

# miR-155-5p regulates macrophage M1 polarization and apoptosis in the synovial fluid of patients with knee osteoarthritis

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**Abstract.** Osteoarthritis (OA) is one of the most prevalent degenerative joint diseases that affects millions of individuals worldwide. During OA, proinflammatory factors (including IL-1, IL-6, IL-17 and TNF- $\alpha$ ) are released from chondrocytes and proliferating synoviocytes potentiate the proinflammatory microenvironment of the synovial fluid (SF). The altered SF microenvironment affects the infiltration, polarization and apoptosis of macrophages, though the underlying mechanisms are not completely understood. In the present study, the hypothesis that the knee synovial fluid of patients with knee osteoarthritis (KOA SF) promotes the polarization of peripheral blood mononuclear cell (PBMC)-derived M1 macrophages and inhibits PBMC-derived macrophage apoptosis was investigated. KOA SF increased PBMC-derived macrophage M1 polarization via the microRNA (miR)-155-5p/suppressor of cytokine signaling 1 signaling pathway. Caspase-3 (CASP3) was identified as a novel target of miR-155-5p, where KOA SF inhibited macrophage apoptosis via the miR-155-5p/CASP3 signaling pathway. The results suggested that the proinflammatory environment of KOA SF promoted macrophage M1 polarization and reduced macrophage apoptosis via miR-155-5p. The results provided a potential explanation for the increased number of M1 macrophages observed in KOA SF during OA. In addition, the present study suggested that miR-155-5p may serve as a potential therapeutic target for KOA.

## Introduction

Osteoarthritis (OA) is one of the most common degenerative joint diseases, affecting millions of individuals worldwide. OA is characterized by chondrocyte apoptosis and hypertrophy (1,2),

formation of osteophytes and inflammation of the synovial membrane (3). An increasing number of studies have reported that chronic inflammation with mild synovitis, elevated proinflammatory cytokine levels and infiltration of inflammatory cells (4) are linked to the progression of OA (5). During OA, cartilage integrity is lost and hypertrophic chondrocytes exhibit increased synthetic activity in an attempt to repair the damage. However, as a result of the increased synthetic activity, proinflammatory mediator products [including tumor necrosis factor (TNF)- $\alpha$  to interleukin (IL)-6 and IL-1 $\beta$ ] deregulate chondrocyte function and act on the adjacent synovium to stimulate proliferative and inflammatory responses (6). Proliferating synoviocytes also release proinflammatory products into the synovial fluid (SF), leading to a proinflammatory microenvironment (6). The altered SF microenvironment affects inflammatory cell infiltration, polarization and apoptosis. In addition, various proinflammatory factors such as IL-1, IL-1 $\beta$ , IL-6 and TNF- $\alpha$  secreted by the increased number of inflammatory cells, including M1 macrophages, promote the proinflammatory microenvironment of SF further to exacerbate OA. Previous studies have reported that mononuclear cells, including macrophages (7), B (8) and T cells (9), which aggregate during OA synovial tissues and fluid (4,6-8), contribute to chondrocyte degradation, local inflammation and activation of innate immune cells. A number of studies have demonstrated that the elevated number of macrophages present in the inflamed synovium and SF serve a major role during OA progression (10-12). Therefore, identifying the potential mechanism underlying the effects of the SF microenvironment on macrophage infiltration, apoptosis and polarization is important for identifying the mechanism underlying OA.

Macrophages serve important roles in innate immunity, displaying a high degree of plasticity and polarize into two main subtypes: Classically activated M1 macrophages and alternatively activated M2 macrophages (13,14). Previous studies have reported that M1 macrophages can be activated by interferon- $\gamma$  and toll-like receptors, which display a proinflammatory phenotype by producing proinflammatory cytokines, including IL-1, IL-6, TNF- $\alpha$  and IL-12 (15,16). Macrophages can also be polarized into an anti-inflammatory phenotype, known as M2 macrophages, which promote T helper cell 2 responses and contribute to tissue repair and healing (17,18). During OA, macrophages can be identified

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by measuring the expression of their respective cell markers using flow cytometry (FCM), including CD14, CD163 and CD68 (19). A previous study reported that local inflammation and activation of innate immune cells, primarily M1 macrophages, are major factors that contribute to the acceleration of OA progression (20). However, few studies have explored the impact of the proinflammatory microenvironment, including synovial tissues and SF, on macrophage infiltration, apoptosis and polarization during OA progression.

MicroRNAs (miRNAs/miRs) are small, single-stranded RNAs that are associated with a number of different diseases, including OA, diabetes and cancer (21-23). miRNAs control the differentiation and function of myeloid and lymphoid cells, among other cell types (24). A recent study on miRNAs in the immune system demonstrated that miR-146a and miR-155-5p are associated with OA (25). Additionally, it has been reported that increasing the expression of miR-155-5p leads to M1 polarization in the RAW264.7 macrophage cell line (26,27). It has also been reported that miR-155-5p promoted M1 macrophage polarization by targeting suppressor of cytokine signaling 1 (SOCS1), which is involved in STAT3 and AKT signaling (28-30). In the present study, the impact of knee SF from patients with knee OA (KOA) on macrophage polarization and apoptosis was investigated. miR-155-5p was identified as a potential target that affected macrophage apoptosis and polarization in KOA SF.

## Materials and methods

**Collection of clinical samples.** A total of 53 SF samples were collected by needle aspiration from the knee joints of patients with KOA (28 males and 25 females; age range, 50-79 years; average age, 66.7 years; admitted from January 2017 to December 2018; last follow up occurred 1 year after admission) receiving treatment at the Yantai Yuhuangding Hospital (Yantai, China). The present study was approved by the Ethics Committee of Yantai Yuhuangding Hospital and written informed consent was obtained from all participants. Blood samples (20 ml) from healthy subjects (20 males, 22 females; age range, 50-79 years; average age, 68.1 years; admitted from January 2017 to December 2018; last follow up occurred 1 year after admission) and patients with KOA were collected into sodium heparin Vacutainer® tubes (cat. no. 366480; Becton-Dickinson and Company). The severity of radiographic X-ray KOA was graded using the Kellgren/Lawrence scoring system (24), which is scored from 0 to 4. The X-ray presentations of the patients were as follows: i) Stage 2 KOA, 9 patients; ii) stage 3 KOA, 28 patients; and iii) stage 3 KOA, 15 patients. KOA SF was carefully cross-examined and processed within 2 h of fluid collection. KOA SF supernatant was collected by centrifugation at 16,000 x g at 4°C and stored at -80°C until further analysis.

**Cell culture and stimulation.** 293T cells (American Type Culture Collection) were cultured in DMEM (Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% heat-inactivated fetal bovine serum (FBS; Gibco; Thermo Fisher Scientific, Inc.) and penicillin and streptomycin (Gibco; Thermo Fisher Scientific, Inc.) at 37°C in a humidified incubator containing 5% CO<sub>2</sub>. Peripheral blood mononuclear cells (PBMCs) were obtained from whole

blood samples by density gradient centrifugation (400 x g at room temperature for 30 min) using Ficoll® 400 (Sigma-Aldrich; Merck KGaA), according to the manufacturer's protocol. After washing twice with PBS, PBMCs were seeded (3x10<sup>6</sup> cells/well) into 12-well plates with RPMI-1640 (Gibco; Thermo Fisher Scientific, Inc.) containing 10% FBS and 10 ng/ml macrophage colony-stimulating factor (R&D Systems, Inc.). After incubation for 5 days at 37°C, PBMC-derived macrophages were obtained, seeded (1x10<sup>6</sup> cells/well) into 12-well plates and co-incubated with 500 µl RPMI-1640-0.4% BSA (Sigma-Aldrich; Merck KGaA) and 500 µl SF for 12, 24 or 48 h at 37°C. Control cells were treated with 1 ml control medium (CM; RPMI-1640-0.4% BSA). CD14<sup>+</sup> monocytes/macrophages (MON/Mc) were isolated from the KOA SF and PBMCs of patients with KOA or normal subjects using human CD14 MicroBeads (cat no. 130-050-201; MACS; Miltenyi Biotec, Inc.), according to the manufacturer's protocols. PBMC-derived macrophage apoptosis was induced by incubation with 1% DMSO (Sigma-Aldrich; Merck KGaA) for 12 h at 37°C.

**Cell transfection.** PBMC-derived macrophages were seeded into 12-well plates at a density of 1.5x10<sup>5</sup> cells per well and transfected with 100 nmol/l miR-155 mimic (5'-UUA AUGCUAAUCGUGAUAGGGGU-3'), 100 nmol/l miR-155 inhibitor (5'-ACCCCUAUCACGAUUAGCAUUA-3') or 100 nmol/l miR-155 mimic/inhibitor negative control (NC; 5'-UCACAA CCUCCUAGAAAGAGUAGA-3'; Shanghai GenePharma Co., Ltd.) using Lipofectamine® 2000 reagent (Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. Following transfection for 24 or 48 h at 37°C, PBMC-derived macrophages were co-incubated with 500 µl RPMI-0.4% BSA and 500 µl SF for a further 24 h at 37°C. Control cells were treated with 1 ml CM.

**Reverse transcription-quantitative PCR (RT-qPCR).** Total RNA was extracted from macrophages and 293T cells used TRIzol® reagent (Thermo Fisher Scientific, Inc.). Total RNA was reverse transcribed into cDNA using the PrimeScript™ RT reagent kit (Takara Bio, Inc.), according to the manufacturer's protocol. Subsequently, cDNA was subjected to qPCR using SYBR® Premix Ex Taq™ (Takara Bio, Inc.). The thermocycling conditions were as follows: 5 min at 95°C, followed by 40 cycles at 95°C for 15 sec and 60°C for 30 sec. The primers used for qPCR are presented in Table I. mRNA and miRNA expression levels were quantified using the 2<sup>-ΔΔC<sub>q</sub></sup> method (31) and normalized to the internal reference genes GAPDH and U6, respectively.

**FCM staining, analysis and apoptosis detection.** PBMC-derived macrophage single-cell suspensions (1x10<sup>6</sup> cells/100 µl) were incubated at 4°C for 30 min with LIVE/DEAD™ Fixable Aqua Dead Cell Stain kit (Thermo Fisher Scientific, Inc.), which are a class of viability dyes suitable for identifying dead cells in samples that are sent for fixation. FcR-blocking reagent (10 µl; 10 min at 4°C) and fluorescently labeled antibodies targeted against CD86 (1:100; cat. no. 12-0869-42; eBioscience; Fluorophores, R-phycoerythrin; Thermo Fisher Scientific, Inc.), inducible nitric oxide synthase (1:100; iNOS; cat. no. 53-5920-82; eBioscience; Fluorophores, Alexa Fluor 488; Thermo Fisher Scientific, Inc.) and CD206 (1:100; cat. no. 12-2069-41; eBioscience; Fluorophores;

Table I. Primers used for quantitative PCR.

Gene	Primer sequence (5'→3')
IL-1 $\beta$	F: ATGATGGCTTATTACAGTGGCAA R: GTCGGAGATTCGTAGCTGGA
IL-6	F: ACTCACCTCTTCAGAACGAATTG R: CCATCTTTGGAAGGTTTCAGGTTG
NOS2	F: TTCAGTATCACAACCTCA R: TGGACCTGCAAGTTAAAAT
IL-10	F: GACTTTAAGGGTTACCTGGGTTG R: TCACATGCGCCTTGATGTCTG
ARG1	F: GTGGAAACTTGCATGGAC R: AATCCTGGCACATCGGGAAT
SOCS1	F: CACGCACTTCCGCACATTC R: TAAGGGCGAAAAAGCAGTT
CASP3	F: CATGGAAGCGAATCAATGGACT R: CTGTACCAGACCGAGATGTCA
Ym1	F: TCACAAACAAAAGG R: GAATATGTAACACATTCAA
miR-155	F: ATTGCCAATTTCTCTACCAC R: AGTAACAGGCATCATACT
GAPDH	F: CAAGGTCATCCATGACAACCTTG R: GTCCACCACCTGTTGCTGTAG
U6	F: CTCGCTTCGGCAGCACA R: AACGCTTACGAATTTGCGT

IL, interleukin; NOS2, nitric oxide synthase 2; ARG1, arginase 1; SOCS1, suppressor of cytokine signaling 1; CASP3, caspase-3; F, forward; R, reverse.

PE-Cy7; Thermo Fisher Scientific, Inc.) were then added. The cell pellet was resuspended in 200  $\mu$ l PBS and reacted with Alexa Fluor<sup>®</sup> 488-labeled goat anti-mouse IgG antibody (1:200; Invitrogen; Thermo Fisher Scientific, Inc.) at room temperature for 30 min in darkness. Apoptotic cells were detected using the Annexin V/PI Cell Apoptosis kit (Sungene Biotech Co., Ltd.), according to the manufacturer's protocol. FCM analyses were performed using a flow cytometer (BD Accuri C6 cytometer; BD Biosciences) and FlowJo software (version 7.6.1; FlowJo LLC).

**Western blotting.** Total protein was isolated from macrophages and cells using protein extraction kit (Bio-Rad Laboratories, Inc.), after which protein concentration was determined by a BCA assay (Beyotime Institute of Biotechnology). Subsequently, samples were separated by 8-12% SDS-PAGE and transferred onto nitrocellulose membranes using a semidry transfer apparatus. PVDF membranes were washed with TBST (0.05% Tween-20) three times before being blocked with 5% skimmed milk at room temperature for 2 h. The membranes were then incubated at 4°C overnight with primary antibodies targeted against: Phosphorylated (p)-STAT1 (1:1,000; cat. no. ab30645; Abcam), STAT1 (1:1,000; cat. no. ab2415; Abcam), p-STAT6 (1:1,000; cat. no. ab54461; Abcam), STAT6 (1:1,000;

cat. no. ab44718; Abcam), SOCS1 (1:1,000; cat. no. 3950; Cell Signaling Technology, Inc.), caspase-3 (CASP3) (1:1,000; cat. no. 9662; Cell Signaling Technology, Inc.) and GAPDH (1:1,000; cat. no. 60004-1-Ig; ProteinTech, Inc.). The membranes were washed with TBST three times, incubated with horseradish peroxidase-conjugated goat anti-rabbit (1:1,000; cat. no. ab6721; Abcam) and goat anti-mouse (cat. no. ab97040; 1:5,000; Abcam) secondary antibodies for 2 h at room temperature. ECL chemiluminescence solution and bicinchoninic acid protein quantitative detection kits were purchased from Beyotime Institute of Biotechnology. Finally, ImageJ (version 1.38; National Institutes of Health) was used to analyze the gray value.

**Luciferase reporter assay.** The interaction sites between miR-155-5p and CASP3 were predicted using miRanda (<http://www.microna.org>), miRwalk (<http://mirwalk.umm.uniheidelberg.de>) and miRTarbase (<http://mirtarbase.mbc.nctu.edu.tw>) databases. The online resource microRNA.org (<http://www.microna.org>) predicted the targeted binding site between miR-155-5p and the 3'-untranslated region (3'-UTR) SOCS1. The wild-type (WT) or mutant (MUT) 3'-UTR of SOCS1 or CASP3 (constructed by Guangzhou RiboBio Co., Ltd.) were cloned into the pMIR vector (Guangzhou RiboBio Co., Ltd.). 293T cells were seeded (4x10<sup>5</sup> cells/well) into plates with 0.5 ml complete growth medium. At 70-90% confluency, 293T cells were co-transfected with the miR-155 mimic or miR-155 inhibitor, negative control (miR-NC) and 50 ng pMIR-SOCS1-3'-UTR-WT or pMIR-SOCS1-3'-UTR-MUT using Lipofectamine 3000<sup>®</sup> reagent (Thermo Fisher Scientific, Inc.). For each well, 1  $\mu$ g/ $\mu$ l DNA was diluted in 50  $\mu$ l Opti-MEM (Gibco; Thermo Fisher Scientific, Inc.), whilst in a separate tube, 2.5  $\mu$ l Lipofectamine<sup>®</sup> 3000 was diluted in 50  $\mu$ l Opti-MEM. Each tube was gently mixed and incubated for 20 min at room temperature. Subsequently, the two solutions were added to each well and gently mixed. Cells were incubated for 48 h before assessing transfection efficiency. The same operating procedures were used for the miR-155-5p/CASP3 luciferase reporter assay using pMIR-CASP3-3'-UTR-WT and pMIR-CASP3-3'-UTR-MUT vectors. At 48 h post-transfection, cells were harvested and luciferase activities were measured using the Dual-Luciferase<sup>®</sup> Reporter System (Promega Corporation), according to the manufacturer's protocol. Firefly luciferase activity was normalized to that of *Renilla* luciferase.

**Statistical analysis.** Statistical analyses were performed using GraphPad Prism software (version 7; GraphPad Software Inc.). Data are presented as the mean  $\pm$  standard deviation. P<0.05 was considered to indicate a statistically significant difference. Comparisons among groups were analyzed using one-way ANOVA followed by Tukey's post hoc test.

## Results

**KOA SF promotes PBMC-derived M1 macrophage polarization and inhibits macrophage apoptosis.** The effects of SF from patients with KOA on PBMC-derived macrophages were assessed using RT-qPCR and FCM. KOA SF significantly promoted the expression of M1-type markers by the macrophages (Fig. 1A), whilst partially reducing the expression of M2-type markers (Fig. 1B) compared with macrophages incubated in CM. The FCM results indicated that SF significantly

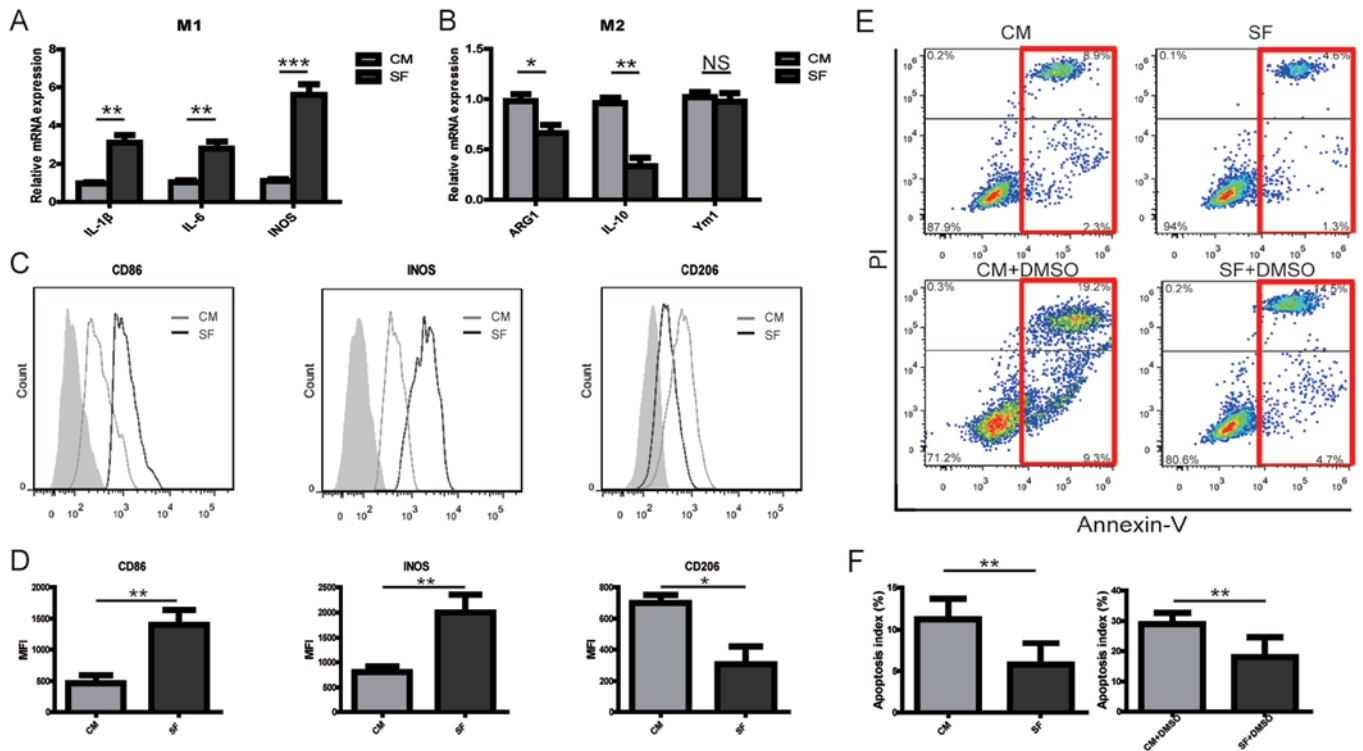


Figure 1. KOA SF promotes PBMC-derived macrophage M1 polarization and inhibits macrophage apoptosis. The relative expression of (A) M1 and (B) M2 markers in PBMC-derived macrophages cultured with either KOA SF or CM for 12 h. CD86, iNOS and CD206 expression levels in PBMC-derived macrophages cultured with KOA SF or CM for 24 h were (C) determined by flow cytometry and (D) quantified. The grey shadow represents the isotype control. Following treatment with or without 1% DMSO for 12 h in RPMI-1640 containing 10% FBS, PBMC-derived macrophages were cultured with KOA SF or CM for 24 h. Apoptotic cells were (E) detected by flow cytometry (red box indicates apoptotic cells) and (F) quantified. \* $P < 0.05$ , \*\* $P < 0.01$  and \*\*\* $P < 0.001$ . KOA, knee osteoarthritis; SF, synovial fluid; PBMC, peripheral blood mononuclear cells; CM, control medium; iNOS, inducible nitric oxide synthase; IL, interleukin; ARG1, arginase 1; Ym1, chitinase-like 3; NS, not significant; PI, propidium iodide.

promoted the expression of CD86 and iNOS, whilst reducing the expression of CD206 in PBMC-derived macrophages compared with those incubated with CM (Fig. 1C and D). During OA, cartilage loses its integrity, where the number of apoptotic chondrocytes and immune cells increases (6). The mechanism underlying this increase in the presence of KOA SF was therefore investigated by assessing its effects on macrophage apoptosis. In the presence and absence of 1% DMSO, KOA SF inhibited PBMC-derived macrophage apoptosis compared with those incubated with CM (Fig. 1E and F). The results suggested that the increase in M1 macrophages in KOA SF may be caused by the proinflammatory SF during the progression of OA.

*Expression of miR-155-5p is upregulated in KOA SF-derived macrophages and KOA SF-cultured PBMC-derived macrophages.* A recent study reported that miR-155-5p was associated with OA (21). The expression level of miR-155-5p was found to be upregulated in CD14<sup>+</sup> MON/Mc derived from the SF of patients with KOA compared with that in CD14<sup>+</sup> MON/Mc isolated from the PBMCs of patients with KOA or healthy individuals (Fig. 2A). In addition, the expression of SOCS1, a target of miR-155-5p, was revealed to be down-regulated in CD14<sup>+</sup> MON/Mc derived from the SF of patients with KOA compared with CD14<sup>+</sup> MON/Mc isolated from the PBMCs of patients with KOA or healthy individuals (Fig. 2B). PBMC-derived macrophages cultured in KOA SF expressed significantly higher levels of miR-155-5p compared with

macrophages cultured in CM (Fig. 2C). These results suggested that miR-155-5p may serve an important role in polarizing macrophages during OA progression.

*KOA SF-induced promotion of PBMC-derived M1 macrophage polarization is associated with the miR-155-5p/SOCS1 signaling pathway.* As a potential target of miR-155-5p, SOCS1-mediated inhibition of p-STAT1 activity and promotion of p-STAT6 activity to induce macrophage M2 polarization whilst inhibiting M1 macrophage polarization has been extensively studied in RAW264.7 cells (16). Therefore, it was hypothesized that miR-155-5p serves a role in KOA-SF-induced PBMC-derived M1 macrophage polarization via SOCS1. Transfection with the miR-155-5p mimic significantly increased miR-155-5p expression, whilst transfection with the miR-155-5p inhibitor significantly reduced miR-155-5p expression in PBMC-derived macrophages (Fig. 2D). Following transfection with NC or miR-155-5p mimic for 24 h, PBMC-derived macrophages were cultured with CM or KOA SF. The expression of miR-155-5p in PBMC-derived macrophages transfected with miR-155-5p mimic was upregulated compared with the CM NC group, while the upregulation of miR-155-5p expression in PBMC-derived macrophages cultured with KOA SF was not significantly different compared with PBMC-derived macrophages cultured with CM (Fig. 2E). Additionally, RT-qPCR results indicated that SOCS1 expression was significantly increased in the SOCS1-WT and SOCS1-MUT groups compared with that in

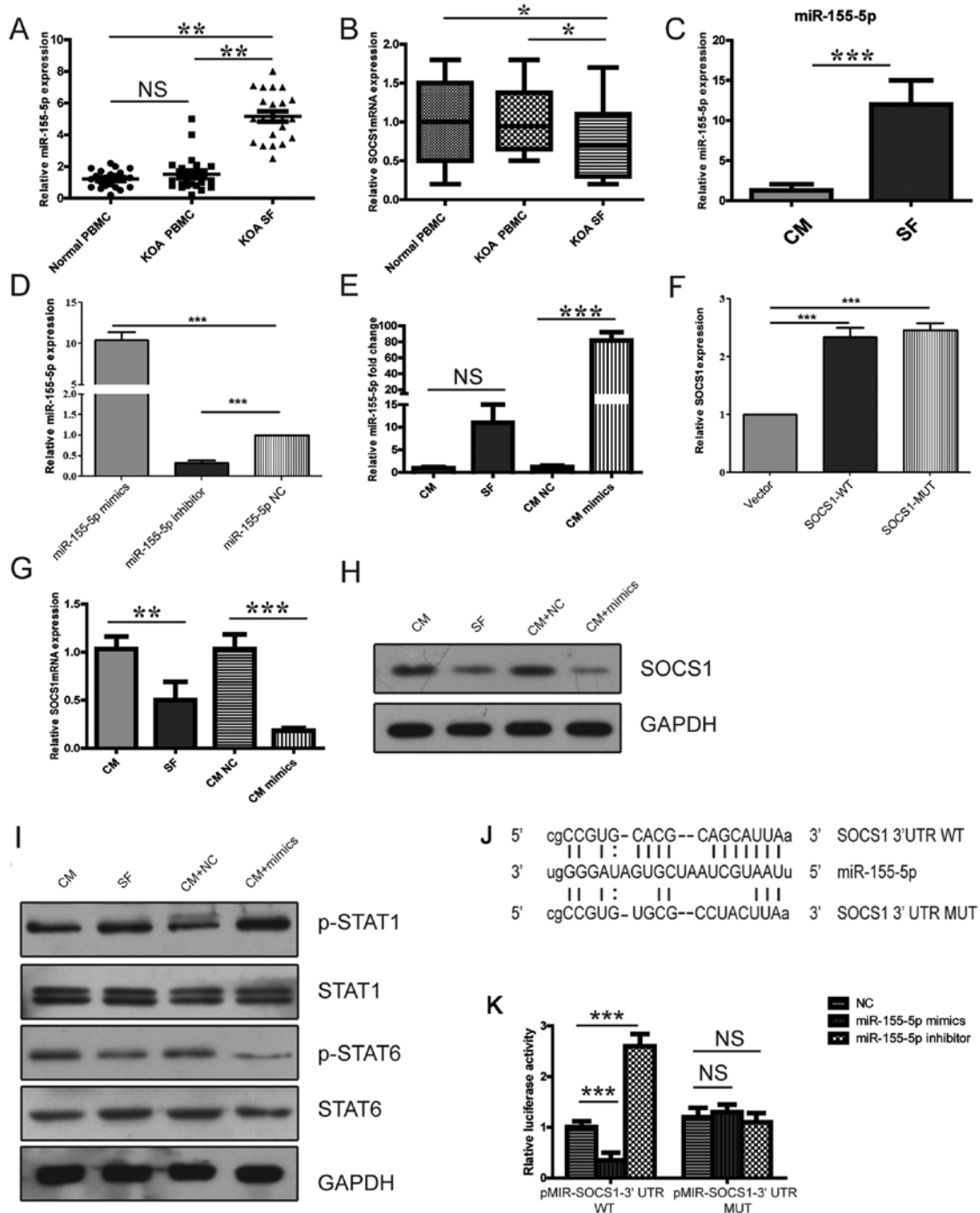


Figure 2. M1 polarization is associated with the miR-155-5p/SOCS1 signaling pathway. The relative expression of (A) miR-155-5p and (B) SOCS1 in PBMC-derived macrophages isolated from the blood samples of normal individuals, patients with KOA and PBMC-derived macrophages isolated from the KOA SF. (C) The expression of miR-155-5p in PBMC-derived macrophages cultured with KOA SF or CM for 12 h was detected by reverse transcription-quantitative PCR. The relative expression of miR-155-5p (D) in cells transfected with either NC, miR-155-5p mimic or the miR-155-5p inhibitor, (E) in transfected cells cultured with CM or KOA and in cells transfected with either NC or miR-155-5p cultured in CM. (F) The relative expression of SOCS1 in PBMC-derived macrophages transfected with control vector, SOCS-WT or SOCS-MUT. The relative expression of SOCS1 (G) mRNA and (H) protein in PBMC-derived macrophages treated and/or transfected with CM, KOA SF, CM + NC or CM + miR-155-5p mimic. (I) Levels of STAT1 and STAT6 phosphorylation in PBMC-derived macrophages treated and/or transfected with CM, KOA SF, CM + NC or CM + miR-155-5p mimic. (J) The potential interaction between miR-155-5p and SOCS1 3'-UTR-WT or SOCS1 3'-UTR-MUT. (K) The interaction between miR-155 and SOCS1 was confirmed using a dual-luciferase reporter assay. \*P<0.05, \*\*P<0.01 and \*\*\*P<0.001. miR, microRNA; SOCS1, suppressor of cytokine signaling 1; KOA, knee osteoarthritis; SF, synovial fluid; PBMC, peripheral blood mononuclear cells; CM, control medium; NC, negative control; p, phosphorylated; 3'-UTR, 3'-untranslated region; WT, wild-type; MUT, mutant; NS, not significant.

the control vector group (Fig. 2F). The expression of SOCS1 mRNA and protein in PBMC-derived macrophages cultured with KOA SF or transfected with the miR-155-5p mimic was found to be downregulated compared with that in the corresponding control groups (Fig. 2G and H). In addition, the

activation of STAT1 was revealed to be increased whereas the activation of STAT6 was found to be reduced in PBMC-derived macrophages cultured with KOA SF or transfected with miR-155-5p mimic compared with those in the corresponding control groups (Fig. 2I). The online resource microRNA.org

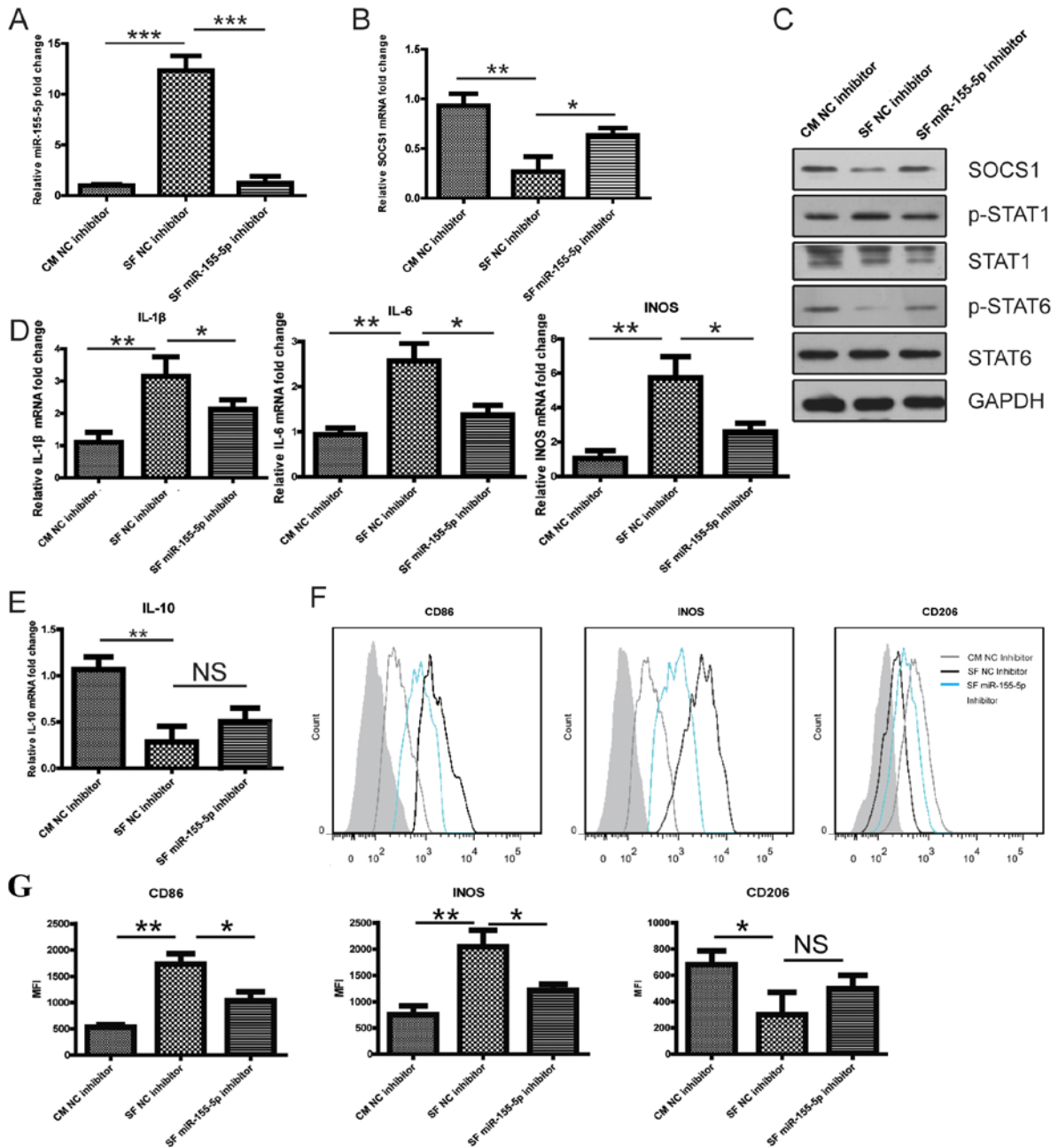


Figure 3. miR-155-5p downregulation promotes SOCS1 expression. (A-G) Following transfection with the NC inhibitor or miR-155-5p inhibitor for 24 h, PBMC-derived macrophages were cultured with CM or KOA SF. The relative expression of (A) miR-155-5p and (B) SOCS1 mRNA in PBMC-derived macrophages. (C) SOCS1 protein expression and the phosphorylation levels of STAT1 and STAT6 in PBMC-derived macrophages. The relative expression of (D) IL-1 $\beta$ , IL-6, iNOS and (E) IL-10 mRNA in PBMC-derived macrophages. The expression levels of CD86, iNOS and CD206 expression in PBMC-derived macrophages was (F) determined by flow cytometry and (G) quantified. The grey shadow represents the isotype control. \* $P < 0.05$ , \*\* $P < 0.01$  and \*\*\* $P < 0.001$ , as indicated. miR, microRNA; SOCS1, suppressor of cytokine signaling 1; NC, negative control; PBMC, peripheral blood mononuclear cells; CM, control medium; KOA, knee osteoarthritis; SF, synovial fluid; p, phosphorylated; IL, interleukin; iNOS, inducible nitric oxide synthase; NS, not significant.

(<http://www.microrna.org>) predicted the targeted binding site between miR-155-5p and the 3'-UTR SOCS1. Subsequently, dual-luciferase reporters encoding the WT or MUT 3'-UTR of SOCS1 were constructed (Fig. 2J). Co-transfection with the miR-155-3p mimic significantly inhibited the dual-luciferase activity of the SOCS1 WT 3'-UTR SOCS1 plasmid but not that of the SOCS1 MUT 3'-UTR plasmid. By contrast, the miR-155-5p inhibitor significantly enhanced the dual-luciferase activity of the SOCS1 WT 3'-UTR plasmid but not that of the SOCS1 MUT 3'-UTR (Fig. 2K). These results suggested a regulatory relationship between miR-155 and SOCS1 mRNA.

*Downregulation of miR-155-5p promotes SOCS1 expression and inhibits M1 polarization of KOA SF-stimulated PBMC-derived macrophages.* Following transfection with the NC inhibitor or miR-155-5p inhibitor for 24 h, PBMC-derived macrophages were cultured with CM or KOA SF. The downregulation of miR-155-5p reversed the reduced SOCS1 mRNA and protein expression by KOA SF (Fig. 3A-C). Furthermore, PBMC-derived macrophages cultured in KOA SF exhibited increased p-STAT1 and decreased p-STAT6 levels compared with those in the CM group. However, in PBMC-derived macrophages transfected with the miR-155-5p inhibitor and cultured

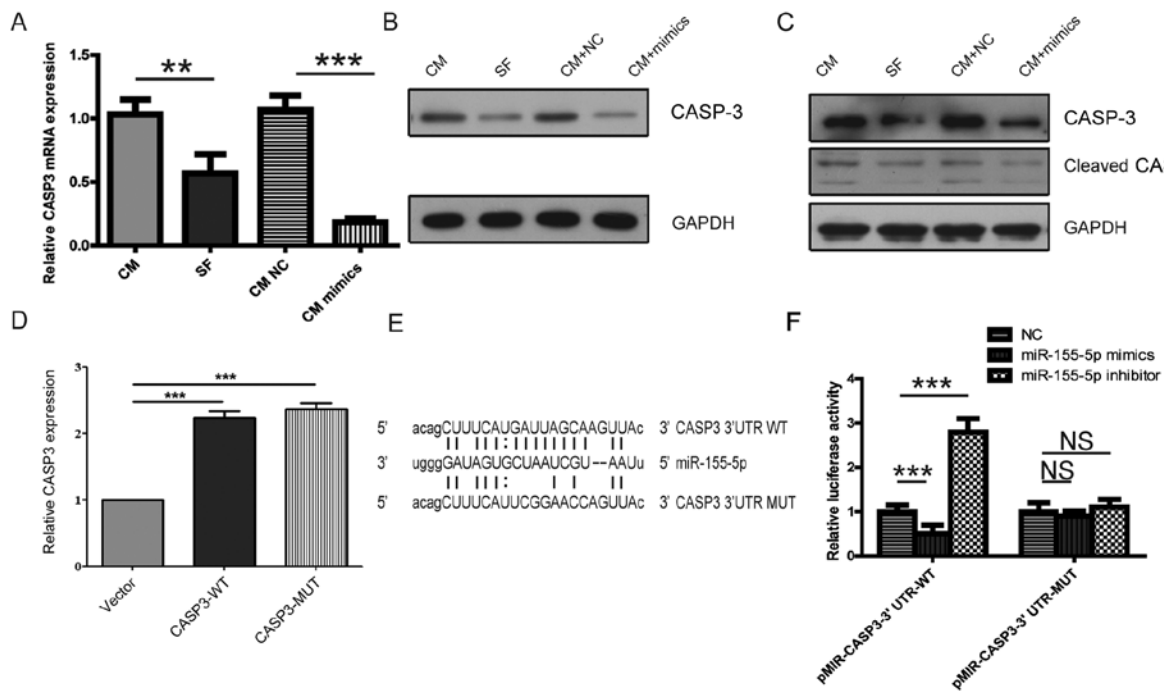


Figure 4. PBMC-derived macrophages are regulated by the miR-155-5p/CASP3 signaling pathway. (A-C) Following transfection with NC or miR-155-5p mimic for 24 h, PBMC-derived macrophages were cultured with CM or KOA SF. The relative expression of CASP3 (A) mRNA and (B) protein. (C) The relative expression of CASP3 and cleaved CASP3 in PBMC-derived macrophages treated with 1% DMSO. (D) The relative expression of CASP3 following transfection with the control vector, CASP-WT or CASP-MUT. (E) The binding sites of miR-155-p on CASP3 3'-UTR-WT or CASP3 3'-UTR-MUT. (F) The potential interaction between miR-155 and CASP3 was verified using a dual-luciferase reporter assay.  $^{**}P<0.01$  and  $^{***}P<0.001$ . PBMC, peripheral blood mononuclear cells; miR, microRNA; CASP3, caspase-3; NC, negative control; CM, control medium; KOA, knee osteoarthritis; SF, synovial fluid; 3'-UTR, 3'-untranslated region; WT, wild-type; MUT, mutant; NS, not significant.

in KOA SF, the KOA SF-induced effects on STAT1 and STAT6 phosphorylation were reversed (Fig. 3C). In PBMC-derived macrophages transfected with the miR-155-5p inhibitor, the KOA SF-induced increased expression of IL-1 $\beta$ , IL-6, iNOS and CD86 was reversed, however the reduced expression of IL-10 and CD206 as a result of KOA-SF treatment was not reversed (Fig. 3D, E, F and G). These observations indicated further that KOA SF promoted M1 macrophage polarization by increasing miR-155-5p, which targeted SOCS1 to upregulate p-STAT1 activity whilst inhibiting p-STAT6 activity.

**KOA SF-induced inhibition of PBMC-derived macrophage apoptosis occurs via the miR-155-5p/CASP3 signaling pathway.** miR-155-5p has been previously associated with apoptosis in several cancer cell lines and carcinomas (23); therefore, it was hypothesized that miR-155-5p may serve an important role in KOA SF-induced suppression of PBMC-derived macrophage apoptosis. The online microRNA, miRanda (<http://www.microrna.org>), miRwalk (<http://mirwalk.umm.uniheidelberg.de>) and miRTarbase (<http://mirtarbase.mbc.nctu.edu.tw>) databases predicted that CASP3 was a target gene of miR-155-5p, whilst the microRNA.org database identified the target binding site for miR-155 on the 3'-UTR of CASP3 mRNA (Fig. 4E). The expression of CASP3 mRNA was found to be significantly reduced in PBMC-derived macrophages cultured with KOA SF or transfected with miR-155-5p mimic compared with that in the corresponding control groups (Fig. 4A). The expression of CASP3 and cleaved CASP3 were also revealed to be decreased in PBMC-derived macrophages cultured with KOA SF or transfected with miR-155-5p

mimic compared with those in the corresponding control groups (Fig. 4B and C). Additionally, the RT-qPCR results indicated that CASP3 expression was significantly increased in CASP3-WT and CASP3-MUT groups compared with that in the control vector group (Fig. 4D). To further investigate the interaction between miR-155-5p and CASP3, WT and MUT CASP3 3'-UTRs were cloned into the pMIR-reporter plasmid (Fig. 4E). The dual-luciferase reporter assay indicated that the luciferase activity of pMIR-CASP3-3'-UTR-WT, but not pMIR-CASP3-3'-UTR-MUT, was significantly decreased by co-transfection with the miR-155-5p mimic and significantly enhanced by transfection with the miR-155-5p inhibitor compared that in the miR-NC group (Fig. 4F). These findings indicated that KOA SF increased miR-155-5p-induced inhibition of macrophage apoptosis by targeting CASP3.

**miR-155-5p downregulation promotes CASP3 expression and enhances macrophage apoptosis.** Following transfection with the NC inhibitor or miR-155-5p inhibitor for 24 h, PBMC-derived macrophages were cultured with CM or KOA SF. miR-155-5p downregulation was found to reverse the KOA SF-induced downregulation of CASP3 mRNA and protein expression (Fig. 5A and B). Furthermore, in 1% DMSO-treated PBMC-derived macrophages, KOA SF reduced the expression of CASP3 and cleaved CASP3 (Fig. 5C). However, in PBMC-derived macrophages transfected with the miR-155-5p inhibitor, the reduced expression of CASP3 and cleaved CASP3 by KOA SF was abolished (Fig. 5C). In addition, both in the presence and absence of 1% DMSO treatment, KOA SF treatment suppressed PBMC-derived macrophage apoptosis,

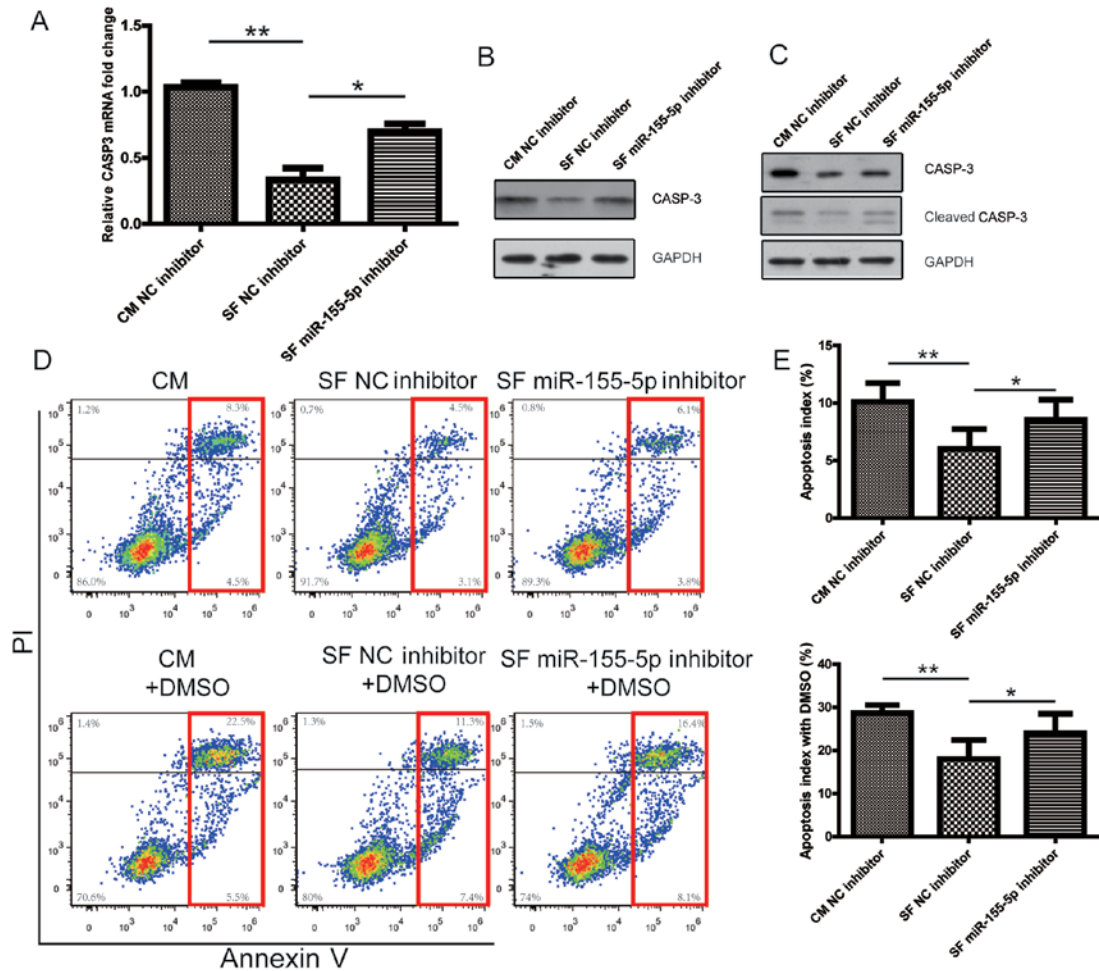


Figure 5. Downregulation of miR-155-5p promotes CASP3 expression and enhances macrophage apoptosis. (A-E) Following transfection with NC or miR-155-5p inhibitor for 24 h, PBMC-derived macrophages were cultured with CM or KOA SF. The relative expression of CASP3 (A) mRNA and (B) in PBMC-derived macrophages. (C) The relative expression of CASP3 and cleaved CASP3 in PBMC-derived macrophages treated with 1% DMSO. PBMC-derived macrophage apoptosis following treatment with or without 1% DMSO was (D) determined by flow cytometry (red box indicates apoptotic cells) and (E) quantified. \* $P < 0.05$  and \*\* $P < 0.01$ . miR, microRNA; CASP3, caspase-3; NC, negative control; CM, control medium; KOA, knee osteoarthritis; SF, synovial fluid; PBMC, peripheral blood mononuclear cells; PI, propidium iodide.

which was reversed by transfection with the miR-155-5p inhibitor (Fig. 5D and E). These results indicated that CASP3 is a novel target of miR-155-5p and that KOA SF inhibited macrophage apoptosis via the miR-155-5p/CASP3 signaling pathway.

## Discussion

OA is a debilitating condition that affects millions of individuals worldwide, where the only effective treatment strategies available are knee and hip replacement surgeries (2). Therefore, identifying the mechanism underlying OA is of substantial importance. OA has a complex pathogenesis where its etiology remains to be completely elucidated (1). A previous study has revealed that macrophages may serve as a major factor in the acceleration of OA progression (7). In particular, miR-155-5p has been previously found to associate with OA, where increased miRNA-155-5p expression leads to the M1 polarization of the RAW 264.7 macrophage cell line (27). It has also been reported that miR-155 is involved in TNF- $\alpha$ -mediated inhibition of osteogenesis differentiation, which directly targets a suppressor of cytokine signaling 1 (SOCS1) to inhibit bone morphogenetic protein 2-induced osteoblast differentiation (32). In the present

study, PBMC-derived macrophages cultured in KOA SF displayed upregulated expression of M1 markers, reduced apoptosis and increased miR-155-5p expression compared with those cultured in CM. The present study investigated the relationship between macrophage phenotype, apoptosis and KOA. The results indicated that the proinflammatory microenvironment of KOA SF may favor M1 macrophage polarization by upregulating miR-155-5p whilst inhibiting M1 macrophage apoptosis. Therefore, the results of the present study may provide an explanation for the increased ratio of total to M1-like macrophages in the granulocyte cell population during KOA progression. In addition, the results of the present study suggested that KOA SF altered PBMC-derived macrophages M1 polarization via the miR-155-5p/SOCS1 signaling pathway. CASP3 was subsequently identified as a novel target of miR-155-5p, where the results indicated that KOA SF inhibited macrophage apoptosis through the miR-155-5p/CASP3 signaling pathway. Based on the results of the aforementioned study (33), the present study suggested that KOA SF promoted PBMC-derived M1 macrophage polarization via the miR-155-5p/SOCS1 signaling pathway and inhibited macrophage apoptosis via the same signaling pathway.



It should be noted that the present study has a number of limitations. Since the infiltration of MON/Mcs in patients with OA was not investigated, the possibility that MON/Mc infiltration may affect the ratio of macrophages in the granulocyte cell population cannot not be ruled out. In addition, miR-155-5p was one of the most important elements that affected macrophage polarization phenotype and apoptosis in KOA SF vs. CM, but other factors can be involved. Further studies are required to explore the mechanism underlying the effect of the KOA SF microenvironment on macrophage polarization, phenotype and apoptosis. Based on the results of the present study, it was hypothesized that targeting the inflammatory KOA SF and macrophage polarization may delay or prevent OA progression. Further investigation into the potential of miR-155-5p as a therapeutic target for KOA is required.

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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

GDW designed the study, collected the data and wrote the manuscript. GSL and LC performed the experiments and interpreted the data. All authors read and approved the final manuscript.

### Ethics approval and consent to participate

The present study was approved by the Ethics Committee of the Yantai Yuhuangding Hospital (Yantai, China). Written informed consent was obtained from all patients.

### Patient consent for publication

Not applicable.

### Competing interests

The authors declare that they have no competing interests.

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