

FTY720 Inhibits the Development of Collagen-Induced Arthritis in Mice by Suppressing the Recruitment of CD4⁺ T Lymphocytes

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Background: Fingolimod (FTY720), a novel immunomodulator, was found to suppress the severity of collagen-induced arthritis (CIA) in mice. However, the potential molecular mechanisms are still unknown, and the effect of FTY720 on the recruitment of immune cells in the affected joints in the CIA model is not clear.

Materials and Methods: Following the oral administration of FTY720 (2 mg/kg) was treated into CIA mice per day for 35 days, intravital microscopy and immunofluorescence assays were performed to examine immune cell recruitment in the affected joints. Human MH7A synoviocytes were stimulated with tumour necrosis factor (TNF)- α and incubated with FTY720. Interleukin-1 β (IL-1 β), interleukin-6 (IL-6), and interleukin-8 (IL-8) mRNA and protein expression were evaluated using RT-PCR and enzyme-linked immunosorbent assay, respectively. Signal transduction pathway protein expression was measured by Western blotting. Nuclear translocation of nuclear factor (NF)- κ B was also analyzed by fluorescence microscopy.

Results: In vivo experiments showed that FTY720 inhibited the recruitment of CD4⁺ lymphocytes in the affected joints of CIA mice. FTY720 reduced the secretion of IL-1 β , IL-6, and IL-8 from TNF- α -stimulated MH7A cells in a dose-dependent manner. FTY720 also inhibited TNF- α -induced phosphorylation of NF- κ Bp65 and I κ B α , as well as NF- κ Bp65 nuclear translocation, in a dose- and time-dependent manner. Interestingly, FTY720 blocked PI3K/Akt, the upstream targets of the NF- κ B pathway.

Conclusion: Our findings demonstrated that oral administration of FTY720 exerted beneficial effects in CIA mice by inhibiting CD4⁺ T lymphocyte recruitment to the affected joints. Our data also indicated that FTY720 inhibited TNF- α -induced inflammation by suppressing the AKT/PI3K/NF- κ B pathway in MH7A cells.

Keywords: FTY720, rheumatoid arthritis, NF- κ B, MH7A, CD4⁺ T lymphocytes

Introduction

Rheumatoid arthritis (RA) is a chronic inflammatory autoimmune disease of unknown aetiology that is characterized by inflammation of the synovial tissues leading to destruction of periarticular bone and cartilage, which can lead to irreparable joint destruction if not properly treated.¹ At present, the main drugs for RA are disease-modifying antirheumatic drugs (DMARDs) and biologics.² However, approximately one-third of RA patients still have disease that is not adequately controlled. Therefore, new antirheumatic drugs with high efficacy are needed to meet RA therapy requirements.³

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Sphingosine-1-phosphate (S1P) is a metabolite of membrane sphingolipids, which can be phosphorylated by sphingosine kinase 1 (Sphk1) and Sphk2,⁴ is a metabolite of membrane sphingolipids. The transport and migration of various types of immune cells can be controlled by the S1P/S1P receptor axis, such as lymphocytes, neutrophils and macrophages. FTY720 is developed from the natural product ISP-1 (myriocin), the extract of *Cordyceps sinensis*. FTY720-P, the phosphorylated form of FTY720, is a structural analogue of S1P and exerts its effects by binding with the S1P receptor in vivo.⁵

The effects of FTY720 which is positive have been measured in a murine rheumatoid arthritis model. Researchers found that FTY720 could inhibit arthritis by sequestering CD4⁺ T lymphocytes in the thymus and inhibiting PGE₂ production in synovial cells.⁶ However, the effect of FTY720 on the recruitment of immune cells in the affected joint was not well understood, and the detailed molecular mechanisms of the therapeutic effects of FTY720 in RA have also not been fully elucidated.

Fibroblast-like synoviocytes (FLSs) play an important role in the development of RA, and the abnormal activation of FLSs eventually leads to synovial inflammation and joint damage.^{7,8} We performed intravital microscopy to evaluate the role of FTY720 in immune cell recruitment in the affected joints of CIA mice in this study. In addition, we examined the anti-inflammatory effect and mechanism of FTY720 in human MH7A synoviocytes as well.

Our results suggested that FTY720 inhibited the recruitment of CD4⁺ T lymphocytes in the affected joints of CIA mice. Our results also showed that the production of inflammatory cytokines was inhibited by FTY720 in human MH7A synoviocytes, which may be related to the inhibition of the PI3K/Akt/NF- κ B pathway.

Materials and Methods

Mice

Male DBA/1J mice were purchased from Shanghai Slac Laboratory Animal (Shanghai, China) at 6–8 weeks of age and maintained under specific pathogen-free conditions at Nanjing Medical University Animal Facility, China. Experimental protocols were approved by the Animal Research Ethics Committee of Nanjing Medical University. The guidelines followed for the welfare of the laboratory animals were the UK Animals Act, 1986.

Induction of Collagen-Induced Arthritis

The induction of collagen-induced arthritis (CIA) was showed by Brand et al⁹ which was described previously. First, collagen was dissolved in 0.1 M acetic acid (2 mg/mL) by gently stirring overnight at 4 °C and then emulsified with an equal volume of CFA containing mycobacteria to produce the inducing agent. On the day 0, 0.1mL of emulsion was injected subcutaneously 1–1.5 cm from the root of tail, and on day 21, near the initial injection site, a booster injection was executed.

Grouping of Animals

The animals were randomly divided into three groups (6 animals each) as follows: Group A: Control; Group B: Collagen-induced arthritis (CIA); and Group C: CIA + FTY720. The mice in group C received 2 mg/kg FTY720 each day by oral from the day of immunization until the mice were sacrificed.^{10,11} In the model group of CIA model and the control group, mice accepted saline instead.

Clinical Assessment of Arthritis

Beginning on day 25, arthritis severity was assessed every two or three days by two specially trained observers in a blinded manner using a 16-point scoring method as follows: no joint redness and swelling, 0 points; joint redness and mild swelling limited to the tarsal or ankle joint, 1 point; joint redness and swelling, mild swelling, swelling from the tarsal bone to ankle joint, 2 points; joint redness and swelling, moderate swelling extending from the tarsal bone to ankle joint, 3 points; and joint redness and swelling, severe swelling extending from toe to ankle joint, joint ankylosis and dyskinesia 4 points.¹² In case of obstruction, 4 points were scored. The total arthritis score was the sum of the limb joint score.

Intravital Confocal Microscopy

The surgical preparations for joint capillary imaging were based on previously described methods.¹³ By using a cocktail of ketamine (50 mg/kg), xylazine (10 mg/kg), and acepromazine (1.7 mg/kg) which were injected intraperitoneally, the mice were anaesthetized. Beginning 45 min later, anaesthesia was maintained by half-dose boosts administered subcutaneously every 30 min. The animals were intravenously (i.v.) administered PE/Dazzle™ 594 anti-mouse CD4 (Biolegend, San Diego, California, USA) (0.5 mg/kg, body weight) via the tail vein to label

CD4⁺ lymphocytes, and FITC anti-mouse CD8 (Biolegend) (0.5 mg/kg, body weight) was used to label CD8⁺ lymphocytes. After anaesthetizing the mice, depilatory cream was used for hair removal and skin preparation, and the mice were sterilized with 75% alcohol on absorbent cotton for 5 seconds. Sterile ophthalmic surgical instruments were used to open the mouse ankle joint tissue and expose the joint capillaries under a stereo dissection microscope. Images were acquired using an inverted epifluorescence Zeiss LSM 880 confocal laser system equipped with a 10×/0.3 Plan Neofluar objective or a Fluor 40×/1.3 objective. The microscope objective was thermostatically controlled to maintain a temperature of 37 °C. Videos were obtained by consecutive frames using appropriate combinations of 488-nm, 561-nm, and 633-nm laser lines. Cells that remained stationary for at least 30 seconds were considered to be adhered to the inner wall of the vessel.

Immunofluorescence Assay

Fresh samples of lower extremity joint synovium of the mice were embedded with OCT, and then frozen sections were prepared. After the frozen sections were naturally dried for 30 min, they were fixed with 4% paraformaldehyde prepared in 0.1 mol/L phosphate buffer (PBS, pH 7.2 ~ 7.4) for 30 min, washed with PBS 3 times, and permeabilized with 1% Triton X-100 for 20 min. The frozen sections were blocked with 3% bovine serum albumin (BSA) and 10% bovine serum albumin (BSA). PBS containing normal goat serum (250 µL) was added and incubated at 4 °C for 1 h in a wet box; the incubation solution was gently removed, and then the primary antibody against CD4⁺ T lymphocytes (1:150) (Abcam, Cambridge, MA, USA) was added. PBS was utilized as a negative control instead of primary antibody and was incubated overnight in a 4 °C wet box. The sections were washed three times with PBS, and FITC-conjugated secondary antibody (1:200) (Alexa Fluor[®] 594, Abcam) and DAPI were added to the wet box and incubated at 4 °C for 1 h. The sections were washed three times with PBS and sealed with an anti-fluorescence quencher. A fluorescence microscope (Nikon Eclipse TE2000-S, Nikon, Tokyo, Japan) was used to observe and photograph the samples in the darkroom. In each slide, three visual fields at ×200 magnification were selected, and the positively stained spots were counted by two pathologists in a blinded manner.

Cell Culture

MH7A cells were provided by Professor David Yu at the University of California, Los Angeles. This was approved by the institutional ethics committee of the Third Hospital of Hebei Medical University. Cells were maintained in high-glucose Dulbecco's modified Eagle's medium (DMEM, GE Healthcare HyClone, Logan, UT, USA) containing 10% FBS (Gibco, Gaithersburg, MD, USA) and 1% penicillin/streptomycin in a 37 °C incubator with 5% CO₂.

Cell Viability Assay

CellTiter-Blue[®] Cell Viability Assay (Promega Corporation, Madison, WI, USA) was applied to assess the cell viability. The 96-well culture plates at 3000–5000 cells/well received MH7A cells for 24 h and then treated with or without the indicated concentrations of FTY720 (Cayman, Ann Arbor, MI, USA) for 24 h. Then, CellTiter 96[®] Aqueous One Solution Cell Proliferation Assay reagent (20 µL/well) was added and incubated at 37 °C for 1–2 h. The absorbance was measured at 490 nm by a microplate reader.

Quantitative Real-Time Polymerase Chain Reaction (PCR)

MH7A cells were pretreated without or with various concentrations of FTY720 for 1 h and then incubated for 8 h in the use of 10 ng/mL TNF-α (PeproTech Rocky Hill, NJ, USA) or not. Total RNA was extracted from the cells in the use of TRIzol reagent (Invitrogen Life Technologies, Gaithersburg, MD, USA), and reverse transcription was applied in order to obtain single-stranded cDNA. In a 20-µL volume system which contained 4 µL of cDNA, 2 µL of each primer (forward and reverse), 10 µL of mix and 2 µL of ddH₂O, real-time PCR was performed in the use of the SYBR green PCR reagent (Takara). The primer sets which were used for amplification were as follows:

β-ACTIN: 5'-TGA CGT GGA CAT CCG CAA AG-3' (forward)

5'-CTG GAA GGT GGA CAG CGA GG-3' (reverse)

IL-1β: 5'- CCT GTC CTG CGT GTT GAA AGA -3' (forward)

5'- GGG AACTGG GCA GAC TCA AA -3' (reverse)

IL-6: 5'- CCT GAC CCA ACC ACA AAT GC -3' (forward)

5'- ATC TGA GGT GCC CAT GCT AC -3' (reverse)

IL-8: 5'-GGT GCA GTT TTG CCA AGG AG -3' (forward)

5'- TTC CTT GGG GTC CAGACA GA -3' (reverse).

Cytokine Quantification by ELISA

In the 6-well plates at 0.5×10^6 cells/well, MH7A cells were seeded for 24 h. The cells were pretreated with various concentrations of FTY720 for 1 h and then the cells were incubated with or without 10 ng/mL TNF- α for another 24 h. And then the culture medium was gathered, and the concentrations of IL-1 β , IL-6 and IL-8 were determined by ELISA using a commercial kit (BD Biosciences, San Diego, CA, USA) according to the manufacturer's instructions.

Immunofluorescence Staining

After pretreatment with or without FTY720 (10 nM) for 1 h, MH7A cells were incubated for 30 min when treated with TNF- α (10 ng/mL). The cells were incubated with anti-NF- κ B-p65 antibody (Cell Signaling Technology, Beverly, MA, USA) overnight and incubated with FITC-conjugated secondary antibody for 1 h at room temperature. After being washed in PBS, the cells were stained with 0.1 mg/mL DAPI. The stained cells were visualized using fluorescence microscope (Nikon Eclipse TE2000-S, Nikon, Tokyo, Japan).

Western Blotting

In order to extract proteins, the cells were lysed in 1 mL of RIPA buffer supplemented with 1% PMSF and 1% phosphatase inhibitor, and the protein concentration was tested by using BCA kit (Beyotime Institute of Biotechnology, Shanghai, China). Then, in the use of 10% SDS-PAGE, 20 μ g of the protein was separated and transferred to a PVDF membrane. Then, at room temperature, the membrane was blocked with 5% skim milk for 2 h and incubated at 4 °C overnight using the following primary antibodies: anti-PI3Kp85 (Cell Signaling Technology), anti-p-PI3K p85 (Cell Signaling Technology), anti-Akt (Cell Signaling Technology), anti-p-Akt (Cell Signaling Technology), anti-NF- κ Bp65 (Cell Signaling Technology), anti-p-NF- κ Bp65 (Cell Signaling Technology), anti-P-I κ B α (Abcam, MA, USA), and anti-GAPDH (Cell Signaling Technology). The membrane was incubated with HRP-labelled secondary antibodies at room temperature for 2 h, after three times washed with PBST. The membranes were washed again (three times) in use of PBST and visualized utilizing enhanced chemiluminescence.

Statistical Analysis

Statistical analysis was performed using GraphPad Prism 5.0 (GraphPad Software Inc., San Diego, CA, USA). A one-way analysis of variance (ANOVA) was used for the statistical analysis among different groups. Each value is presented as the mean \pm standard deviation (SD). The results with $P < 0.05$ were considered statistically significant.

Results

FTY720 Suppressed the Development of CIA

The FTY720 group were immunized with CII/CFA and received oral FTY720 (2 mg/kg/d), in order to observe the inhibitory effect of FTY720 on CIA, in DBA/1 mice. In the CIA group, signs of arthritis came into develop 4–6 days after the booster immunization. FTY720 treatment reduced the mean articular scores. On days 40 and 42 after the first immunization, there were significant differences in the mean articular scores between the groups (Figure 1).

T Lymphocytes into the Affected Joints of CIA Mice but Did Not Affect the Recruitment of CD8 T Lymphocytes

By using intravital microscopy, we found that CD4⁺ T lymphocyte adhesion in the peripheral vessels of the lower extremity joint was obviously increased in the CIA group compared with the normal control group. Notably, after FTY720 treatment, CD4⁺ T lymphocyte recruitment in the peripheral vessels of the lower extremity joint was significantly reduced (Supplementary Video S1, Figure 2A). There were no significant differences in the recruitment of CD8⁺ T lymphocytes in the peripheral blood vessels of the lower extremity joint in any of the groups (Supplementary Video S2, Figure 2B).

The Influence of FTY720 Treatment on the Number of CD4⁺ T Lymphocytes in the Synovial Tissue of CIA Mice

The immunofluorescence results showed that the number of CD4⁺ T lymphocytes in the CIA model group (69.7 ± 6.9) was higher than that in the normal control group. Compared with that of the CIA group, the number of CD4⁺ T lymphocytes in the FTY720 group was significantly decreased (Figure 3).

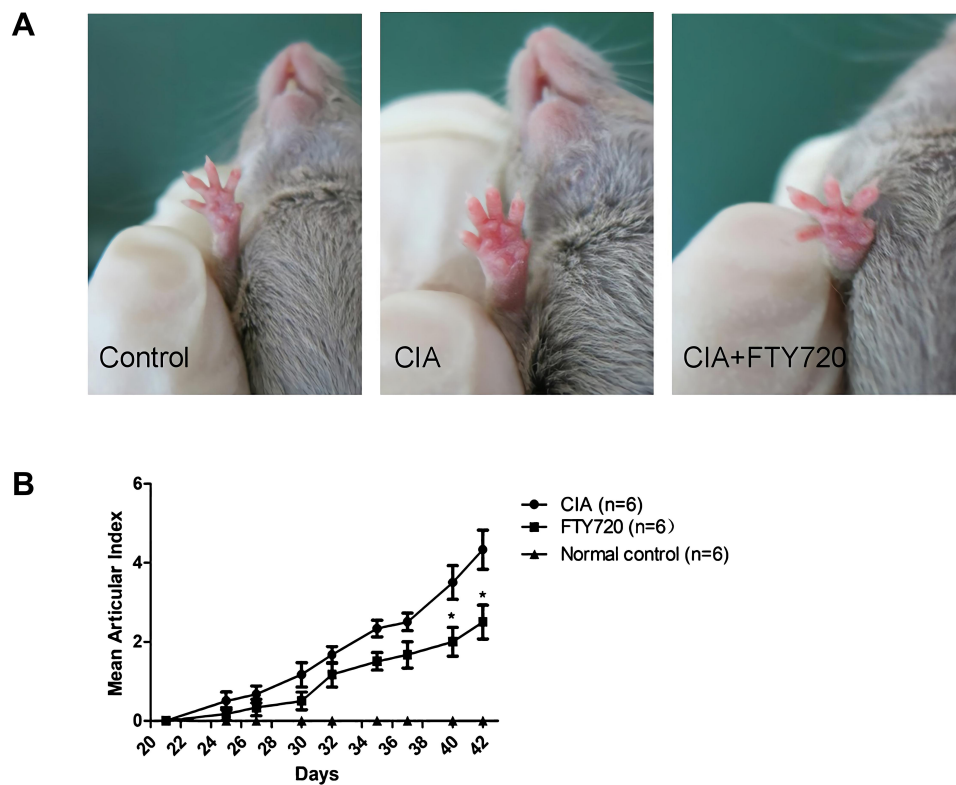


Figure 1 Influences of FTY720 treatment on CIA. Rheumatoid arthritis was induced in DBA/1 mice by a CFA injection at the base of the tail on day 0, followed by a booster injection at day 21, and divided in CIA model group (n = 6) that received saline treatment (CIA), or FTY720 treatment (CIA +FTY720; 2 mg/kg/d). Treatments started the day before immunization until the end of the experiment. Representative images of articular index recorded every 2–3 days after the booster injection until the end of the experiment (**A** and **B**). *P < 0.05 for comparisons between the CIA model group and the FTY720 treatment group.

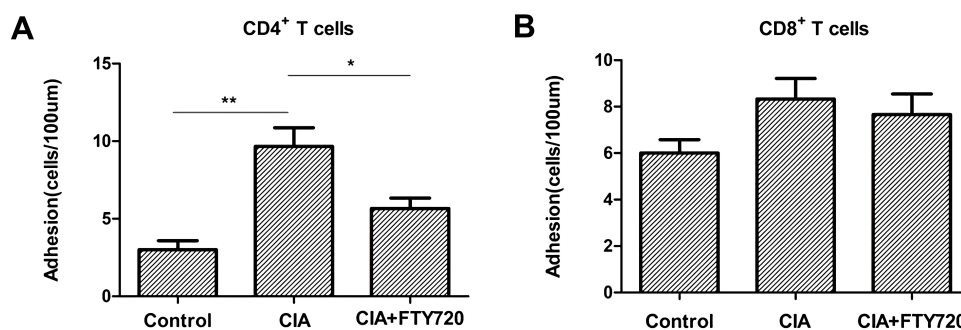


Figure 2 (A) The recruitment of CD4⁺ T lymphocytes in the peripheral vessels of lower extremity joints in three group mice. **P < 0.01 for comparisons between the CIA model group and the control group, *P < 0.05 for comparisons between the CIA model group and the FTY720 treatment group. **(B)** The recruitment of CD8⁺ T lymphocytes in the peripheral vessels of lower extremity joints in three group mice.

Cytotoxicity of FTY720 on MH7A Cells (IC₅₀)

We investigated the cytotoxicity of FTY720 in MH7A cells by using the MTS assay (Promega, Madison, WI, USA). The cytotoxicity was not observed until the cells were exposed to up to 2 μ M FTY720 for 24 hours (**Figure 4**). Treatment with increasing concentrations of FTY720 for 24 or 48 h resulted in a dose- and

time-dependent reduction in MH7A cell viability (**Table 1**).

Effects of FTY720 on IL-1 β , IL-6, IL-8 mRNA and Protein Expression in MH7A Cells Induced by TNF- α

As shown in **Figure 5A–C**), TNF- α (10 ng/mL) significantly increased the mRNA expression of IL-1 β , IL-6,

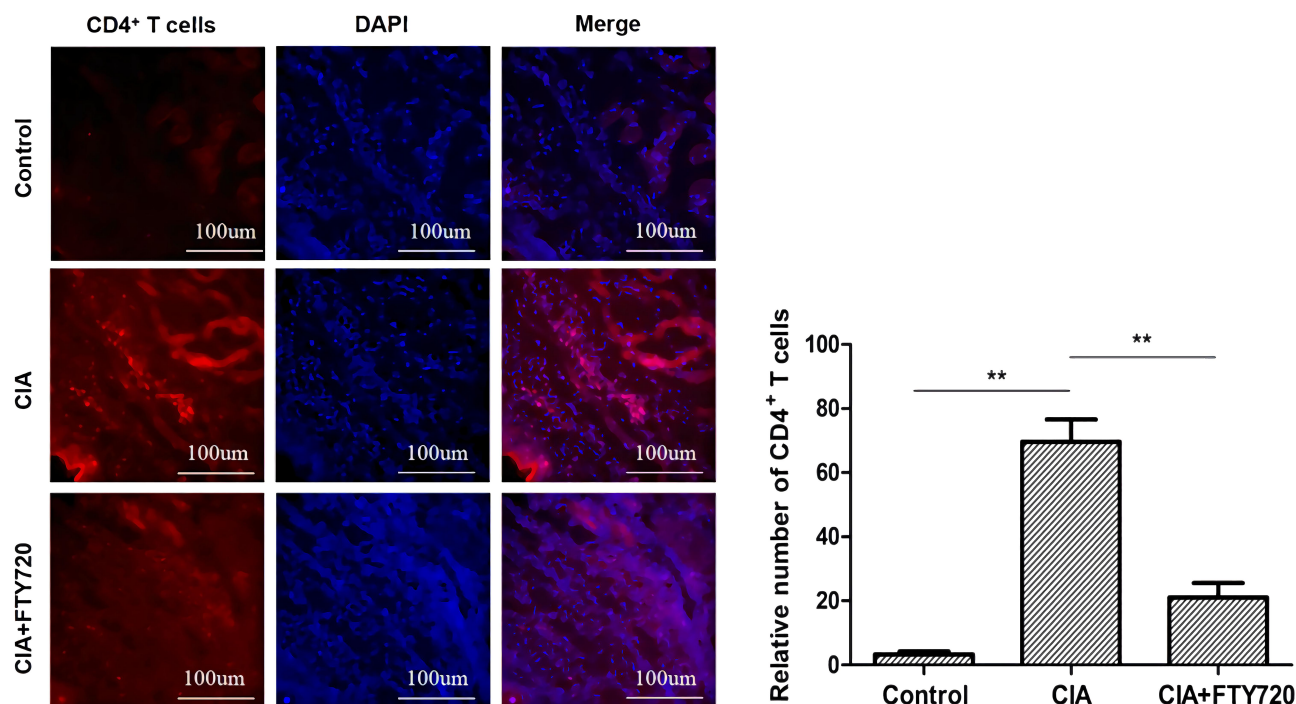


Figure 3 Immunofluorescence analysis of CD4⁺ T lymphocytes in the synovial tissue of each group of mice. The level of CD4⁺ T lymphocytes in the synovial tissue of the CIA model group was significantly higher than that in the normal control group. After FTY720 treatment, the number of CD4⁺ T lymphocytes decreased significantly compared with that in the CIA group. Magnification $\times 200$. The values are expressed as the means \pm SD ($n = 6$). ** $P < 0.01$.

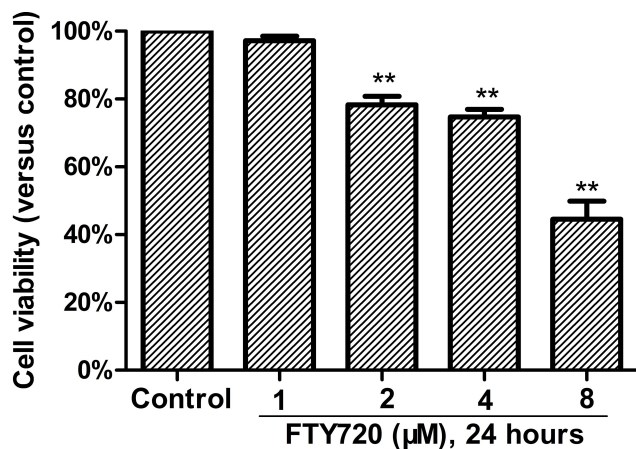


Figure 4 Cytotoxicity of FTY720 on MH7A cells. Cells were treated with different concentrations of FTY720 for 24 h, and the viability was determined using the CellTiter-Blue[®] Cell Viability Assay. ** $P < 0.01$ versus the group without FTY720 treatment. The data are presented as the means \pm standard deviation of three independent experiments.

and IL-8 in MH7A cells compared with that of the control group, which could be attenuated by FTY720 pretreatment in a dose-dependent manner. Similarly, the protein expression of IL-1 β , IL-6 and IL-8 in the supernatant of the control group was low but increased significantly with TNF- α stimulation (10 ng/mL), and FTY720 decreased the TNF- α -induced increases in

these factors in a dose-dependent manner (Figure 5D–F).

FTY720 Attenuated the TNF- α -Induced Activation of NF- κ B in MH7A Cells

NF- κ B pathway plays a key role in inflammatory responses and NF- κ B is activated by phosphorylated I κ B- α in previous studies. Therefore, the expression levels of p65 and p-p65 in whole cell extracts and phosphorylated I κ B- α in the cytosolic fraction were detected by Western blotting. As shown in Figure 6A and B, FTY720 treatment inhibited the TNF- α -induced expression of p-p65 and p-I κ B α in MH7A cells compared with that in control cells. We also found that TNF- α -induced NF- κ B p65 nuclear translocation was significantly inhibited by treatment with FTY720 (Figure 6C)

Table 1 IC₅₀ and IC₈₀ of FTY720 for MH7A Cells

Time	IC ₅₀ (μM)	IC ₈₀ (μM)
24 h	7.38	11.87
48 h	5.77	8.91

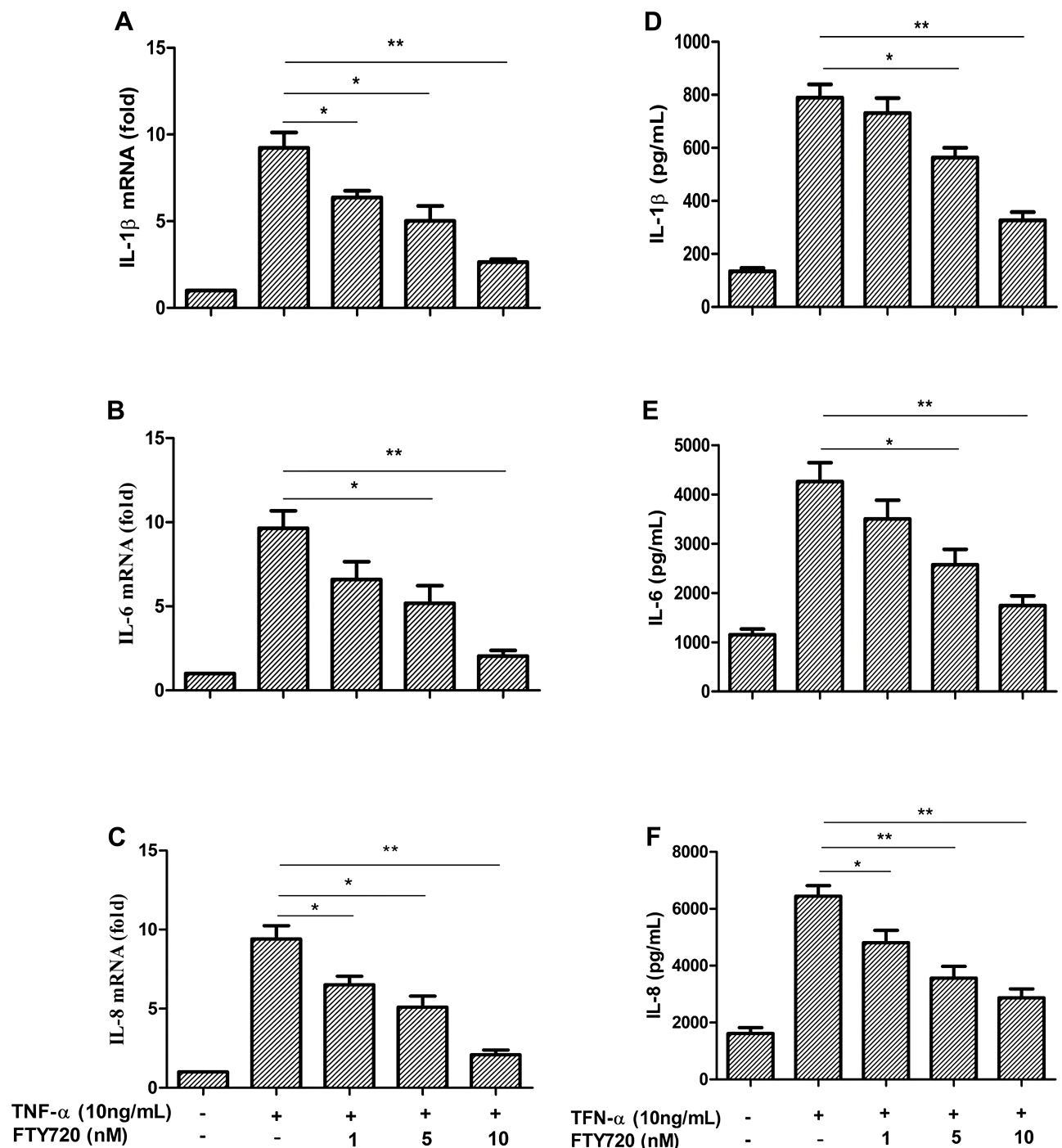


Figure 5 The TNF- α -induced IL-1 β , IL-6, and IL-8 mRNA were suppressed by FTY720 and protein could be found in MH7A cells.

Notes: (A–C) MH7A cells were pretreated with FTY720 or not for 2 h and then stimulated with TNF- α (10 ng/mL). The mRNA levels of IL-1 β , IL-6, and IL-8 were monitored by qPCR after 8 hours. The results are showed as the mean \pm SD of three independent experiments. * P < 0.05, ** P < 0.01 versus the group which was treated with TNF- α alone. (D–F) MH7A cells were pretreated with FTY720 or not for 2 h and then stimulated with TNF- α (10 ng/mL). The protein levels of IL-1 β , IL-6, and IL-8 were monitored in the culture medium by enzyme-linked immunosorbent assay after 24 hours. The results are presented as the mean \pm SD of three independent experiments. * P < 0.05, ** P < 0.01 versus the group were treated with TNF- α alone.

FTY720 Inhibited TNF- α -Induced PI3K/Akt Activation in MH7A Cells

We further examined the effect of FTY720 on the PI3K/Akt pathway by measuring the expression levels

of Akt, p-Akt, p85, and p-p85 using Western blotting. As shown in Figure 7, FTY720 treatment decreased the expression levels of p-Akt and p-p85 compared with those of the control. These results indicated that

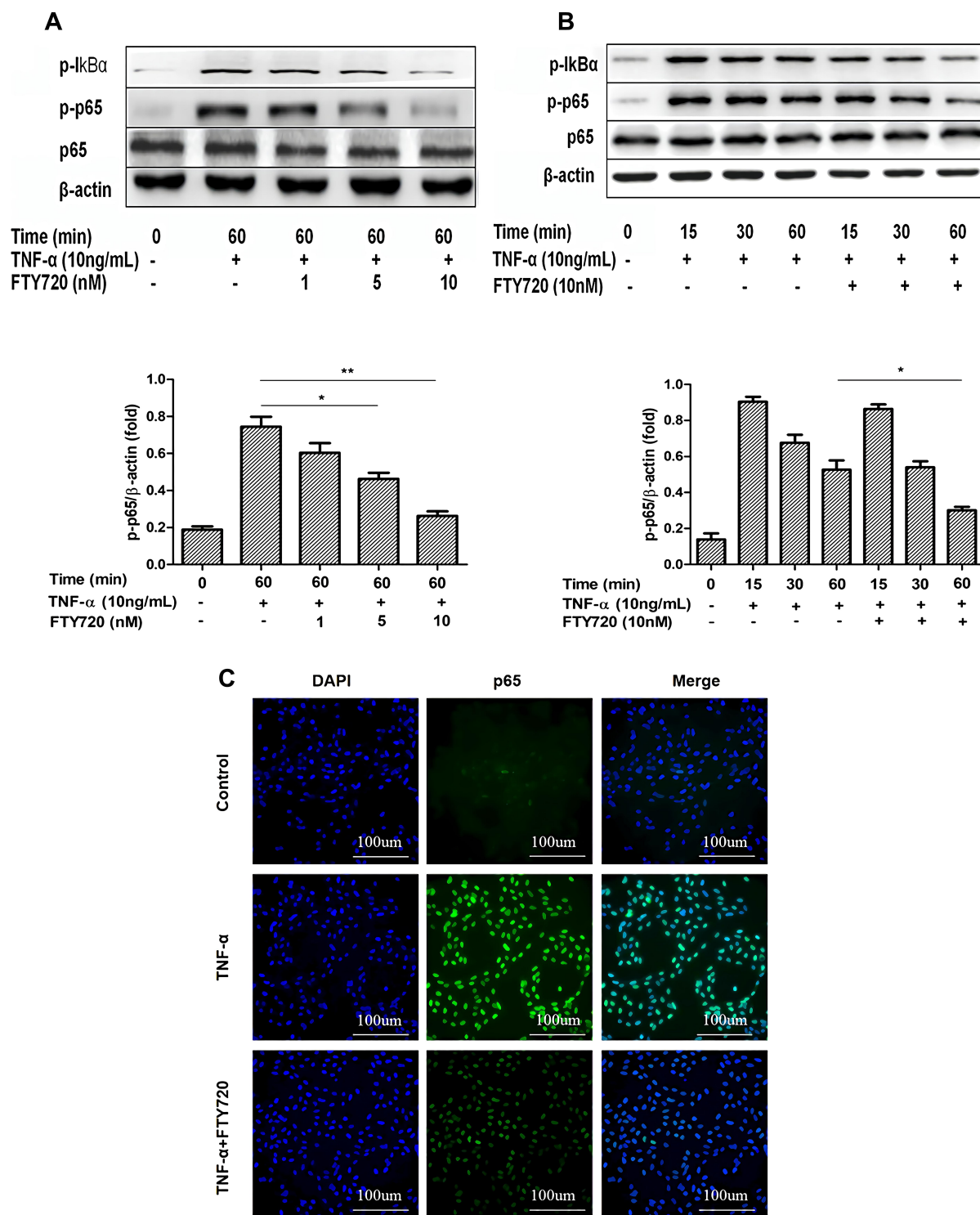
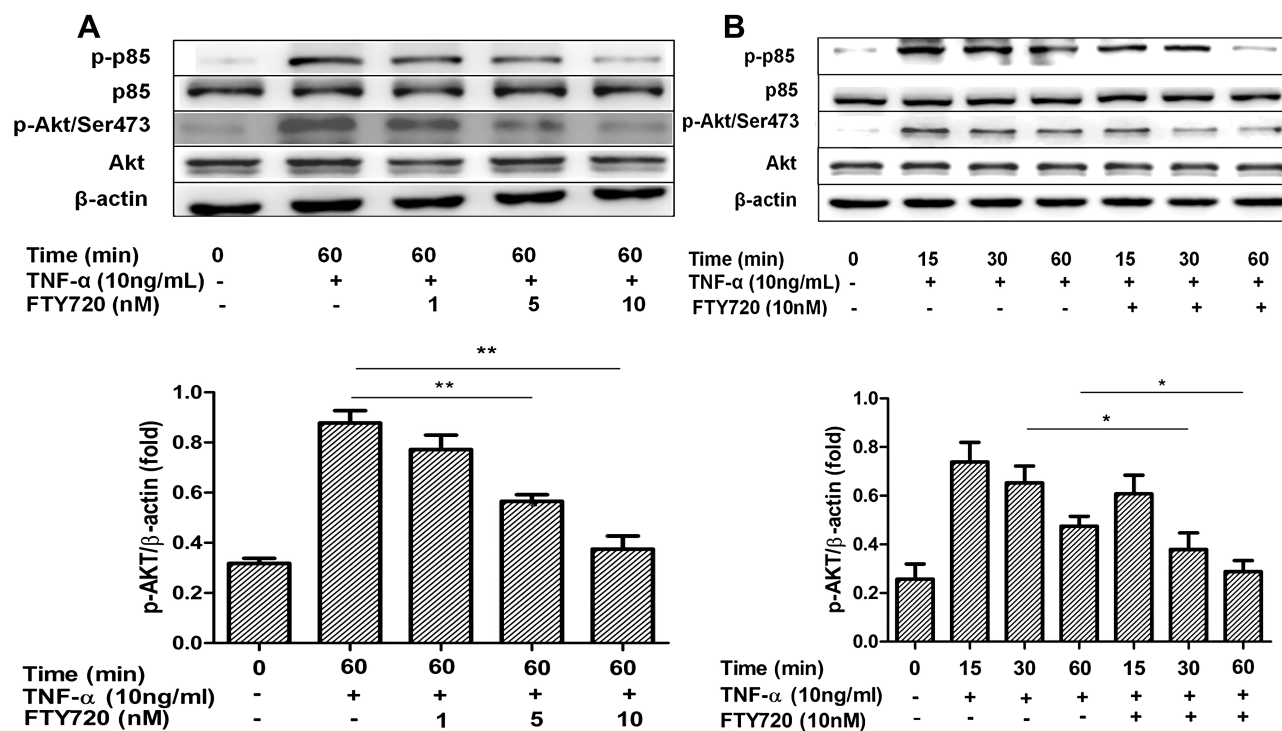


Figure 6 (A–C) FTY720 attenuates the TNF- α -induced activation of NF- κ B in MH7A cells.

Notes: (A) MH7A cells were pretreated with FTY720 or not (in a dose-dependent manner) for 2 h, MH7A cells were then incubated with TNF- α (10 ng/mL) for 60 min. Using Western blotting measured p65, phospho-p65 and phospho-I κ B α . The results are shown as the mean \pm SD of three independent experiments. * P < 0.05, ** P < 0.01 versus the group treated with TNF- α alone. (B) MH7A cells were pretreated with FTY720 or not (10 nM) for 2 hours, the MH7A cells were then incubated with TNF- α (10 ng/mL) for 15, 30, and 60 min. p65, phospho-p65 and phospho-I κ B α were monitored with Western blotting. The consequences are shown as the mean \pm SD of three independent experiments. * P < 0.01 versus the group treated with TNF- α alone. (C) After pretreatment with FTY720 or not (10 nM) for 2 h, and then MH7A cells were incubated with TNF- α (10 ng/mL) for 30 min. The nuclear translocation of NF- κ Bp65 was measured by confocal laser fluorescence microscopy. Cells were immunostained with NF- κ Bp65 antibodies labelled with FITC (green), and nuclei were stained with DAPI (blue). The consequences are representative of three independent experiments. Bar: 10 μ m.



FTY720 treatment inhibited the PI3K/Akt pathway in MH7A cells.

Discussion

Sphingosine is one of the components of the cell membrane. Sphingosine is phosphorylated to form sphingosine-1-phosphate (S1P), which is catalysed by sphingosine kinase 1 (SphK1) and sphingosine kinase 2 (SphK2) *in vivo*.⁴ S1P is synthesized in the cell and transported to the extracellular environment or cell surface by transporters.¹⁴ S1P can bind to sphingosine-1-phosphate receptors (S1P1-S1P5) on the cell membrane surface by autocrine or paracrine signalling and mediate various biological effects, such as cell activation, proliferation, differentiation, migration and angiogenesis.^{15,16}

There was a high level of S1P receptor in the peripheral blood of rheumatoid arthritis patients.¹⁷ Lai et al found that the expression of S1P in the synovial fluid of RA patients was 5 times higher than that of the synovial fluid of normal patients.¹⁸ Kitano found that S1P could independently enhance TNF- α - and IL-1 β -induced production of COX-2 and PGE2 in MH7A cells;¹⁹ Zhao found

that S1P could induce rheumatoid synovial cells to secrete inflammatory cytokines (IL-6 and IL8), and S1P could promote the proliferation of rheumatoid synovial cells through anti-apoptotic effects.²⁰ Takeshita found that S1P/S1P1 signalling plays a role in the pathogenesis of RA by regulating the expression of RANKL. S1P can independently enhance the expression of RANKL in MH7A cells and has a synergistic effect with TNF- α .²¹ Because of the role of S1P/S1P1 signalling in the occurrence and development of RA, S1P/S1P1 is a new target in terms of the treatment of rheumatoid arthritis.²² The phosphorylated form of FTY720 is a structural analogue of S1P, which can be recognized by the S1P receptor at very low concentrations (10^{-9} M) and can suppress the downstream signal of S1P1.²³ In this study, FTY720 was shown to inhibit CIA development through both *in vitro* and *in vivo* experiments.

In the process of inflammation, circulating immune cells roll, adhere, crawl and cross the vascular wall on the surface of the vascular endothelium and finally migrate to the tissue parenchyma.²⁴ This process widely exists in the migration of immune cells, tumour cells and other

cells. We can directly observe and analyze the behaviours of immunocytes labelled by fluorescence through visualization by exposing the blood vessels of the affected joints of anaesthetized animals to intravital microscopy.

The mechanism by which FTY720 is believed to play a strong immunosuppressive role is through the reversible and dose-dependent reduction in the number of peripheral blood lymphocytes, especially CD4⁺ T lymphocytes.⁶ FTY720 has immunomodulatory effects on dendritic cells (DCs), innate lymphoid cells (ILCs) and has important consequences for cytokines that have major impact on RA pathogenesis.^{25,26} Several studies have presented that FTY720 can decrease inflammation in a murine rheumatoid arthritis model. Tsunemi et al found that FTY720 could inhibit arthritis in an SKG mouse RA model via the sequestration of autoimmune CD4⁺ T lymphocytes in the thymus.⁶ Han et al demonstrated that FTY720 inhibited arthritis in CIA mice by hindering DC migration to local lymph nodes.¹² However, the recruitment of immune cells in the affected joint after FTY720 treatment in CIA mice was not well understood. Our results suggested that FTY720 inhibited the recruitment of CD4⁺ T lymphocytes in the lower extremity joint in CIA mice. Increasing evidence shows that the occurrence and development of RA is the result of many abnormal lymphocyte effects, among which the imbalance in CD4⁺ T lymphocyte subsets is an important factor leading to the occurrence and development of RA.^{27,28} We also confirmed that the infiltration of CD4⁺ T lymphocytes in the synovium of CIA mice was significantly higher than that of normal subjects. The mechanism of action of FTY720 is believed to be mainly through the reversible and dose-dependent reduction in the number of peripheral blood lymphocytes, especially CD4⁺ T lymphocytes, downregulating the ratio of CD4⁺/CD8⁺ lymphocytes. The S1P/S1P receptor axis has been found to control the transport and migration of various types of immune cells, such as lymphocytes, neutrophils and macrophages.^{29,30} Eken et al found that when S1P receptor 1 is knocked out other disease models such as EAE does not develop and the Th17 cells are unable to home to tissues.²⁵

In our study, by using intravital microscopy, we observed a significant reduction in the number of infiltrating CD4⁺ lymphocytes in the peripheral vessels of the lower extremity joint in CIA mice after FTY720 treatment. We also found that after FTY720 treatment, the number of CD8⁺ lymphocytes adhering to the vessels around the

joints in CIA mice decreased, but the difference was not statistically significant.

FLSs play a key role in the development of RA.^{31,32} Tsunemi et al found that FTY720 inhibited PGE2 production by MH7A cells and cytokine expression in synovial tissues of a mouse RA model.⁶ The results of our study showed that in MH7A cells, FTY720 inhibited TNF- α -induced production of inflammatory cytokines. Furthermore, our results also demonstrated that the anti-inflammatory properties of FTY720 were mediated by the inhibition of the Akt/PI3K/NF- κ B signalling pathway.

The significance of RA treatment is to control inflammation and pain.³³ The pathogenesis of rheumatoid synovitis is closely related to the release of cytokines such as TNF- α , IL-1 β , IL-6 and IL-8. The higher the concentration of cytokines, the more serious the clinical symptoms and the more active the RA. The levels of TNF- α , IL-1 β , IL-6 and IL-8 were significantly reduced after DMARD treatment. Our results showed that FTY720 significantly inhibited the production of TNF- α -activated inflammatory cytokines, such as IL-1 β , IL-6 and IL-8, at the mRNA and protein levels in MH7A cells. In addition, we found that the effective concentration of FTY720 on MH7A cells was 0.001 μ M to 0.01 μ M, far lower than the IC50 value of FTY720 against the proliferation of MH7A.

The transcription factor NF- κ B plays an important role in inflammation.³⁴ Several studies have shown that NF- κ B is stimulated in the development of RA.^{35,36} It is well established that NF- κ B is activated via the canonical pathway in RA pathogenesis. In the resting stage, the NF- κ B protein dimer (p65/p50) is complexed with the inhibitory subunit I κ B α and is sequestered in the cytoplasm. After stimulation with TNF- α , I κ B α is rapidly phosphorylated by IKK α / β at Ser32 and Ser36 residues and subsequently ubiquitinated and degraded by the proteasome. Once released from I κ B, free NF- κ B translocates to the nucleus and initiates the transcription of numerous inflammatory gene products. In this study, we found that FTY720 remarkably restrained I κ B α and NF- κ Bp65 phosphorylation induced by TNF- α and abated nuclear NF- κ Bp65 levels in MH7A cells.

The PI3K/Akt pathway is a major regulator of inflammation in inflammatory diseases.³⁷ Moreover, PI3k/Akt axis has also been shown in the context of Treg cells.³⁸

Several studies have shown that the PI3K/Akt signalling pathway probable the principal upstream pathway of the NF- κ B signalling cascade.^{37,39} Previous studies have shown that TNF- α could activate the PI3K/Akt pathway in synovial cells.⁴⁰ Interestingly, our results showed that

FTY720 could inhibit the PI3K/Akt pathway, which was accompanied by the suppression of I κ B α and NF- κ B phosphorylation. Other studies have shown that the phosphorylation of NF- κ Bp65 was also hindered by the PI3K inhibitor LY294002 in TNF- α -activated MH7A cells. Based on these results, we hypothesised that FTY720 suppressed NF- κ B activity through the PI3K/Akt signalling pathway in MH7A cells.

In an inflammatory environment, endothelial cells express high levels of adhesion molecules. P65 and Akt are key signalling molecules that mediate endothelial cell activation. Zhao et al found that in an LPS-induced central nervous system (CNS) inflammation model, the phosphorylation of FTY720 could inhibit the phosphorylation of Akt and NF- κ B in vivo and in vitro by binding to S1P1 on the surface of endothelial cells, thus significantly inhibiting the activation of mouse cerebral vascular endothelial cells.⁴¹ Kitano et al also found that S1P1 was upregulated in vascular endothelial cells in RA.¹⁹ Based on these conclusions and our experimental results, we speculated that FTY720 could inhibit the recruitment of CD4⁺ lymphocytes by inhibiting S1P/S1P1 signalling and restraining the activation of vascular endothelial cells in CIA mice, but further research is needed to confirm this hypothesis.

In conclusion, our study showed that FTY720 reduced CD4⁺ lymphocyte adhesion in peripheral vessels of lower extremity joints in CIA mice, and in vitro experiments showed that FTY720 reduced the production of inflammatory cytokines by inhibiting the PI3K/Akt/NF- κ B signalling pathways in MH7A cells. We speculated that the inhibitory effect of FTY720 on CD4⁺ lymphocyte recruitment in RA was related to the inhibition of p65 and Akt phosphorylation; however, further experiments are needed to clarify the mechanism.

Acknowledgments

The study was supported by the Non-profit Central Research Institute Fund of the Chinese Academy of Medical Sciences (2019PT320001) and the Major Research Plan of National Natural Science Foundation of China (91949203).

Disclosure

The authors reported no conflicts of interest for this work.

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