1 association has been widely reported, especially in OA.^{26,27}

Isovitexin (otherwise called homovitexin or saponaretin) is a class of flavonoid, commonly found in the passion flower, Cannabis and, and the palm.²⁸ Multiple studies have demonstrated an anti-inflammatory role of IVX.^{29,30} Moreover, IVX was also reported to protect cells from inflammation via its regulation of Nrf2 signaling pathway.³¹ Given what is known about IVX, it is possible that its anti-inflammatory action may serve as a protective measure against OA, especially through the Nrf2/NF- κ B pathway.

This paper aims to assess the protective aspect, if any, of IVX on OA. OA inflammation is activated via Nrf2/NF- κ B signaling. Therefore, the IVX impact on the Nrf2/NF- κ B axis will also be evaluated. The data presented here may aid in the advancements of future high efficacy OA therapy.

Materials and Methods

Statement of Ethics

The experimental mice are acquired from the Animal Experimental Research Center of Zhejiang Chinese Medical University (SYXK 2021-0012). The experimental scheme and ethical application are approved by the Animal Welfare and Ethics Review Committee of Zhejiang Chinese Medical University (IACUC-20,190,826-09), all operations are performed strictly in accordance with the guidelines of this Committee.

Reagents and Antibodies

Reagents used and the companies they were purchased from are listed as follows: IVX (purity \geq 98%, Meilun Biotechnology, China and MedChemExpress LLC, USA), recombinant human IL-1 β (PeproTech, USA), Collagenase Type II and dimethylsulfoxide (DMSO, Sigma-Aldrich, USA), Cell Counting Kit-8 (CCK-8, Dojindo, Japan). The primary (1 $^{\circ}$) antibodies used and the companies they were obtained from are listed below: against iNOS, AggreCan, ADAMTS-5, and collagen II (COL II, Abcam, USA), against GAPDH, MMP-3, COX-2, and MMP-13 (Proteintech, China), and against I κ B α , p-I κ B α , p65, p-p65, Nrf2, HO1 (Cell Signaling Technology, USA). The secondary (2 $^{\circ}$) antibodies used and their companies are provided below: goat anti-mouse and anti-rabbit IgG-HRP (Bioss, China), Alexa Fluor[®]488-labelled as well as Alexa Fluor[®]594-labelled goat anti-rabbit IgG (H+L) (Jackson ImmunoResearch, USA). The cell culture medium came from Gibco (Grand Island, USA), and the 4',6-diamidino-2-phenylindole (DAPI) came from Beyotime (Shanghai, China).

Isolation and Cultivation of Primary Mice Chondrocytic Cells

To isolate primary mouse chondrocytic cells, cartilages of knee belonged to 30 immature C57BL/6 mice were diced into cubes, approximately 1 mm³, before Wash with PBS, and digested in 0.25% type II collagenase at 37°C for 2 hr. The resulting isolated cells were grown in DMEM/F12 containing 10% FBS (Gibco) and 1% penicillin-streptomycin solution in a humidified environment at 5% CO₂ at 37°C. The media was replaced once in 2 days and chondrocytes from the 2nd passage were utilized for further investigation.

Experimental Design and Cell Transfections

The chondrocytes were seeded in a six-well plate and cultured for 24 hr until they were 50–70% confluent. Then, the cells were administered 10 ng/mL IL-1 β , together with differing concentrations of IVX, namely 0, 25, 50 μ g/mL for 24 hr. Subsequently, we employed PCR, Western blot analysis, immunofluorescence, or ELISA assays.

As for Cell transfections, the Nrf2-siRNA and Control-siRNA (negative control) were purchased from Santa Cruz Biotechnology (Ribo, Guangzhou, China). Then, the cells were transfected with 50 nM siRNA using Lipofectamine 2000 siRNA transfection reagent (Thermo Fisher) for 24 hr. And the validity of siRNA was verified by Western blot in Figure 1A.

Cell Viability Assay

The Cell Counting Kit-8 (CCK-8; Dojindo Co, Japan) was employed for the assessment of cellular survival. Briefly, chondrocytic cells (50,000 cells/cm²) were grown in 96-well plates for 24 hr before a 24 hr treatment with IL-1 β and IVX, as described previously. The cells next underwent PBS-wash, and exposure to 100 μ L of DMEM/F12 and 10 μ L of CCK-8, in 37°C for 1 hr. Lastly, absorbance at 450 nm was recorded by using a microplate reader (Thermo, USA).

Real-Time PCR

TRIzol (Invitrogen, USA) was employed for total RNA isolation, following manufacturer guidelines. In the next step, a total amount of 1000 ng RNA was used for reverse

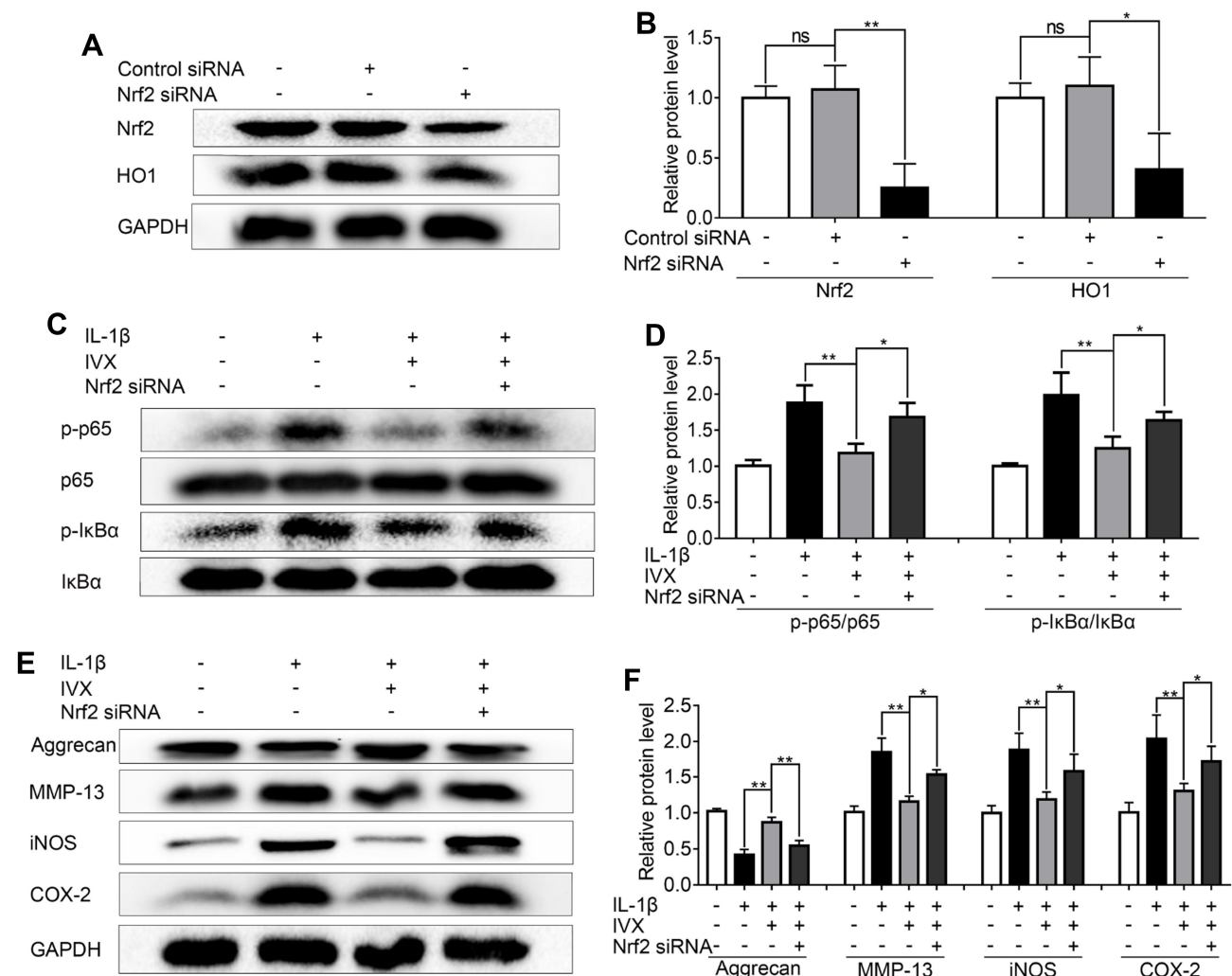


Figure 1 Nrf2 deficiency compromises IVX action on IL-1 β -exposed chondrocytes. Nrf2 was knocked down by siRNA in IL-1 β -exposed chondrocytes (**A**, **B**) Evaluation of Nrf2 and HO1 protein expression, using Western blot. Protein quantification was done using Image J. (**C**, **D**) Assessment of phosphorylated and unphosphorylated forms of I κ B α and p65, using Western blot. Protein quantification was performed using Image J software. (**E**, **F**) Determination of the Nrf2 signaling-related protein expression. Data is average \pm SD of 3 separate examinations. * p < 0.05, ** p < 0.01.

Abbreviation: Ns, not significant.

transcription to form cDNA (MBI Fermentas, Germany) with the kit of PrimeScript-RT reagent (Japan), as well as the System of CFX96 Real-Time PCR (Bio-Rad Laboratories, USA). cDNA amplification was done by SYBR Premix Ex Taq (Takara, Japan) in the System of CFX96 Real-Time PCR (Bio-Rad Laboratories, USA). The normalization of the cycle threshold (Ct) of examined transcripts was carried out to the endogenous control (GAPDH). Finally, the relative levels of relevant genes were measured with the $2^{-\Delta\Delta C_t}$ method.³² Lastly, the primers employed to amplify inflammatory transcripts were described previously.¹⁹

Western Blot Analysis

Chondrocyte lysis was done with radio immunoprecipitation assay (RIPA) (Beyotime, China) buffer and 1 mM phenylmethanesulfonyl fluoride (PMSF) (Beyotime, China), before centrifugation at 4°C and 12,000 rpm for 15 min. The levels of protein were quantified using the BCA protein assessment kit (Beyotime, China). Next, 40 µg of the protein was separated on an 8–12% (w/v) sodium dodecyl sulfate polyacrylamide gel (Beijing Solarbio science and technology co., Ltd, China) electrophoresis and blotted onto the membranes of polyvinylidene fluoride (Bio-Rad, Hercules, USA). Moreover, the blocking was performed with 5% milk with zero fat within 2 hr, then 1° antibody incubation occurred at 4°C overnight (O/N). The 1° antibodies used were against iNOS (1:1000), Aggrecan (1:800), COX-2 (1:1000), ADAMTS-5 (1:1000), p65 (1:1000), pp65 (1:800), MMP-3 (1:1000), MMP-13 (1:1000), IκB-α (1:1000), p-IκB-α (1:800), Nrf2 (1:1000), HO1 (1:1000), or GAPDH (1:5000). The following day, after wash 3 times with PBS, the corresponding 2° antibodies were introduced for 2 hr at room temperature (RT) before Tris-buffered saline containing Tween® 20-wash 3X, followed by visualization with Plus reagent (Invitrogen, NY, USA) and ChemiDoc XRS+ Gel Imaging System (Bio-Rad, CA, USA). Lastly, the band intensities were meticulously explored using Image J 2.1 (Bethesda, USA), standardized by GAPDH.

Immunofluorescence

Chondrocytic cells were grown and treated in the plates including 6 wells as mentioned earlier. Next, they underwent ice-cold PBS-wash before fixation with 4% (v/v) paraformaldehyde for 15 min, permeabilization with 0.1% Triton X-100 diluted in PBS for 10 min, blocked with 5% bovine serum albumin for 1 hr at 37°C, exposed to 1° antibodies against COL II (1:200) p65 (1:200) or Nrf2 (1:200) at 4°C O/N, exposed to Alexa Fluor® 594-conjugated or Alexa Fluor® 488-

labelled 2° antibodies (1:300) for 1 hr at RT, and stained by using 4',6-diamidino-2-phenylindole (DAPI) for 5 min. Next, 5 random fields per slide were selected for analysis using an ECLIPSE Ti microscope of Nikon (Nikon, Japan). Lastly, scientists blinded to the sample groups were employed to quantify fluorescence intensity with Image J 2.1 (Bethesda, USA). Protein expression was quantified by integrated optical density (IOD) with the Image J 2.1 image analysis system.

ELISA Assay

Culture supernatants were collected and stored at −20°C until analysis. The concentration of TNF-α, IL-6, and PGE2 in cell culture supernatants was measured using ELISA kits (R&D Systems, Minneapolis, MN, USA) according to the manufacturer's instructions.

Molecular Modelling

The IVX 3-D molecular structure was constructed utilizing Discovery Studio 2016 and minimized with the CHARMM force field. Next, the Keap1-Nrf2 structural composite (PDB ID: 3WN7)³³ was retrieved from the RCSB Protein Data Bank (<https://www.rcsb.org/>) and opened up in Discovery Studio 2016. Following this, the Keap1-Nrf2 composite structure was manually altered by eliminating water and introducing hydrogen before the partially flexible CDOCKER program was utilized to localize special binding locations and the receptor radius. Finally, the molecular docking data were evaluated to assess CDOCKER energy scores, interaction site, and interaction force types.³⁴

Statistical Analysis

Data presented as average ± SD of 3 separate examinations. Data analysis was executed with GraphPad Prism (USA) (one-way analysis of variance (ANOVA) as well as Tukey's post hoc test). Statistically significant data was given to p value of <0.05. The variations were deemed statistically significant at p < 0.05.

Results

Evaluation of IVX-Mediated Cellular Toxicity and Protection in Chondrocytic Cells

The IVX chemical composition is illustrated in Figure 2A. To identify if IVX exerts cytotoxicity, chondrocytic cells were given increasing concentrations of IVX, namely, 0, 12, 25, 50, 100, 200, 400 µg/mL, for 24 or 48 hr before cell survival measurement, as described above. Based on

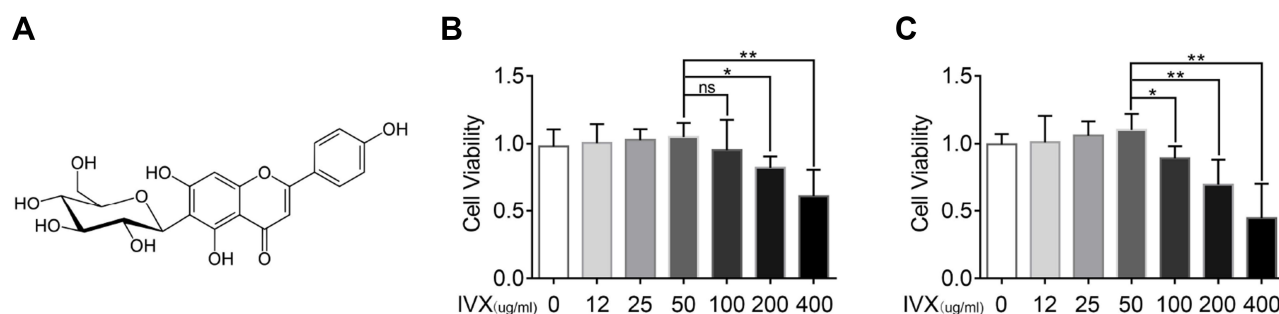


Figure 2 Evaluation of IVX-induced cellular toxicity and protection in IL-1 β -exposed chondrocytic cells. **(A)** Chemical composition of IVX. **(B, C)** Assessment of IVX chondrocytic cells cytotoxicity, using differing concentrations of IVX for 24 hr and 48 hr. Data = average \pm SD of 3 separate examinations. * $p < 0.05$, ** $p < 0.01$.

Abbreviation: Ns, not significant.

our result, concentrations up to 50 μ M produced no observable cytotoxicity. However, IVX concentrations >100 μ M markedly diminished cell survival (Figure 2B and C). Therefore, in further experimentation, IVX concentrations of 0, 25, and 50 μ M were used to analyze its effect on IL-1 β -treated chondrocytes.

IVX Impact on the Anabolic and Catabolic Nature of ECM in IL-1 β -Exposed Chondrocytes

In order to determine whether IVX preserves ECM from IL-1 β -mediated anabolic events, anabolic proteins were evaluated in IVX- and IL-1 β -treated cells. We demonstrated a strong protection in the levels of Aggrecan and COL II in the IVX-treated cells, as opposed to cells receiving IL-1 β alone (Figure 3A and E). We, further, tested IVX's effect on the ECM catabolic events and showed a strong suppression of ECM catabolic proteins, MMP-3, MMP-13, and ADAMTS-5 in the IVX-treated cells, relative to IL-alone (Figure 3A–D).

IVX Modulation of Pro-Inflammatory Factors in IL-1 β -Exposed Chondrocytes

Since pro-inflammatory factors play a major role in OA, we next examined whether IVX modulates the expression of members of the pro-inflammatory pathway. To do this, we employed PCR and ELISA assays to demonstrate significant increases in TNF- α and IL-6 levels upon IL-1 β treatment, which were attenuated with exposure to IVX (Figure 4A and E). Similarly, using PCR and Western blot analysis, we revealed dramatic IL-1 β -stimulated rise in iNOS and COX-2 generation, which were suppressed in the presence of IVX (Figure 4B–F). Collectively, these

results suggest that IVX represses pro-inflammatory pathways.

IVX Impact on NF- κ B Signaling in IL-1 β -Exposed Chondrocytic Cells

Given that IVX strongly suppressed pro-inflammatory factors, we next investigated whether the cytoprotective actions of IVX involved the NF- κ B signaling pathway. Using immunofluorescence, we demonstrated p65 translocation to the nucleus in IL-1 β -exposed chondrocytic cells. Comparatively, IVX exposed cells exhibited more p65 localization in the cytoplasm (Figure 5A). Moreover, Western blot analysis revealed a larger population of phosphorylated p65 and I κ B α in IL-1 β -exposed chondrocytic cells, as opposed to cells with no treatment, suggesting stimulation of the NF- κ B signaling. Comparatively, the IVX-treated chondrocytes displayed remarkably low quantities of phosphorylated p65 and I κ B α (Figure 5B–D). We further isolated chondrocyte nucleus and examined p65 protein expression in the nucleus using Western blot. Similar to our immunofluorescence data, IVX suppressed p65 presence in the nucleus (Figure 5B and E). In all, these results suggest a strong NF- κ B signaling repression in the IVX-exposed chondrocytic cells.

Molecular Docking of IVX and Nrf2

Multiple upstream regulators are known to modulate NF- κ B signaling. Among them, Nrf2 was shown to be a major regulator of this pathway²⁶ and serve as a potential target for OA therapy.³⁵ To identify possible interaction between IVX and protein(s) involved with the Nrf2/NF- κ B pathway, we conducted molecular docking analysis. Our analysis predicted that the Nrf2 protein contains a binding site for IVX (Figure 6A and B). In fact, we discovered high interaction energy between IVX and Nrf2, measuring at

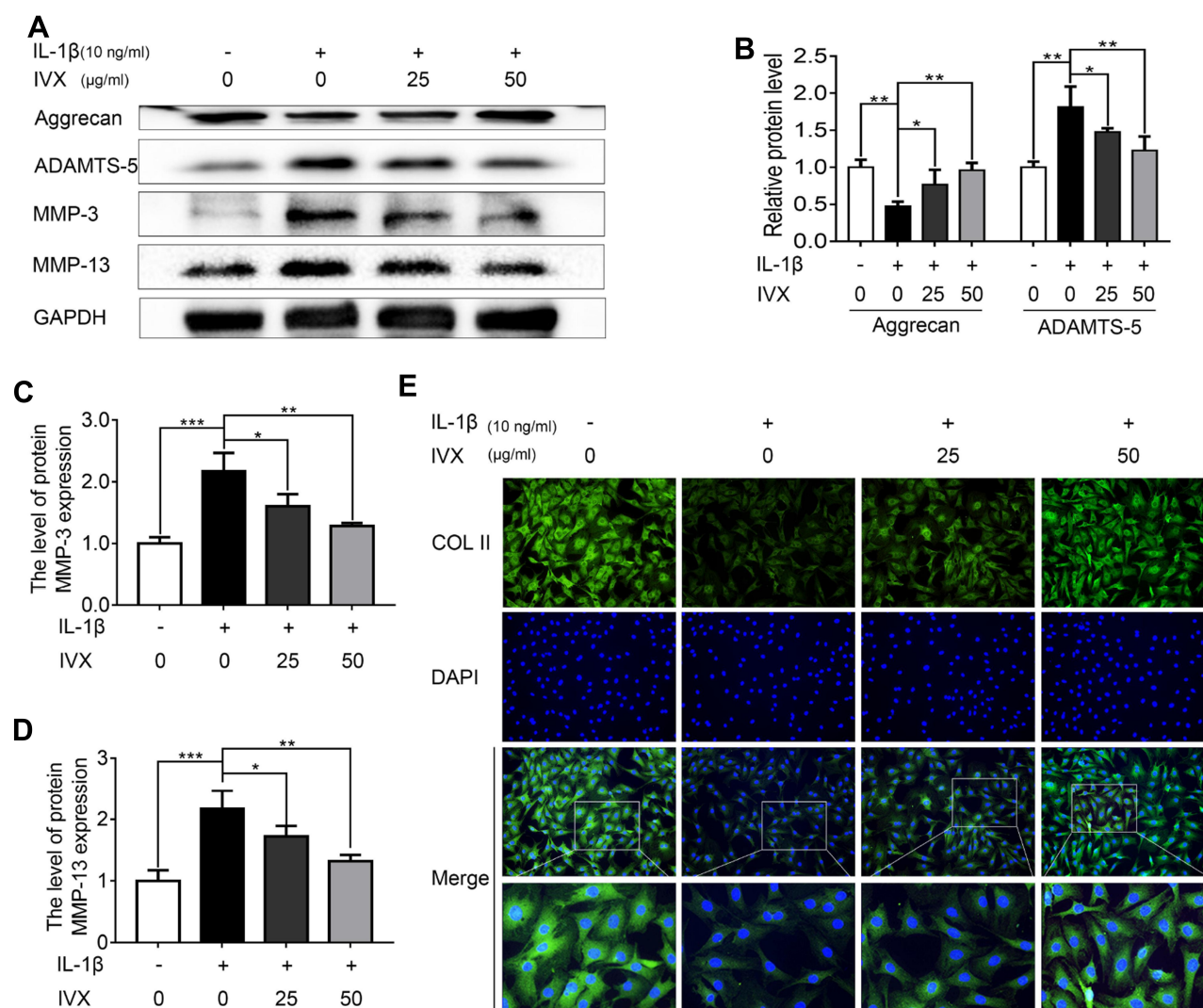


Figure 3 IVX impact on ECM anabolic and catabolic actions in IL-1 β -exposed chondrocytic cells, in presence or absence of IVX. **(A)** Evaluation of ECM catabolic and anabolic protein expression in IL-1 β - and IVX-treated chondrocytes, relative to control. **(B–D)** Image J employed quantification of protein expression. **(E)** Detection of COL II by fluorescence microscopy, in conjunction with nuclear DAPI stain (scale bar: 50 μ m). Data is average \pm SD of 3 separate examinations. * p < 0.05, ** p < 0.01, *** p < 0.001.

42.95 kcal/mol (Figure 6C). Additionally, according to the 2D-binding model, Nrf2 interacted with IVX via van der Waals, hydrogen bonds, and pi-alkyl interaction (Figure 6D and E).

IVX Modulates NF- κ B Signaling via Nrf2

To validate the involvement of the Nrf2 pathway in the IVX-mediated cytoprotection of IL-1 β -exposed chondrocytes, we first explored Nrf2 localization using immunofluorescence. We demonstrated significant Nrf2 translocation to the nucleus, upon IVX treatment (Figure 7A). Similarly, Nrf2 levels in the nuclear fractions of chondrocytes were significantly upregulated in IVX-treated chondrocytes, relative to IL-1 β alone. Moreover, Nrf2-targeted HO1 protein expression in whole cells extracts was also elevated, relative to IL-1 β -exposed

cells (Figure 7B, D and E). These data were also verified in terms of HO1 transcripts levels, which remained remarkably high in IVX-exposed chondrocytic cells versus IL-1 β alone (Figure 7C). Next, we generated Nrf2 deficiency in the isolated chondrocytes, using siRNA. Using Western blot, we revealed low expression of Nrf2 and HO1 proteins upon Nrf2 deficiency, relative to the negative control (Figure 1A and B). Moreover, Nrf2 deficiency produced more phosphorylated forms of p65 and I κ B α proteins, as compared to the control (Figure 1C and D). In all, our data indicates that IVX modulates NF- κ B pathway through its regulation of Nrf2. Furthermore, we showed that IVX exposure raised the amount of Aggrecan protein, as opposed to IL-1 β alone (Figure 1E). Conversely, Aggrecan protein levels were strongly suppressed under conditions of Nrf2 deficiency, while MMP13 protein

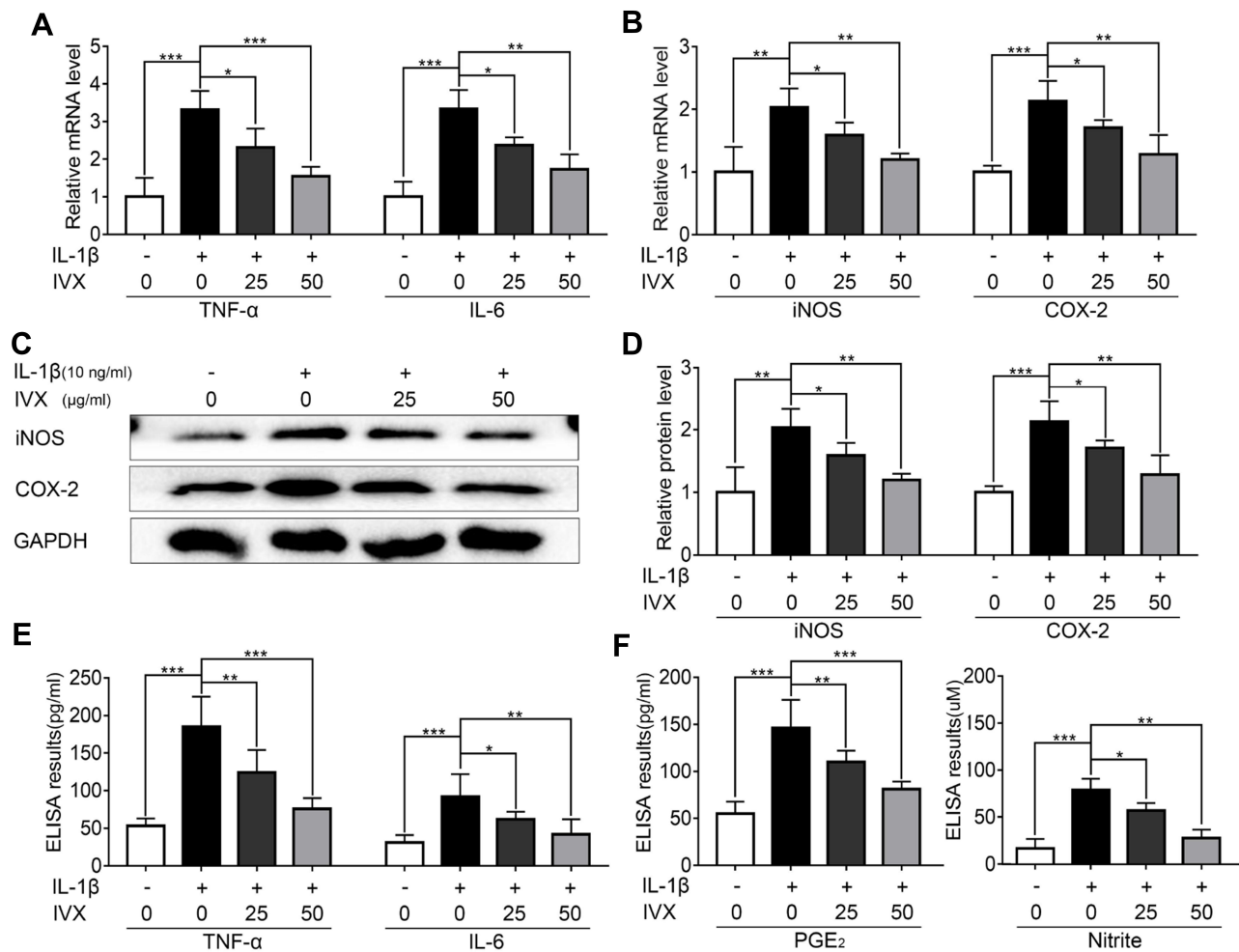


Figure 4 IVX modulation of pro-inflammatory factors in IL-1 β -exposed chondrocytes. (A, B) PCR analysis of inflammation-related transcripts. (C) Evaluation of inflammation-related protein expression, using Western blot. (D) Quantification of the protein expression using Image J software. (E, F) Detection of inflammation-related protein expression, using ELISA. Data presented as average \pm SD of 3 separate examinations. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$.

was markedly elevated (Figure 1F). In all, these results suggest that IVX protects ECM via Nrf2 regulation. We, also, examined iNOS and COX-2 levels in the IVX-exposed chondrocytic cells, using Western blot analysis. Based on our results, IVX exposure decreased iNOS and COX-2 protein levels, whereas Nrf2 deficiency increased both protein expression (Figure 1E and F), suggesting IVX regulates inflammation via Nrf2.

Discussion

OA is a wide-spread, incurable, degenerating illness, affecting joints, and is associated with severe discomfort, limited mobility, and disability. Currently, there is no effective therapy for OA. This disease is usually characterized by severe inflammatory attack on the joints. One therapeutic agent, IVX, has shown promise in suppressing inflammation in

other tissues. Therefore, this study explored whether IVX can modulate OA progression. Based on our results, we demonstrated that IVX indeed slows down OA progression by Nrf2-mediated inhibition of the NF- κ B signaling. This new finding can provide further insight into the future development of therapeutics against OA.

Multiple studies have demonstrated protective effects of IVX on inflammatory-related diseases like Alzheimer's disease,³⁶ neuroinflammation,³⁷ osteosarcoma,³⁸ and liver cancer.³⁹ To test IVX-induced protection of OA, isolated chondrocytes were exposed to a range of working and non-toxic IVX concentrations. We confirmed that IVX was able to suppress both excessive catabolism and inadequate anabolism of ECM in IL-1 β -exposed chondrocytes. This was consistent with another study where IVX was shown to decrease matrix metalloproteinases MMP-2 expression.⁴⁰

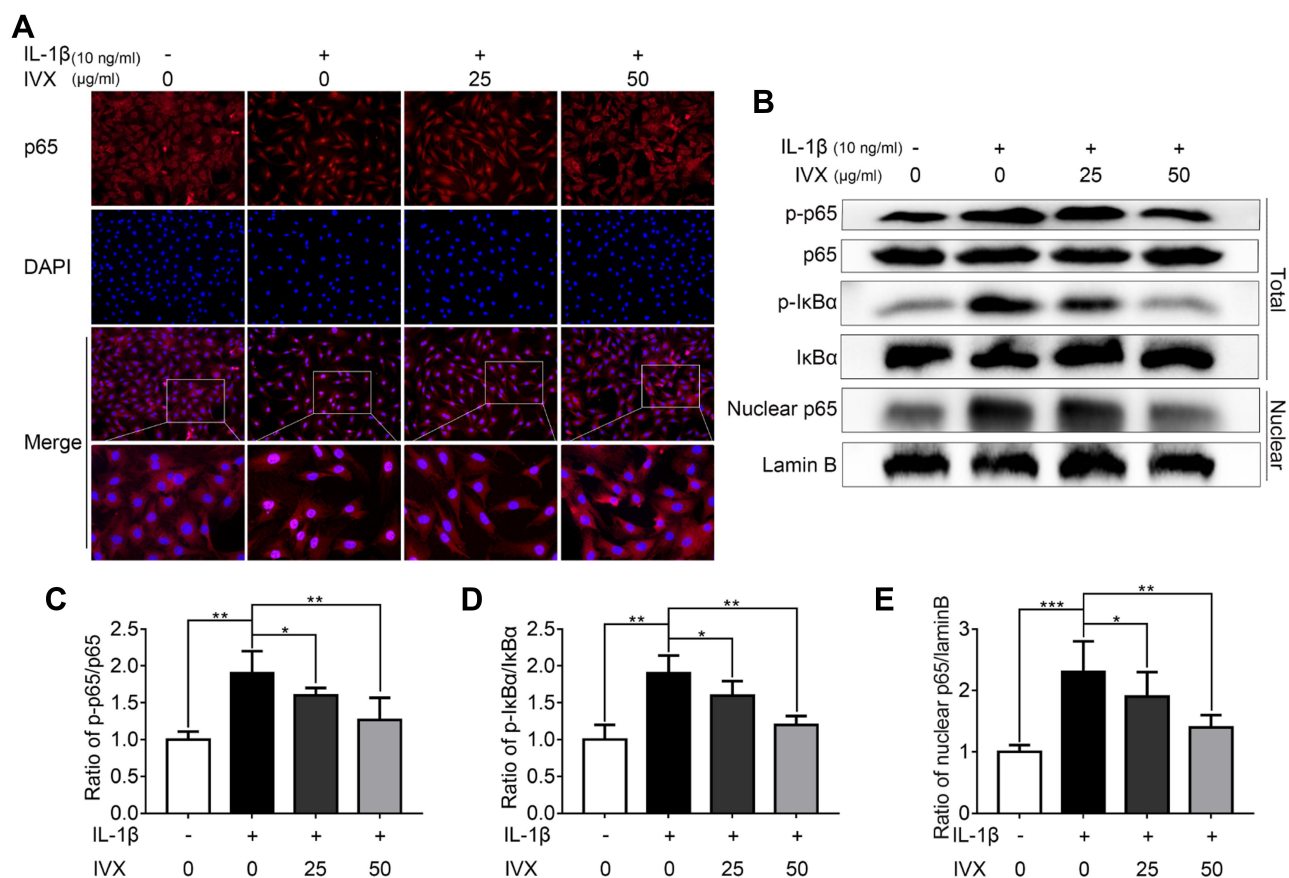


Figure 5 IVX impact on NF- κ B signaling in IL-1 β -exposed chondrocytic cells, with or without IVX. **(A)** P65 localization, detected by fluorescence microscopy, in conjunction with nuclear DAPI stain (scale bar: 50 μ m). **(B)** Evaluation of protein expression, using Western blot. **(C–E)** Image J employed quantification of protein expression. Data is average \pm SD of 3 separate examinations. * p < 0.05, ** p < 0.01, *** p < 0.001.

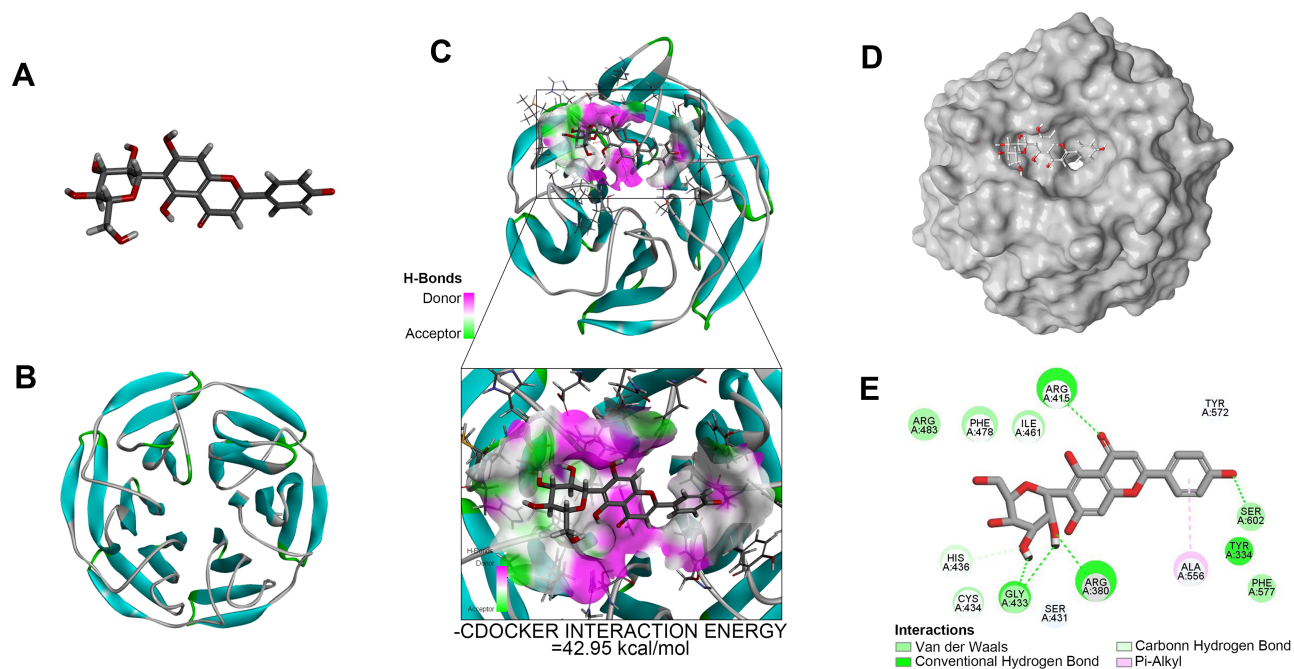


Figure 6 Molecular docking of IVX on its binding site in Nrf2, using docking studies analysis. **(A)** The IVX modular structure. **(B)** The Nrf2 ribbon modular structure. **(C)** The high interaction energy (-42.95 kcal/mol) interaction between IVX and Nrf2. **(D, E)** The 2D binding model between IVX and Nrf2, showing interaction through van der Waals, Hydrogen Bonds, and Pi-Alkyl.

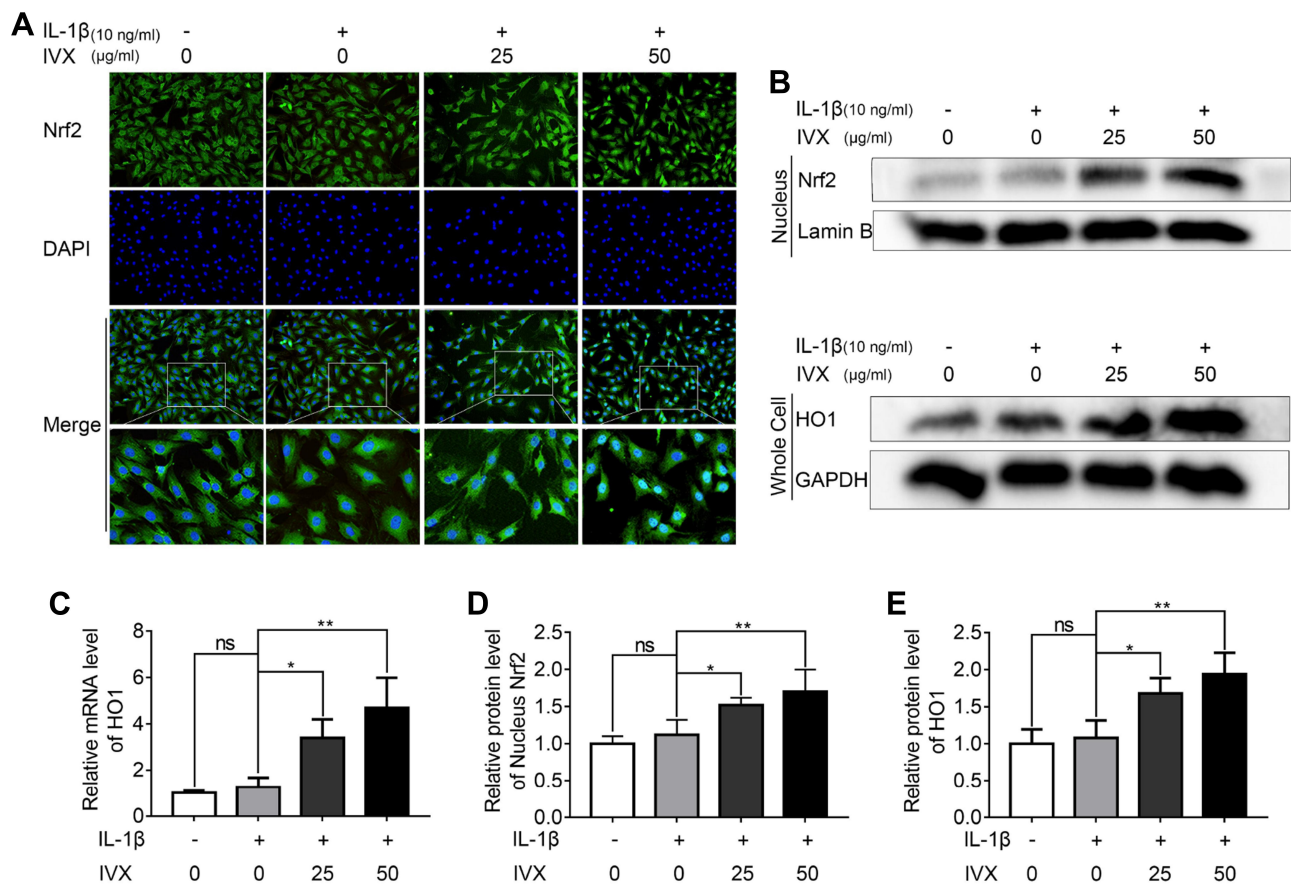


Figure 7 IVX impact on the Nrf2 signaling pathway in IL-1 β -exposed chondrocytes, in the presence of or absence of IVX. **(A)** Detection of Nrf2, using immunofluorescence in conjunction with nuclear DAPI stain (scale bar: 50 μ m). **(B, D and E)** Evaluation of Nrf2 levels in the nuclear fraction and downstream HO1 levels in the whole extract, using Western blot. Quantification of the protein was done using Image J. **(C)** Evaluation of HO1 transcript levels, using PCR. Data presented as average \pm SD of 3 separate examinations. * $p < 0.05$, ** $p < 0.01$.

Abbreviation: Ns, not significant.

Furthermore, inflammation is known to accelerate OA development,⁹ and existing evidences show that artificially suppressed inflammation can abrogate the progression of OA.^{27,41,42} In fact, Lv et al reported that IVX administration can notably reduce iNOS and COX-2 levels in monocyte/macrophage-like cells (RAW 264.7).³¹ Similarly, in another study, TNF- α and IL-6 were strongly suppressed upon IVX treatment.²⁹ Our findings corroborated previous studies in that IVX presence markedly lowered the amount of pro-inflammatory factors in IL-1 β -exposed chondrocytes.

The NF- κ B pathway controls IL-1 β -mediated inflammation.^{16,43} In this study, IVX administration was shown to significantly suppress the NF- κ B signaling by reducing the occurrence of phosphorylated p65, thereby, sequestering it in the cytoplasm. In agreement with this,

earlier studies have also shown marked IVX-mediated suppression of NF- κ B and its downstream signaling.^{30,44}

Moreover, past studies have also demonstrated the Nrf2 signaling pathway to be a major modulator in IL-1-mediated inflammation in chondrocytic cells.²² Moreover, the NF- κ B signaling can be effectively targeted by the Nrf2 pathway thus delaying the advancement of OA.^{26,27} Here, we showed strong evidence that IVX exposure can activate the Nrf2 signaling pathway in order to repress OA development. We were also able to demonstrate that IVX directly interacts with the AA residues in the inhibitory interacting pocket of Nrf2. Similar to our findings, Lv et al³¹ and Liu et al²⁹ also reported that IVX can suppress Nrf2 signaling.

In conclusion, we investigated the role of IVX in modulating OA and explored its underlying mechanism.

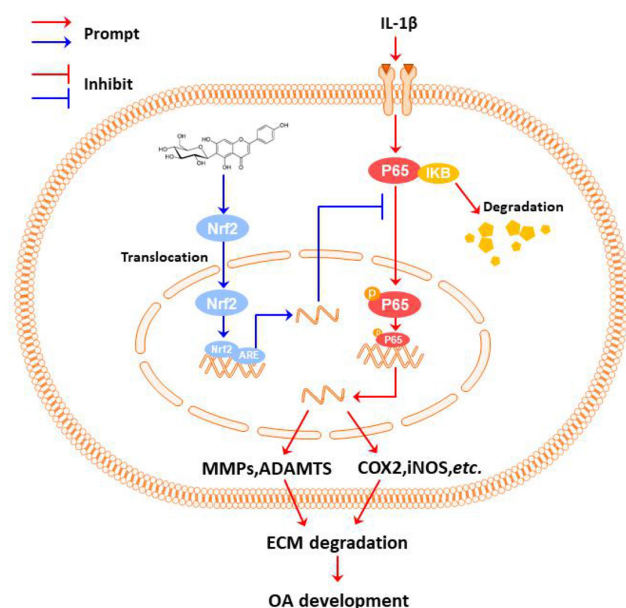


Figure 8 The proposed IVX-induced OA repression scheme. Under the stimulation of IL-1 β , the chondrocytic cell undergoes inflammation and ECM degeneration; IVX can suppresses chondrocytic cell inflammation and ECM destruction via the NF- κ B signalling pathway in its regulation of OA; Additionally, the IVX mechanism of action involved the Nrf2 activation.

We reported that IVX can indeed suppress IL-1 β -mediated inflammatory activation and maintain ECM balance in OA chondrocytic cells via the Nrf2/NF- κ B axis (Figure 8).

Data Sharing Statement

The data used to support the findings of this study are available from the corresponding author upon request.

Author Contributions

Quanqing Wan conceived of this research and helped modified the manuscript. Xiaofen Hu and Ruijie Li performed the animal model experiments and collected the samples. Haifeng Zhu and Fujiang Wang participated in the experimental design, guided the animal surgical operations, prepared reagents, materials and analytical tools. Xiaofen Hu wrote the manuscript. Ming Sun, Ying Kong participated in the statistical analysis, prepared all figures and embellished the manuscript. All authors made substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data; took part in drafting the article or revising it critically for important intellectual content; agreed to submit to the current journal; gave final approval of the version to be published; and agree to be accountable for all aspects of the work.

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Disclosure

The authors have no conflicts of interest.

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