

MAPK inhibitors protect against early-stage osteoarthritis by activating autophagy

CHUN-NA LAN¹, WEI-JUN CAI², JIE SHI³ and ZHONG-JIE YI⁴

¹Department of Rehabilitation, The Second Xiangya Hospital of Central South University, Changsha, Hunan 410011;

²Department of Histology and Embryology, School of Basic Medicine, Central South University;

³Department of General Surgery, The Third Xiangya Hospital of Central South University, Changsha,

Hunan 410013; ⁴Department of Burn and Plastic Surgery, The Second Xiangya Hospital of Central South University, Changsha, Hunan 410011, P.R. China

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Abstract. Osteoarthritis (OA) is a chronic, age-related osteoarthropathy that causes a considerable decline in quality of life, as well as economic losses due to its high incidence and poor prognosis. Mitogen-activated protein kinases (MAPKs) regulate multiple cellular processes, including proliferation, differentiation and apoptosis, in certain diseases, such as cancer, diabetes and Alzheimer's disease. The present study aimed to investigate the regulatory role of the MAPK signaling pathway in early-stage OA. A rabbit model of early-stage OA was induced by treatment with the enzyme papain. U0126 [an extracellular signal-regulated kinase (ERK) inhibitor], SP600125 [a Jun NH2-terminal kinase (JNK) inhibitor] and SB203580 (a p38 inhibitor) were administered to the rabbits via intra-articular injection. The severity of OA was assessed by histological examination using H&E, toluidine blue and safranin-O/fast green staining, as well by analyzing the glycosaminoglycan (GAG) content and determining the OA Research Society International (OARSI) score. Western blotting was used to detect the protein expression levels of

matrix metalloproteinase-3 (MMP3), ERK, phosphorylated (p)-ERK, p38, p-p38, JNK, p-JNK, Beclin1, UNC-51-like kinase 1 (ULK1) and microtubule-associated protein 1 light chain 3 (LC3)II/I. U0126, SP600125 or SB203580 treatment significantly decreased the OARSI scores and significantly increased the GAG levels in the cartilaginous tissues of OA model rabbits. These results indicated that the MAPK inhibitors reduced the severity of OA-induced injury at the early stage. Western blotting results demonstrated that MAPK inhibition significantly decreased the protein expression levels of MMP3 in OA cartilage. The protective effect of MAPK inhibitors in OA was mediated via the activation of autophagy, as demonstrated by the increased protein expression levels of LC3II/I, ULK1 and Beclin1. Overall, the data indicated that MAPK inhibitors may exert a protective effect against OA by restoring compromised autophagy. Furthermore, the present study suggested that MAPK inhibitors may represent a potential pharmacological strategy for treating OA in the future.

Introduction

Osteoarthritis (OA) is a chronic, age-related osteoarthropathy characterized by loss of cartilage, remodeling of subchondral bone, formation of articular marginal osteophytes and the presence of synovitis (1,2). OA in joints is a major cause of deformity, pain and disability, which results in considerable socioeconomic costs worldwide due to its high incidence (3,4). Degeneration of the cartilage structure is associated with changes in the extracellular matrix (ECM), including matrix metalloproteinase-3 (MMP3), collagen and glycosaminoglycan (GAG), which are all regarded as biological markers of OA (5). The severity of OA is correlated with MMP3 accumulation and GAG reduction. OA can be induced by various physical and biochemical triggers (6,7), with pathological changes being irreversible at the middle and late stages of the disease. Therefore, it is important to develop therapeutic strategies to treat the early stage of the disease (8).

Autophagy is a metabolic process in which cytoplasmic organelles and cytosolic components are degraded to protect cells from stress, and it is a mechanism that is present in numerous diseases (9-12). The expression levels of UNC-51-like

Correspondence to: Dr Zhong-Jie Yi, Department of Burn and Plastic Surgery, The Second Xiangya Hospital of Central South University, 139 Renmin Road, Changsha, Hunan 410011, P.R. China
E-mail: yizhongjie312@csu.edu.cn

Dr Jie Shi, Department of General Surgery, The Third Xiangya Hospital of Central South University, 138 Tongzipo Road, Changsha, Hunan 410013, P.R. China
E-mail: shijie3451@csu.edu.cn

Abbreviations: MAPK, mitogen-activated protein kinase; OA, osteoarthritis; MMP3, matrix metalloproteinase-3; GAG, glycosaminoglycan; ULK1, UNC-51-like kinase 1; LC3, microtubule-associated protein 1 light chain 3; ERK, extracellular signal-regulated kinase; JNK, Jun NH2-terminal kinase; OARSI, Osteoarthritis Research Society International

Key words: osteoarthritis, MAPKs, autophagy, JNK, ERK, MMP3

kinase 1 (ULK1), Beclin1 and microtubule-associated protein 1 light chain 3 (LC3), which function as the initiator, regulator and executor of autophagy, respectively, are commonly used as a biomarker of the autophagic process (13). Autophagy has been considered to serve as a protective mechanism in maintaining normal cartilage function, and compromised autophagy has been demonstrated to lead to OA and numerous other age-related diseases (14-18). Therefore, targeting autophagy may serve as a promising therapy for OA.

Mitogen-activated protein kinases (MAPKs) are a family of Ser/Thr protein kinases that are widely found in eukaryotes and include extracellular signal-regulated kinase (ERK), p38 and c-Jun NH₂-terminal kinase (JNK). MAPKs regulate multiple cellular processes, such as proliferation, differentiation, survival and apoptosis (19,20). Previous studies have demonstrated that JNK and p38 contribute to apoptosis, whereas ERK mediates proliferation in diseases such as OA, cancer, diabetes and Alzheimer's disease (19,21-23). However, the role of MAPKs in autophagy varies among diseases (24-31). Previous studies have also reported that p38 acts synergistically with autophagy, whereas other studies have demonstrated that MAPKs attenuate autophagy under certain conditions (32-34). The effect of MAPKs on autophagy in the pathogenesis of OA remains unclear.

In our previous study, compromised autophagy was demonstrated to be closely related to the progression of OA, and JNK and p38 MAPKs increased MMP3 expression and inhibited autophagy (18). Selective inhibition of the MAPK/MMP3 signaling pathway and restoration of impaired autophagy may serve as a novel therapeutic strategy for OA and other degenerative disorders.

In the present study, the effects of MAPK inhibitors on autophagy in a rabbit model of early-stage OA, using intra-articular injections of U0126 (an ERK inhibitor), SP600125 (a JNK inhibitor) and SB203580 (a p38 inhibitor), were explored.

Materials and methods

Ethics. All animal experiments were approved by the Institutional Animal Care and Use Committee of The Second Xiangya Hospital of Central South University (Changsha, China; approval no. 2018005). All experimental procedures were performed in accordance with the Animal Research: Reporting of *in vivo* Experiments guidelines (35) and the National Institutes of Health Guide for the Care and Use of Laboratory Animals (36).

Animals. Healthy male New Zealand rabbits (age, 4 months; weight, 2 kg; purchased from Hunan Pacific Biological Technology Co., Ltd.) were used in the present study. The animals (n=40) were randomly assigned into the following five groups: i) the control group; ii) the OA group; iii) the OA + U0126 group; iv) the OA + SP600125 group; and v) the OA + SB203580 group. In the control group, the animals received an articular injection of 0.9% NaCl (0.1 ml) once a day for 3 consecutive days. In the OA groups, an articular injection of 0.1 ml 4% papain enzyme was administered once a day for 3 consecutive days to establish the respective OA model. For the OA + U0126, OA + SP600125 and OA + SB203580 groups,

at 7 days after the last injection of papain, animals received an intra-articular injection of 0.1 ml (100 μ mol/l) of U0126 (an ERK inhibitor; Abcam; cat. no. ab120241), SP600125 (a JNK inhibitor; Abcam; cat. no. ab120065) or SB203580 (a p38 inhibitor; Abcam; cat. no. ab120162), respectively, once a day for 3 days. All animals were fed in cages at the Animal Center of Second Xiangya Hospital with a 12-h light/dark cycle, with free access to water and standard chow, and were monitored every day. The environmental temperature was maintained at 20-25°C and the relative humidity was 50 \pm 5%. Animals were euthanized at 15 days after the last intra-articular injection of papain by 20 ml/kg of air embolism under anesthesia (30 mg/kg of 3% pentobarbital sodium via the ear vein). Cartilage of the medial condyles of the femur was then removed for general observation, histological examination and western blotting. Changes in rabbit weight were calculated before and after the experiment.

General observation of OA severity. The distal epiphyses of the femurs were collected following animal euthanasia and washed in normal saline at room temperature. For the macroscopic observations, photographs of the rabbit knees were recorded using a digital camera from the same distance for each sample, and descriptive methods were used for examining the thickening of cartilage, cartilage rough surface and exudation of synovial fluid that were observed by the naked eye. OA cartilage was characterized by joint swelling, cartilage hyperemia, thickening of cartilage, a rough cartilage surface and the abundant exudation of synovial fluid.

Histology. The medial condyles of the femurs were fixed in 4% paraformaldehyde solution for 24 h and then decalcified in 11% EDTA disodium buffer. The buffer was replaced every 3 days until the cartilage was completely decalcified after 15 days. The condyles were then dehydrated and embedded in paraffin. Frontal serial sections (thickness, 6 μ m) were cut and mounted on polysine adhesion glass slides. Toluidine blue staining, H&E staining and safranin-o/fast green staining were performed to evaluate morphological changes in chondrocytes. Sections were observed using a light microscope. All the procedures carried out at room temperature.

To evaluate the pathological changes of early-stage OA, cartilage was subjected to histological examination of lesion depth, lesion extent over cartilage surface, matrix staining, chondrocyte clustering, surface fibrillation and abrasion quantification of GAG. For semi-quantitative analysis of GAG, Image-Pro Plus version 6 (Media Cybernetics, Inc.) software was used to measure the average optical density of the toluidine blue staining (at wavelength of 255 nm). Five randomly selected fields of view were analyzed from each slice and the average optical density of each group was compared. The severity of OA was evaluated based on the OA Research Society International (OARSI) scoring system as previously described (35,36).

Western blotting. Total protein was extracted from the medial condyles in lysis buffer (Cell Signaling Technology, Inc.) and used to detect the protein expression levels of phosphorylated (p)-ERK, ERK, p-p38, p38, p-JNK, JNK, MMP3, Beclin1, ULK1 and LC3B (including LC3-II and LC3-I). Total protein

concentration was quantified using the BCA Kit (Thermo Fisher Scientific, Inc.). Equal amounts of protein (20 μg) were separated via 10% SDS-PAGE and transferred onto PVDF membranes via electroblotting. The membranes were blocked for 1 h at room temperature in 5% BSA solution (cat. no. 9048-46-8; GENVIEW) in 1 mol/l TBS-Tween 20 [TBST; pH 7.5; 0.1% (v/v) Tween-20]. The membranes were subsequently incubated overnight at 4°C with the following primary antibodies: Anti- β -actin (1:5,000; cat. no. ab8227; Abcam), anti-ERK (1:500; cat. no. bs-0022R; BIOSS), anti-p38 (1:500; cat. no. bs-0637R; BIOSS), anti-JNK (1:500; cat. no. bs-2900R; BIOSS), anti-p-ERK (1:500; cat. no. bs-3016R; BIOSS), anti-p-p38 (1:500; bs-0636R; BIOSS), anti-p-JNK (1:1,000; cat. no. ab124956; Abcam), anti-ULK1 (1:1,000; cat. no. bs-3602R; BIOSS), anti-LC3B (1:500; cat. no. NB100-2220; Novus Biologicals, LLC), anti-Becn1 (1:1,000; cat. no. LS-C117076; LifeSpan Biosciences, Inc.) and anti-MMP3 (1:500; cat. no. 17873-1-AP; ProteinTech Group, Inc.). To visualize the bands following the primary incubation, the membranes were incubated with horse-radish peroxidase-conjugated secondary antibodies (1:6,000; cat. no. SA00001-2; ProteinTech Group, Inc.) for 1 h at room temperature. Membranes were washed three times, for 10 min each time, in 1 mol/l TBST after incubation with the primary and secondary antibodies. Protein bands were visualized using chemiluminescence (Thermo Fisher Scientific, Inc.). Grayscale densitometry analysis was performed using Image Pro-Plus version 6 software with β -actin as the loading control.

Statistical analysis. GraphPad Prism 7.0 (GraphPad Software, Inc.) was used to analyze the data. Analysis of three or more independent groups was performed by one-way ANOVA followed by Tukey's post hoc test. $P < 0.05$ was considered to indicate a statistically significant difference. Experiments were repeated ≥ 3 times and all animals were included in the analysis to avoid biased parameter estimates.

Results

General observation of OA severity. Table I displays the average body weight of the animals in each group before the study and at the time of sacrifice. Within each group, the change in body weight between the two time points was < 0.5 kg, which was not statistically significant. Compared with the healthy cartilage in the control group, the cartilage in the OA group exhibited joint swelling, cartilage hyperemia, thickening of cartilage, rough cartilage surfaces and abundant exudation of synovial fluid. The severity of OA was markedly reduced in animals receiving an intra-articular injection of U0126, SP600125 or SB203580 compared with that in the OA group (Fig. 1).

U0126, SP600125 and SB203530 ameliorate OA severity. Histological OA damage was evaluated by H&E, toluidine blue and safranin-o/fast green staining. Cartilage from the OA group exhibited pathological changes characterized by thickening of the cartilage surface, cell disorientation, cell clustering and uneven matrix distribution compared with the control group. However, these pathological changes were markedly attenuated in OA model animals treated with the MAPK inhibitors U0126, SP600125 and SB203530 (Fig. 2A-C).

Table I. Changes in rabbit weight before and after the experiment.

Group (n=8)	Before study, kg	After sacrifice, kg
Control	1.93 \pm 0.05	2.36 \pm 0.08
OA	1.99 \pm 0.05	2.38 \pm 0.07
OA + U0126	2.00 \pm 0.06	2.44 \pm 0.07
OA + SP600125	2.01 \pm 0.06	2.39 \pm 0.08
OA + SB203580	2.00 \pm 0.06	2.39 \pm 0.03

Data are presented as the mean \pm SD. Each of the two groups were compared using a paired Student's t-test. Results were not statistically significant ($P > 0.05$). OA, osteoarthritis.

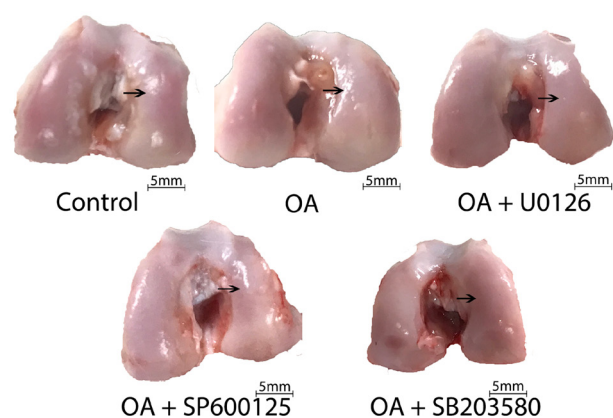


Figure 1. General observation of OA severity. Compared with the healthy cartilage in the control group, the cartilage in the OA group exhibited joint swelling, cartilage hyperemia, thickening of cartilage, rough cartilage surfaces and abundant exudation of synovial fluid. The severity of OA was markedly reduced in animals receiving an intra-articular injection of U0126, SP600125, or SB203580 compared with that in the OA group. Scale bar, 5 mm. OA, osteoarthritis. Arrows showed the thickening of cartilage and rough cartilage surfaces.

Semi-quantitative analysis of GAG was performed based on the optical density of the toluidine blue stain and was analyzed using Image-Pro Plus. The GAG staining in the OA group was significantly lower compared with that in the control group. However, the OA-induced decreases in GAG staining were significantly inhibited by U0126, SP600125 and SB203530 treatment (Fig. 2D). The OARSI scoring system was used to assess the severity and extent of OA based on H&E staining and safranin-O/fast green staining. The results demonstrated that U0126, SP600125 and SB203530 treatment significantly ameliorated the severity of OA compared with that in the OA group (Fig. 2E).

Inhibition of the ERK signaling pathway by U0126 restores autophagy in OA. Western blotting was used to analyze protein expression levels in cartilaginous tissues (Fig. 3). p-ERK protein expression levels were significantly increased in OA animals compared with those in the control group. Moreover, the OA-induced increase in p-ERK protein expression levels was significantly reduced by the

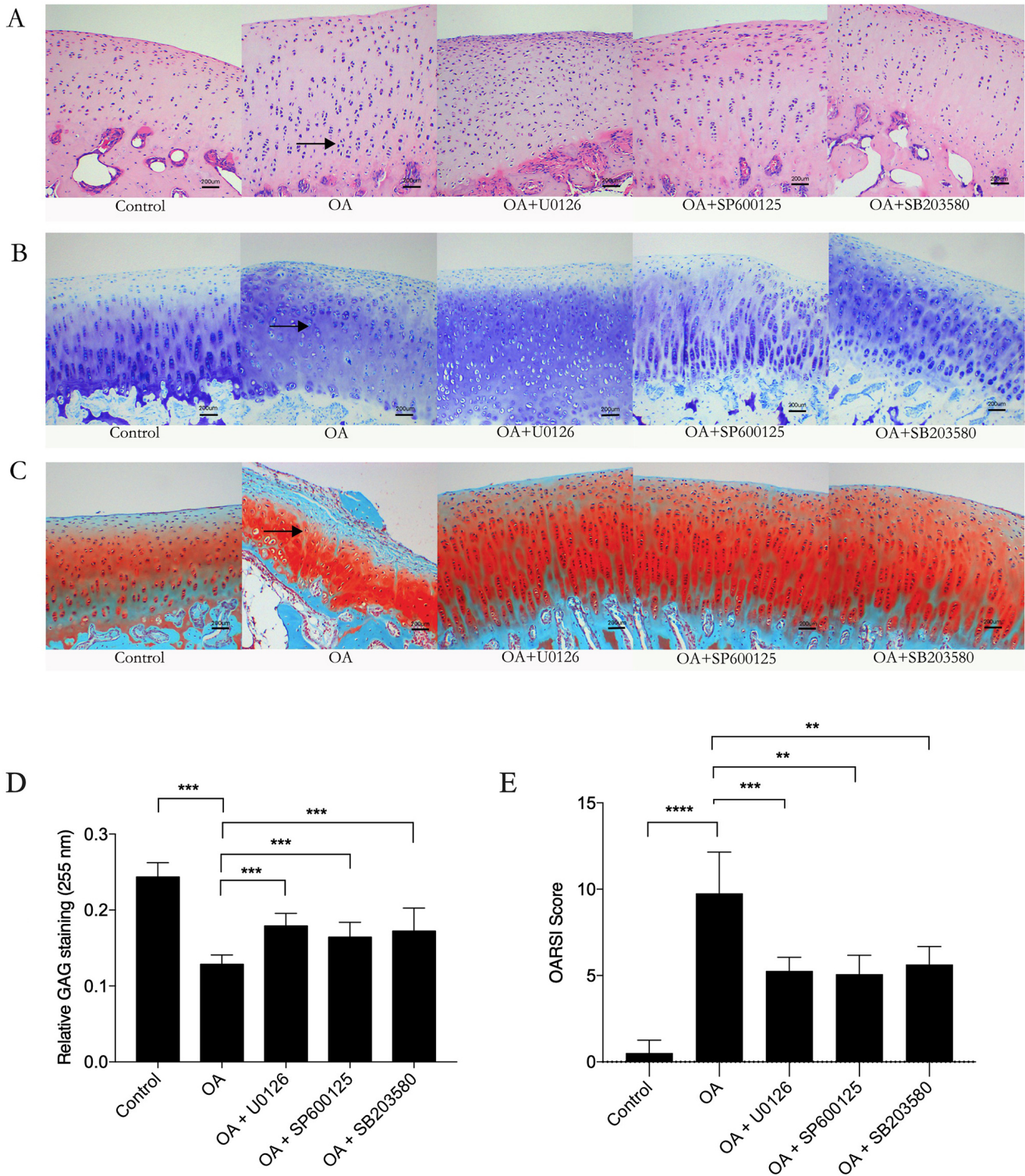


Figure 2. Histological observation of OA severity. (A) H&E, (B) toluidine blue and (C) safranin-o/fast green staining were used to determine OA severity (magnification, $\times 10$). OA changes included chondrocyte disorientation, loss of extracellular matrix and fibrillation and fissure formation. (D) Semi-quantitative analysis of GAG staining using the mean optical density of toluidine blue staining. (E) OA severity was evaluated using the OARSI scoring system. Arrows in (A) and (B) showed the cluster and disorientation of chondrocytes. Arrow in (C) showed the loss of extracellular matrix and fibrillation and fissure formation. Data are presented as the mean \pm SD. ** $P < 0.01$, *** $P < 0.001$ and **** $P < 0.0001$. OA, osteoarthritis; GAG, glycosaminoglycan; OARSI, Osteoarthritis Research Society International.

ERK inhibitor U0126. However, no statistically significant differences were observed in the total ERK protein expression levels among the control, OA and OA + U0126 groups. MMP3 protein expression levels were significantly increased in the OA group compared with those in the control group.

Administration of U0126 significantly suppressed MMP3 protein expression levels compared with those in the OA group, confirming that U0126 had a protective effect against OA pathogenesis. The cartilaginous tissue of OA model rabbits had significantly reduced LC3II/I protein expression

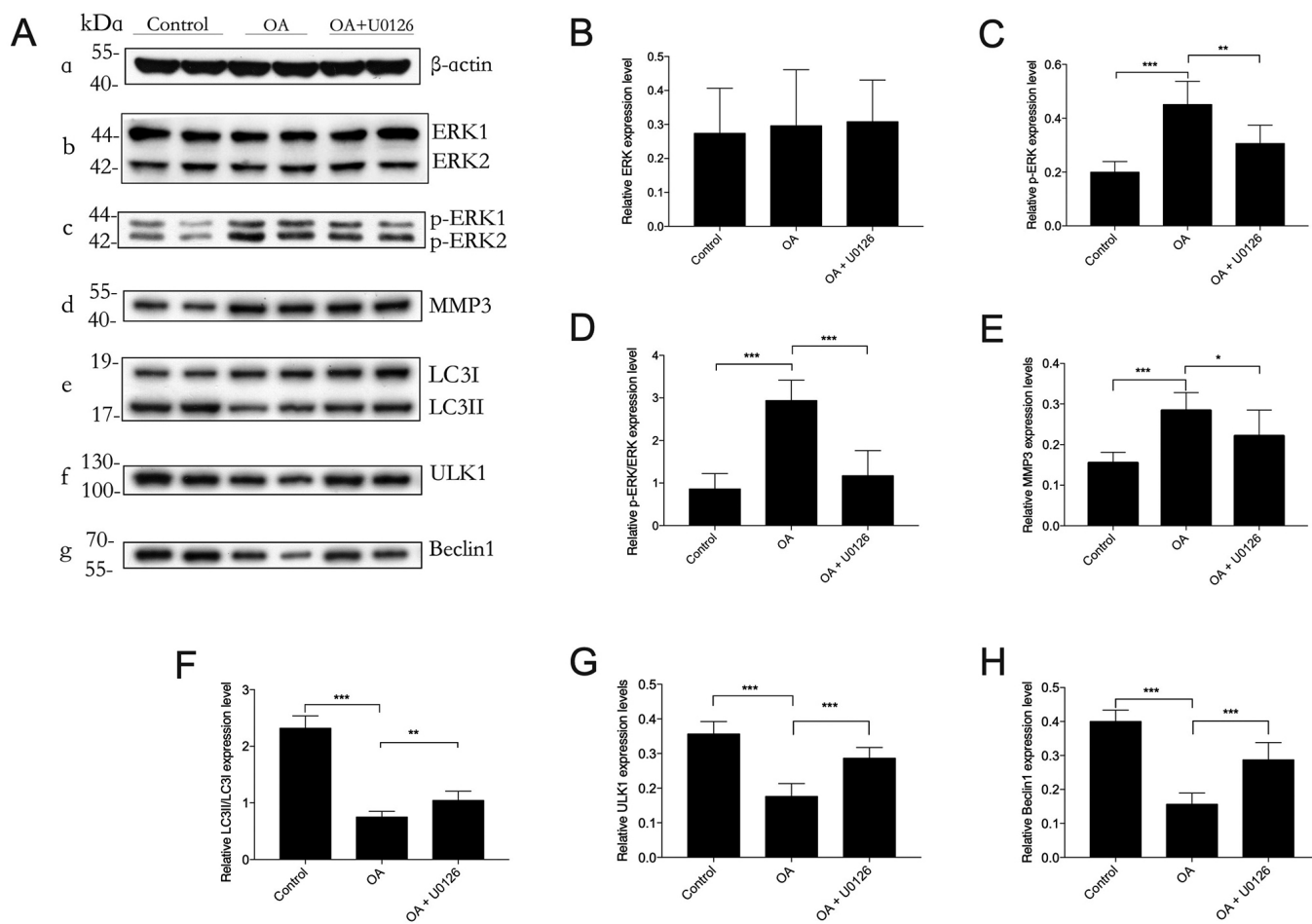


Figure 3. Effect of U0126 on autophagy in OA. Western blotting was performed to determine the protein expression levels (A-a) β -actin (loading control), (A-b) ERK, (A-c) p-ERK, (A-d) MMP3, (A-e) LC3II/LC3I, (A-f) ULK1 and (A-g) Beclin1 in cartilage lysates in each experimental group of rabbits. (B-H) Semi-quantification of band density. p-ERK protein expression levels were normalized to β -actin. n=3. *P<0.05, **P<0.01, ***P<0.001. OA, osteoarthritis; p, phosphorylated; ULK1, UNC-51-like kinase 1; MMP3, matrix metalloproteinase-3; LC3, microtubule-associated protein 1 light chain 3; ERK, extracellular signal-regulated kinase.

levels compared with those in the control group, which suggested that autophagy was compromised in OA. However, U0126 rescued the impaired autophagy process in OA, as demonstrated by the significantly increased LC3II/I protein expression levels in the OA + U0126 group compared with those in the OA group. U0126 treatment also significantly increased the expression of ULK1 and Beclin1 in the OA group compared with that in the control group. Overall, these results suggested that selectively blocking the ERK/MMP3 signaling pathway may protect against compromised autophagy induced by OA.

Inhibition of the JNK signaling pathway by SP600125 restores autophagy in OA. Western blotting demonstrated that the protein expression levels of MMP3 and p-JNK were significantly upregulated in the OA group compared with those in the control group. These upregulated protein expression levels were significantly reduced by SP600125 treatment (Fig. 4). The significant downregulation of the protein expression levels of autophagy markers LC3II/I, ULK1 and Beclin1 in the OA group, compared with those in the control group, were significantly attenuated by SP600125 treatment. These results suggested that the protective effect of the JNK inhibitor

SP600125 against OA may be mediated via restoration of impaired autophagy in OA.

Inhibition of the p38 signaling pathway by SB203580 restores autophagy in OA. Western blotting demonstrated that the protein expression levels of MMP3 and p-JNK were significantly increased in the OA group compared with those in the control group. This effect was significantly decreased by SB203580 treatment (Fig. 5). The protein expression levels of LC3II/I, ULK1 and Beclin1 were significantly downregulated in the OA group compared with those in the control group, which was significantly reduced by SB203580 treatment. These results demonstrated that the selective p38 inhibitor SB203580 potentially protected against OA by positively regulating autophagy in cartilaginous tissues with OA.

Discussion

OA is highly prevalent and has become a leading cause of disability (37). The healthcare resources and costs associated with managing OA can be substantial. Early intervention with the purpose of preventing disease onset and minimizing its progression is of vital importance (4,6). In the present

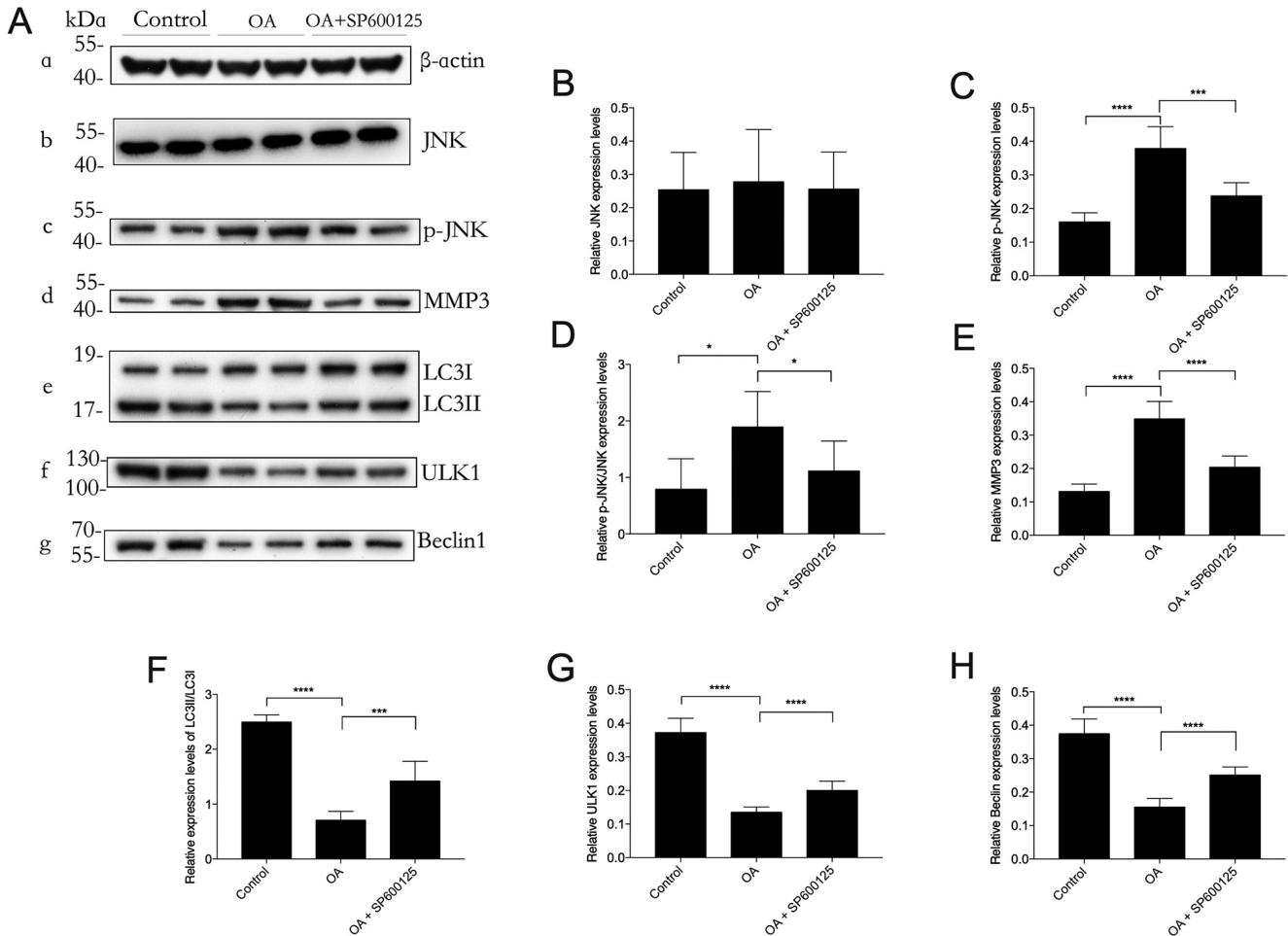


Figure 4. Effect of SP600125 on autophagy in OA. Western blotting was performed to determine the protein expression levels (A-a) β -actin (loading control), (A-b) JNK, (A-c) p-JNK, (A-d) MMP3, (A-e) LC3II/LC3I, (A-f) ULK1 and (A-g) Beclin1 in cartilage lysates in each experimental group of rabbits. (B-H) Semi-quantification of band density. p-JNK protein expression levels were normalized to β -actin. $n=3$. * $P<0.05$ *** $P<0.001$ and **** $P<0.0001$. OA, osteoarthritis; p, phosphorylated; ULK1, UNC-51-like kinase 1; JNK, Jun NH2-terminal kinase; MMP3, matrix metalloproteinase-3; LC3, microtubule-associated protein 1 light chain 3.

study, a rabbit model of early-stage OA was established via intra-articular injection of papain, and papain is a canonical reagent for inducing OA model (38,39). The current results indicated that early-stage OA could be characterized by joint swelling, cartilage hyperemia, cartilage thickening, rough cartilage surfaces and abundant exudation of synovial fluid. Although cartilage lesions could be easily identified in the early stages, we found there were no obvious symptoms or behavioral changes observed in the rabbits. The successful establishment of the OA model was further confirmed by subsequent histopathological assessment and the presence of chondrocyte proliferation, cell disorientation, loss of ECM, cartilage fibrillation and fissure formation. The OARSI scoring system was first proposed by Pritzker *et al* (40) in 2006 and is a useful methodology for assessing histopathological changes in OA cartilage. In contrast to the Mankin grading system (41), which primarily reflects the late stages of OA, the OARSI system, which is based on six grades reflecting the depth of the lesion and four stages reflecting the extent of OA over the joint surface, can identify differences in early or mild OA. An OARSI score <12 is considered to indicate early-stage OA. In 2016, a new OARSI grading

system was established that reflects the differences in cartilage stiffness between OARSI grade 0 (intact surface and no signs of degeneration) and grade 1 (intact surface and early signs of arthritis) (42). Based on this new scoring system and our previous findings (18), the OARSI score in the rabbit OA model on the 15th day after papain injection was determined to be ~ 12 . Therefore, in the present study, rabbits were euthanized and cartilage was obtained at this time point for all experiments. The results demonstrated that early-stage OA was successfully established in rabbits. Moreover, the administration of MAPK pathway inhibitors (U0126, SP600125, or SB203580) significantly ameliorated OA-induced injury, as demonstrated by significantly diminished OARSI scores and significantly increased GAG levels.

Autophagy is an evolutionarily conserved catabolic process required for cellular homeostasis that determines cell viability in response to stress. Autophagic systems in mammals are broadly categorized into three types: Macroautophagy, microautophagy and chaperone-mediated autophagy (43). Macroautophagy is the most common and best characterized form of autophagy in mammalian cells (9-12). The process of autophagy initiates with the

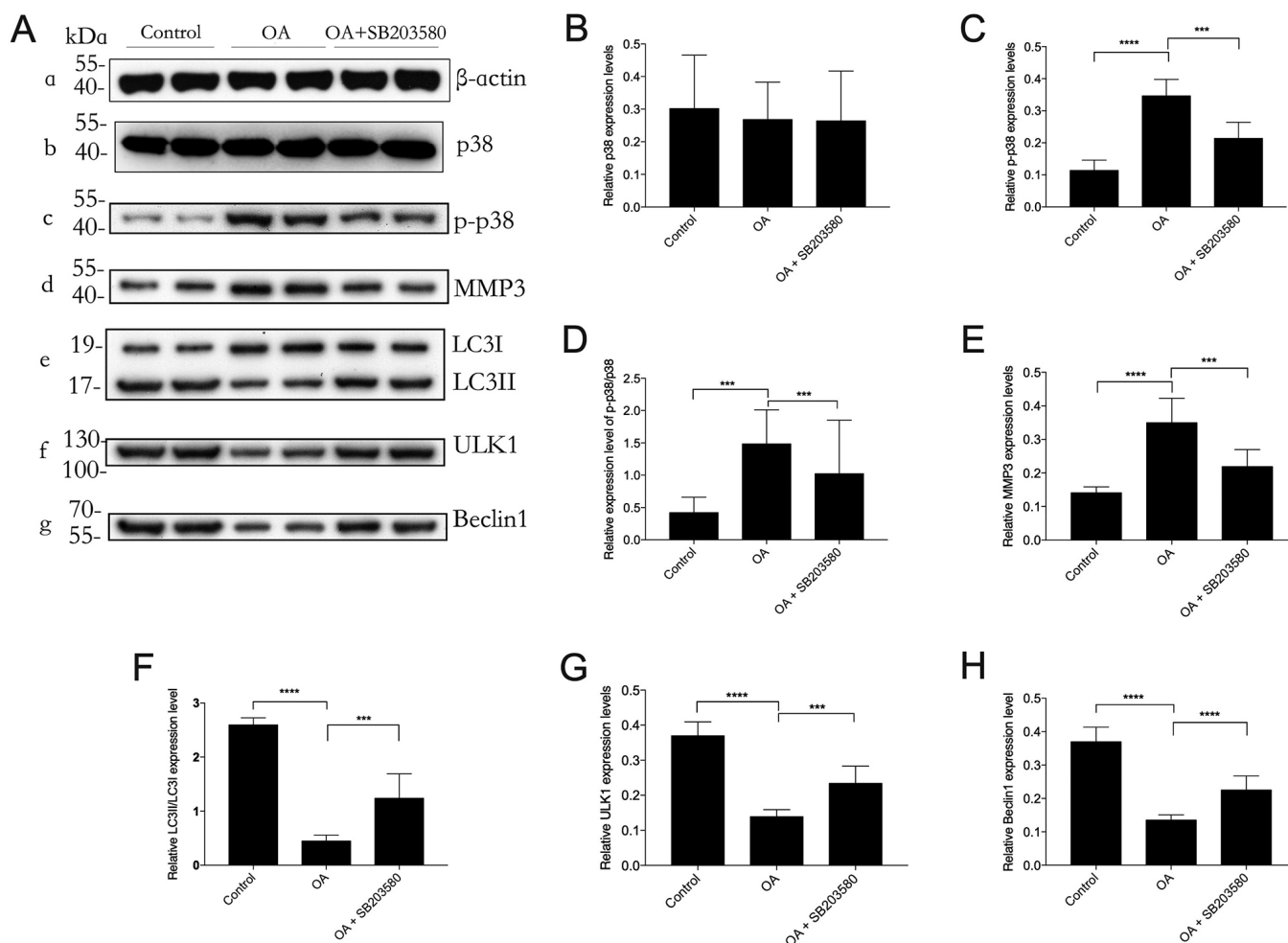


Figure 5. Effect of SB203580 on autophagy in OA. Western blotting was performed to determine the protein expression levels (A-a) β -actin (loading control), (A-b) p38, (A-c) p-p38, (A-d) MMP3, (A-e) LC3II/LC3I, (A-f) ULK1 and (A-g) Beclin1 in cartilage lysates in each experimental group of rabbits. (B-H) Semi-quantification of band density. p-p38 protein expression levels were normalized to β -actin. n=3. ***P<0.001 and ****P<0.0001. OA, osteoarthritis; p, phosphorylated; ULK1, UNC-51-like kinase 1; MMP3, matrix metalloproteinase-3; LC3, microtubule-associated protein 1 light chain 3.

formation of the phagophore. Under stressful conditions, mTOR is inactivated, which consequently facilitates the formation of the ULK complex, namely ULK1-FAK family-interacting protein of 200 kDa-autophagy-related protein (Atg) 13. The ULK complex subsequently recruits other autophagy-related proteins, and Beclin1 actively forms a complex with vacuolar protein sorting (VPS) 34 and the associated protein VPS15 to induce the formation of a phagophore. The elongation of the phagophore is facilitated by two ubiquitin conjugation systems: i) The Atg12-Atg5-Atg16 complex; and ii) the phosphatidylethanolamine (PE)-conjugated LC3II system. LC3I is formed by the cleavage of pro-LC3 at the C-terminus by Atg4 and is subsequently conjugated to PE with the help of Atg7 and Atg3 to form LC3II (9-12). LC3II binds to the autophagosome membrane and enables its fusion with the lysosome to form autophagolysosomes that subsequently degrade the accumulated cytosolic contents. In this way, autophagy serves an important role in cellular homeostasis and has been considered a protective mechanism in maintaining the function of cartilage (14-18). The present study demonstrated that the inhibition of ERK, JNK and p38 resulted

in a significant increase in the protein expression levels of ULK1, Beclin1 and LC3II/I in OA model rabbits, which suggested that the inhibition of MAPKs exerted a protective effect against OA by restoring impaired autophagy in OA.

Numerous signaling pathways and kinases regulate autophagy, including Atg9, BCL-2, the forkhead box O family of transcription factors, AMP-activated protein kinase, Akt, protein kinase C and the MAPKs (ERK, p38 and JNK) (9,30). The MAPK cascades are intracellular transduction pathways that respond to various extracellular stimuli and control multiple fundamental cellular processes, including growth, proliferation, differentiation, motility, stress responses and cell death (20). MAPKs not only serve an important role in regulating cell death by apoptosis but have also been implicated in autophagy (30). Hence, the MAPK signaling pathway has become an area of high interest in the field of autophagy research in recent years. However, the effect of MAPKs on autophagy varies among different diseases (24-31). Our previous study of early-stage OA demonstrated that with the development of OA, the key markers of the JNK and p38 MAPK pathways were elevated and the biomarkers of autophagy were decreased, suggesting

that JNK and p38 may negatively regulate the autophagic process but positively regulate MMP3 in OA (18). These findings are shown in the current study as well. ERK was also included in the present study in order to fully elucidate MAPK pathways in regulating autophagy in early-stage OA. It was found that p-ERK1/2 was also increased in OA, while the ERK inhibitor, similar to JNK and p38 inhibitors, enhanced the autophagic process in OA.

To the best of our knowledge, the present study was the first to investigate the effect of MAPK signaling on autophagy in early-stage OA via intra-articular injection of specific MAPK inhibitors in rabbits. The results indicated that selective inhibition of the ERK, JNK, or p38 MAPK signaling pathways significantly increased the protein expression levels of autophagy biomarkers (Beclin1, ULK1 and LC3II/I) in OA model rabbits. Overall, these results suggested that the inhibition of MAPKs may restore impaired autophagy in OA. Therefore, MAPK inhibitors exerted a protective effect against OA by restoring impaired autophagy, suggesting a potential therapeutic strategy for OA, as well as other degenerative disorders.

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Availability of data and materials

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

Authors' contributions

ZJY and CNL designed the study, CNL and JS conducted the experiments and JS and WJC analyzed the data. ZJY and WJC interpreted the results of the experiments. CNL generated the figures and drafted the manuscript and ZJY and JS edited and revised the manuscript. All authors reviewed and approved the final version of the manuscript. ZJY and JS confirm the authenticity of all the raw data.

Ethics approval and consent to participate

All animal experiments were approved by the Institutional Animal Care and Use Committee of The Second Xiangya Hospital of Central South University (Changsha, China; approval no. 2018005). All experimental procedures were performed in accordance with the Animal Research: Reporting of *in vivo* Experiments guidelines and the National Institutes of Health Guide for the Care and Use of Laboratory Animals.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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