Kirenol inhibits TNF-α-induced proliferation and migration of HaCaT cells by regulating NF-κB pathway

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Abstract

Psoriasis is a common chronic, inflammatory skin disease possessing properties of inflammatory cell infiltration and excessive proliferation of keratinocytes, the occurrence and development of which remain fully elucidated. Therefore, the study was designed to determine the effects of kirenol (50, 100 and 200 μg/mL) on Cultured Human Keratinocytes (cells) (HaCaT) in vitro and reveal its molecular mechanism. The in vitro psoriasis model was established utilizing tumor necrosis factor-α (TNF-α)-stimulated HaCaT cells. Kirenol, a diterpenoid compound, was applied at different concentrations (50, 100 and 200 μg/mL) to HaCaT cells for 24 h. The Cell Counting Kit-8 (CCK-8) and thymidine monobromodeoxyuridine (BrdU) assays were used to assess cell viability and proliferation, followed by assessment of cell migration by Transwell assay. Subsequently, inflammatory cytokines were measured by enzyme-linked immunosorbent assay (ELISA), and Western blot assay was used to evaluate expressions of p65, p-p65, IκBα and p-IκBα. Activities of superoxide dismutase (SOD), catalase (CAT), glutathione (GSH) and malondialdehyde (MDA) contents were measured spectrophotometrically. The results demonstrated that TNF-α induced a significant increase in cell viability and inflammatory cytokines, including expressions of Interleukin (IL)-6, IL-8, IL-22 and IL-1β in HaCaT cells, which was dose-dependently inhibited by kirenol. Similarly, TNF-α-induced cell migration was also suppressed by kirenol treatment. Furthermore, TNF-α stimuli induced the upregulation of phosphorylation levels of p65 and IκBα as well as p-p65–p65 and p-IκBα–IκBα ratios, whereas kirenol significantly suppressed the activation of cellular nuclear factor-kappa B (NF-κB) signaling pathway. In addition, kirenol significantly decreased the level of MDA but increased the levels of SOD, CAT and GSH in a dose-dependent manner. These results proposed that kirenol could inhibit the proliferation, migration, expression of inflammatory factors, and oxidative stress in HaCaT cells via suppressing NF-κB signaling pathway.

Keywords: kirenol; psoriasis; proliferation; migration; inflammation

Introduction

Psoriasis is a recurrent, noninfectious erythema scaly skin disease characterized by epidermal proliferation and inflammation. Its etiology is related to heredity, infection, allergy, metabolic disorders, and autoimmunity (Allergologia et immunopathologia, 2020; Aquino et al., 2021; Evans et al., 2020; Srivastava et al., 2021; Zhou et al., 2020). It is a typical chronic inflammatory disease with an increasing incidence ranging from 1 % to 3 % worldwide (Griffiths & Barker, 2007; Hoegler et al., 2018; Langley et al. 2005). At present, psoriasis is an extremely complicated pathophysiologic process, the occurrence and development of which remain unclear (Boehncke & Schön, 2015). Currently, psoriasis is believed to mainly involve genetic, psychological, immunologic, and...
environmental factors (Yang & Zheng, 2020). Moreover, the principle obsessive progressions from psoriasis hold numerous different features in keratinocyte dysplasia, keratosis, neovascularization as well as inflammatory cell infiltration (Kim et al., 2014a). At the same time, excessive proliferation and abnormal migration of keratinocytes leading to the microenvironment of psoriasis lesions are vital features of psoriasis (Winchell & Watts, 1988). As psoriasis is prone to relapse and challenging to cure, it is conducive to exploring new therapeutic targets related to keratinocyte proliferation as well as migration for the treatment of psoriasis.

As the primary pathogenic cells in psoriasis, keratinocytes is considered as a promoter for amplification of inflammatory response with the additional production of tumor necrosis factor-α (TNF-α) in the initiation of disease (Nickoloff, 2006). Following stimulation with TNF-α, keratinocytes may provoke the abbrrent expression of pro-inflammatory cytokines and chemokines such as IL-6, IL-8, IL-22 and IL-1β, which contribute to the subsequent recruitment of immune cells into inflammatory lesions in the skin (Cho, Lee, & Kim, 2007; Lee, Chung, Kang, Park, & Hyun, 2019; Wolk et al., 2009). It is believed that there are multiple signaling pathways involved in the inflammatory response, among which the nuclear factor-kappa B (NF-κB) pathway plays a decisive role (Nair et al., 2009; A. Wang et al., 2019; Xu, Xu, Xiong, & Deng, 2019). During the signaling pathway, inflammatory stimuli activate IKK, promoting phosphorylation and degradation of IκB and subsequent activation of NF-κB signaling pathway, which results in a series of inflammatory responses (Lawrence, 2009; Zhang, Lenardo, & Baltimore, 2017). Recognizing the certainty that NF-κB signaling pathway is involved in the progression of psoriasis, inhibition of the pathway is beneficial to reduce hyperproliferation and hyper inflammation in keratinocytes. Therefore, inflammatory factor (IL-6/IL-8/IL-22/IL-1β) and NF-κB were chosen to investigate the effect of kirenol on psoriasis in this study.

Kirenol is a diterpenoid characteristic product compound extracted from herba siegesbeckiae (Ibrahim et al., 2021; Wang et al., 2021a). In addition, kirenol attenuates the inflammatory response to collagen-induced arthritis by inhibiting the activation of signaling pathways and reducing the expression of pro-inflammatory cytokines such as IL-1β (Wang et al., 2011). Moreover, kirenol inhibits inflammatory factors and improves ulcerative colitis in mice (Xiu Hong et al., 2019). Besides, kirenol inhibits rheumatoid arthritis (RA)-associated synovial fibroblasts proliferation, migration, invasion, and inflammation (Wu et al., 2019). Although kirenol has also been reported to have immunomodulatory and anti-inflammatory properties, the efficacy of kirenol on psoriasis has not been further investigated. In the present study, we investigated the effects of kirenol on psoriasis development and its mechanism of function. The results revealed that kirenol significantly inhibited the proliferation capacity and migration ability of Cultured Human Keratinocytes (cells) (HaCaT) through regulating the NF-κB pathway.

Materials and Methods

Cell culture

Cultured human immortalized keratinocytes (HaCaT; Procell, Wuhan, China) were cultured in Dulbecco’s Modified Eagle medium (DMEM), supplemented with 10% fetal bovine serum and 1% penicillin–streptomycin in a humidified incubator at 37°C and 5% CO₂. HaCaT cells in a logarithmic growth phase were induced with 10 ng/mL TNF-α for 48 h to establish an in vitro cell model for psoriasis at 37°C and 5% CO₂. The kirenol (Yuan-ye Bio-Technology Co. Ltd, Shanghai, China) concentration was increased to 50, 100 and 200 μg/mL for 24 h at 37°C and 5% CO₂. Furthermore, the cells were separately divided in the following five groups: the control group; the TNF-α group; the TNF-α + kirenol (50 μg/mL) group, the TNF-α + kirenol (100 μg/mL) group and the TNF-α + kirenol (200 μg/mL) group. Dimethyl sulfoxide (DMSO; Merck, Germany) was used in equal concentrations as a negative control to dilute kirenol.

Cell viability assay

The antiproliferative impact of kirenol concerning HaCaT was determined utilizing Cell Counting Kit-8 reagent (CCK-8; Bestbio, Shanghai, China). For a short period of time, HaCaT cells were inoculated in 96-well plates at a density of 5 × 10⁵ cells/well overnight, treated with different concentrations of kirenol (50, 100, 200 and 400 μg/mL) for 24 h, and incubated with CCK-8 reagent in dark for 1 h at 37°C. After TNF-α treatment for 48 h at 37°C and 5% CO₂, kirenol at concentrations of 50, 100, 200 μg/mL was applied to HaCaT cells. The absorbance of the tests was recorded at a wavelength of 450 nm utilizing a Microplate Reader (Infinite M200, Tecan, Austria).

Measurement of proliferation by incorporation of thymidine monobromodeoxyuridine (BrdU)

Cell proliferation was investigated by thymidine monobromodeoxyuridine (BrdU) under de novo-synthesized DNA conditions utilizing the BrdU Cell Proliferation assay kit (Santa Cruz Biotechnology, Santa Cruz, CA). Briefly, HaCaT cells were seeded into 96-well plates at 2 × 10³ cells/well, and incubated at 37°C for 48 h. Subsequently, 20-mL BrdU was added, and incubated...
at 37°C for 4 h. After incubation with anti-BrdU antibody and peroxidase-conjugated goat anti-mouse immunoglobulin G (IgG), successively, 100-μL 3,3′,5,5′-Tetramethylbenzidine (TMB) peroxidase substrate was added and the mixture was kept in dark at room temperature for 30 min. The absorbance at 450/550 nm was obtained using a microplate reader.

**Measurement of superoxide dismutase (SOD) activity**

The mixture containing 0.033-mM ethylenediaminetetraacetic acid (EDTA), 3.3-mM methionine, 0.33-μg/mL riboflavin and 0.01-mM KCN (Sigma, St. Louis, MO, USA) was applied to supernatant. Cuvette containing the reaction mixture was placed in a box illuminated with 20-W neon lamp for 10 min. One unit of SOD enzyme activity is defined as the amount of enzyme required for inhibiting the chromogen production (optical density at 560 nm) by 50% in 1 min under assay conditions and expressed as specific activity in units/min/mg protein.

**Measurement of glutathione (GSH) activity**

The GSH level was measured by reacting with 5,5′-dithiobis(2-nitrobenzoic) acid (DTNB; Sigma, St. Louis, MO, USA), producing a yellow-colored compound, 5′-thio-(2-nitrobenzoic) acid (TNBA). Concentration of TNBA was measured at 412-nm wavelength, and the amount of GSH was expressed in terms of nmol GSH/mg protein.

**Measurement of catalase (CAT) activity**

The CAT activity was determined by splitting H$_2$O$_2$ for 1 min. The reaction was stopped by dichromate/acetate acid reagent, and the remaining H$_2$O$_2$ was measured at 570 nm for chromic acetate. The CAT activity was expressed as μmol/L H$_2$O$_2$ decomposed/min/mg protein.

**Measurement of malondialdehyde (MDA) activity**

The MDA activity was determined by a Lipid Peroxidation MDA assay kit (Beyotime Institute of Biotechnology, Haimen, China) as recommended by the manufacturer. After treatment with kirenol (50, 100, 200 μg/mL) for 24 h, HaCaT cells were harvested, lysed by cell lysis buffer and centrifuged at 12,000 g for 15 min at 4°C. The obtained supernatant was reacted with thiobarbituric acid, and the reaction products were measured with a microplate reader at 532 nm. Total protein level was determined by BCA protein detection kit (Beyotime Biotechnology Institute, Haimen, China), and MDA level was normalized.

**Enzyme-linked immunosorbent assay (ELISA)**

The TNF-α-induced HaCaT cells were treated with kirenol (50, 100, 200 μg/mL) for 24 h. Then, inflammatory factors (IL-6, IL-8, IL-22 and IL-1β) in the supernatant were detected by ELISA kit (eBioscience, CA, USA) following the manufacturer’s instructions. Briefly, 50 mL of standard, samples, and blank control were added to a 96-well plate coated with primary antibodies against cytokines, including IL-6 (Cat No. ab178013; Abcam, Cambridge, MA, USA), IL-8 (Cat No. ab214030; Abcam, Cambridge, MA, USA), IL-22 (Cat No. ab216170; Abcam, Cambridge, MA, USA) and IL-1β (Cat No. ab214025; Abcam, Cambridge, MA, USA), followed by incubation with HRP-conjugated detection antibodies (100 mL) for 1 h at 37°C. After washing for five times, substrates A and B were added for further 15-min incubation. Finally, the absorbance was measured at 450 nm in a microplate reader. A standard curve was produced to quantify the concentration of cytokines from different samples.

**Transwell assay**

To induced cell migration, CoStar Transwell chambers (8-μm pore size; Corning, Costar, NY, USA) were used. TNF-α-induced HaCaT cells (1 × 10$^5$/well) were inoculated in the wells of upper chambers filled with 200-μL serum-free DMEM medium, and lower chambers were filled with 500-μL medium, which contained 20% fetal bovine serum. After incubation at 37°C for 4 h. After incubation with anti-BrdU antibody and peroxidase-conjugated goat anti-mouse immunoglobulin G (IgG), successively, 100-mL 3,3′,5,5′-Tetramethylbenzidine (TMB) peroxidase substrate was added and the mixture was kept in dark at room temperature for 30 min. The absorbance at 450/550 nm was obtained using a microplate reader.

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**Western blotting assay**

For Western blot analysis, the TNF-α-stimulated HaCaT cells were treated with kirenol (50, 100, 200 μg/mL) for 24 h. Proteins were extracted from the cells grown on 35-mm culture dishes using the Trizol protein isolation method. Bradford’s assay quantified the isolated proteins. The membrane was blocked at room temperature for 2 h and incubated overnight at 4°C with the following primary antibodies: anti-p65 (ab19870, 1:500; Abcam, Cambridge, MA, USA), anti-p-p65 (ab194726, 1:500; Abcam, Cambridge, MA, USA), anti-IκBα (ab247825, 1:500; Abcam, Cambridge, MA, USA), and anti-β-actin (ab8226, 1:500; Abcam, Cambridge, MA, USA). Finally, the absorbance was measured at 450 nm in a microplate reader. A standard curve was produced to quantify the concentration of cytokines from different samples.
Molecular mechanism of kirenol in psoriasis

Results

Kirenol restrained TNF-α-induced proliferation of HaCaT cells in a dose-dependent manner

Cultured human keratinocytes were first treated with 10-ng/mL TNF-α for 48 h to simulate the epithelial state of psoriasis. The proliferation of HaCaT cells was detected by CCK-8 and BrdU assays to verify the effects of kirenol on TNF-α-induced proliferation of HaCaT cells. Then HaCaT were incubated with the following gradient concentrations of kirenol: 50, 100 and 200 μg/mL for 24 h. The following five groups were used in the experiment, including the control group: TNF-α group, TNF-α + kirenol (50 μg/mL) group, TNF-α + kirenol (100 μg/mL) group and TNF-α + kirenol (200 μg/mL) group. Figure 1A shows that kirenol concentrations of 50, 100 and 200 μg/mL were non-toxic to HaCaT cells. The CCK-8 assay results indicated

Statistical analysis

The mean ± standard deviation (SD) was calculated for each experimental group. Statistical comparisons of the data were analyzed using the SPSS software (SPSS 26, Chicago, IL, USA). Student’s t-test was used to compare the two groups. Differences between the groups were statistically significant (P < 0.05).

Figure 1. Kirenol inhibited TNF-α-induced proliferation of HaCaT cells in a dose-dependent manner. (A) Kirenol at concentrations of 50, 100 and 200 μg/mL was nontoxic to HaCaT cells. (B) Kirenol reduced cell viability in a dose-dependent manner. (C) Kirenol increased BrdU positive cells in a dose-dependent manner. (D) The percentage of cells with BrdU-positive (BrdU+) staining was determined. **P < 0.01 vs. control; ***P < 0.001 vs. control; #P < 0.05, ##P < 0.01 vs. TNF-α alone. TNF-α: tumor necrosis factor-α.
that kirenol markedly inhibited the viability of HaCaT cells in a dose-dependent manner in comparison to the control group (Figure 1B). In addition, the BrdU assay illustrated that TNF-α-induced proliferation of HaCaT cells was significantly suppressed by kirenol in a dose-dependent manner compared to the control group (Figure 1C).

Kirenol improved TNF-α-induced inflammation of HaCaT cells

In order to determine the impact of kirenol (50, 100, and 200 μg/mL) on TNF-α-induced inflammation of HaCaT cells, ELISA was performed to measure inflammatory cytokines associated with the development of inflammation, such as IL-6, IL-8, IL-22 and IL-1β. The ELISA data revealed that compared with the control group, the expressions of inflammation factors were significantly increasing in the TNF-α alone group. In contrast, they were reduced considerably by kirenol in a concentration-dependent manner. The results revealed that kirenol improved TNF-α-induced inflammation of HaCaT cells in a dose-dependent manner (Figure 2).

Kirenol inhibited oxidative stress in TNF-α-stimulated HaCaT cells

Oxidative stress is the main factor for psoriasis. Therefore, oxidative stress assay was performed and the results displayed that TNF-α increased MDA but decreased SOD, CAT and GSH, indicating that TNF-α stimuli promoted oxidative stress. However, application of kirenol significantly decreased the level of MDA but increased the levels of SOD, CAT and GSH in a dose-dependent manner (Figure 3). Collectively, kirenol inhibited oxidative stress of HaCaT cells.

Kirenol suppressed TNF-α-induced migration of HaCaT cells

In order to investigate the role of kirenol (50, 100, and 200 μg/mL) in the process of TNF-α-induced migration of HaCaT cells, we further assessed its migration capacity utilizing Transwell assay. Correspondingly, the Transwell migration assay demonstrated that TNF-α enhanced the migratory ability of HaCaT cells compared to the control group. However, it was markedly suppressed by kirenol in a dose-dependent manner. The results established

Figure 2. Kirenol alleviated TNF-α-induced inflammation of HaCaT cells. Kirenol decreased the levels of Interleukin-8 (IL-8), IL-1β, IL-6, and IL-22. ***P < 0.001 vs. control; #P < 0.05 vs. TNF-α alone, ##P < 0.01 vs. TNF-α alone, and ###P < 0.001 vs. TNF-α alone.
Molecular mechanism of kirenol in psoriasis

Discussion

Psoriasis is an extremely complicated pathophysiological process, and the primary pathological characteristics incorporate the inordinate proliferation of keratinocytes and local inflammatory response (Callis Duffin et al., 2017; Cooper, 1990). Currently, the mechanism of etiology and pathogenesis of psoriasis remain unclear. As reported, kirenol has various medicinal properties, including antibacterial, antiviral, anti-inflammatory, anti-arthritic as well as anti-adipogenesis effects (Kim et al., 2014b; Lu et al., 2012; Ren et al., 2020; Wang et al., 2012). Accumulating evidence has proved the biological effects of kirenol on a wide range of cells. For instance, its ability to inhibit cell proliferation and differentiation induces cell apoptosis and alleviates inflammatory response (Wang et al., 2021b; Wu et al., 2019). However, whether kirenol affects psoriasis remains fully understood. The present study found that kirenol inhibited TNF-α-induced phenotypes of HaCaT cells by modulating the NF-κB pathway. To further validate the molecular mechanism by which kirenol (50, 100, and 200 µg/mL) projects its effects of both inhibiting cell activity and migratory capacity by decreasing the activation of NF-κB pathway.

Kirenol decreased the activation of NF-κB pathway in TNF-α-stimulated HaCaT cells

NF-κB pathway was related to various pathological and physiological changes such as cell proliferation, immunity as well as inflammation. For one thing, accumulating evidence has been reported that NF-κB pathway exhibits a vital role in TNF-α-induced proliferation and migration of HaCaT cells. For another, the inflammatory factors such as IL-1β and IL-6 critically rely on the activation and nuclear translocation of NF-κB. Therefore, we speculated that kirenol performs its function by regulating the NF-κB pathway. To further validate the molecular mechanism by which kirenol (50, 100, and 200 µg/mL) projects in the cell, we measure the levels of NF-κB pathway-related proteins such as p65, p-p65, IκBα and p-IκBα. The TNF-α-induced HaCaT cells were lysed in RIPA buffer, which contained phosphatase inhibitors as well as protease, and then analyzed by Western blotting assay. The Western blotting assay established that the phosphorylation levels of p65 and IκBα, and p-p65–p65 and p-IκBα–IκBα ratios increased markedly, and kirenol reduced their phosphorylation levels in a dose-dependent manner (Figure 5). This finding demonstrated that kirenol exerted

Figure 3. Kirenol inhibited the oxidative stress in TNF-α-stimulated HaCaT cells. Kirenol decreased the level of MDA but increased the levels of SOD, CAT, and GSH in a dose-dependent manner. ***P < 0.001 vs. control; #P < 0.05 vs. TNF-α alone, ###P < 0.01 vs. TNF-α alone, and ####P < 0.001 vs. TNF-α alone. TNF-α: tumor necrosis factor-α; MDA: malondialdehyde; SOD: superoxide dismutase; CAT: catalase; GSH: glutathione.

that kirenol suppressed the TNF-α-induced migration of HaCaT cells in a dose-dependent manner (Figure 4).
a previous study that TNF-α stimulation considerably improved the viability of HaCaT cells and upregulated inflammatory factors such as IL-8, IL-1β, IL-6 and IL-22 (Na Takuathung et al., 2017). A recent study has found that the levels of cytokines, including IL-1α/β, IL-6 and IL-8, were remarkably upregulated after TNF-α stimuli, indicating that TNF-α induced significant inflammatory response in HaCaT cells. Mevastatin was also demonstrated to alleviate inflammation (Kim et al., 2021). Meanwhile, the expression of inflammatory factors in the supernatant of TNF-α-stimulated cells was significantly down-regulated by kirenol. The results of ELISA assay

Figure 4. Kirenol suppressed TNF-α-induced migration of HaCaT cells. Kirenol inhibited TNF-α-induced HaCaT cell migration, and representative images of the HaCaT cell migration assay. ***P < 0.001 vs. control; ###P < 0.001 vs. TNF-α alone. TNF-α: tumor necrosis factor-α.

Figure 5. Kirenol inhibited the activation of NF-κB pathway in TNF-α-induced HaCaT cells. Kirenol prevented TNF-α-induced activation of HaCaT cell in NF-κB signaling pathway. Western blotting assay was performed to detect the expression of p-p65, p65, p-IκBα, and IκBα in HACaT cells. Expression of p-p-65 relative to p65, p-p-IκBα relative to IκBα, and p65 or IκBα relative to GAPDH. ***P < 0.001 vs. control; **P < 0.01 vs. TNF-α alone, ***P < 0.001 vs. TNF-α alone, and ###P < 0.001 vs. TNF-α alone. TNF-α: tumor necrosis factor-α; I-κBα: inhibitor of NF-κB; p-IκBα: phosphorylated (p)-inhibitor of NF-κBα.
were consistent with the findings of previous studies, which indicated an excellent anti-inflammatory effect on in vitro model of psoriasis (Lin et al., 2018). To further investigate the role of kirenol, stimulation of HaCaT cells was performed with TNF-α to establish an in vitro model of psoriasis (Liu et al., 2020). In the present study, results of CCK-8 and BrdU assays demonstrated that kirenol restrained cellular phenotypes in a dose-dependent manner, which revealed that the increased cell viability of HaCaT cells induced by TNF-α was considerably reduced by kirenol treatment. Moreover, the finding of Transwell migration assay has pointed out that kirenol could suppress the migration of TNF-α stimulated HaCaT cells.

More importantly, the activated keratinocytes could generate plenty of inflammatory factors which induced cell proliferation and viability of HaCaT cells and, in turn, elevated keratinocyte proliferation and epidermal thickening to form psoriasis (Benhadou et al., 2019). It included positive feedback that excessive proliferation and epidermal thickening appeared alternately to perpetuate psoriatic lesions. NF-κB, a protein transcription factor has salient regulatory roles in immune and inflammatory pathways as well as cellular events, such as apoptosis and proliferation (Hayden & Ghosh, 2011; Patel et al., 2018; Yi et al., 2019). Furthermore, the NF-κB signaling pathway has been reported to assume an essential part in the pathogenesis of psoriasis (Wang et al., 2011). Previous study has revealed that TNF-α stimuli resulted in significant increase of inflammatory cytokines and NF-κB in keratinocytes, whereas application of genistein remarkably repressed TNF-α which induced the nuclear translocation of NF-κB and the phosphorylation of IκBα (p-IκBα), suggesting that inhibition of NF-κB expression and phosphorylation of IκBα could attenuate psoriasis (Wang et al., 2019). In addition, Liu et al. (2020) demonstrated that cimifugin exerted protective effects on psoriasis through inhibiting oxidative stress and inflammatory responses by inactivating NF-κB–MAPK pathway. Meanwhile, the fish scale collagen peptides suppressed inflammatory cytokine expression, reactive oxygen species as well as MAPK–NF-κB pathway (Subhan et al., 2017). The results of Western blot assay demonstrated that increased concentrations of kirenol (50, 100 and 200 μg/mL) significantly down-regulated the expression of p-p65 and p-IκBα, and increased the levels of p65 and IκBα, indicating the increased levels of active phosphorylated NF-κB in psoriasis. However, the impact of kirenol on NF-κB pathway and TNF-α-stimulated HaCaT cells has not been reported. Results of the present investigation revealed that the NF-κB pathway might have been included in the impact of kirenol on TNF-α-induced HaCaT cells. TNF-α stimulation intensified the activation of the NF-κB pathway. At the same time, kirenol considerably repressed NF-κB pathway activation on TNF-α stimulated HaCaT cells.

Generally speaking, the present study discovered that kirenol activity for human keratinocytes played a protective role by inhibiting cell viability and migration as well as inflammatory response through NF-κB signaling pathway in vitro. However, some limitations associated with the study must be mentioned. For one thing, the main limitation is that the study focused more on the results of in vitro examination without in vivo and clinical analysis, which must be attained in the future investigation. For other thing, whether the effects of kirenol on NF-κB signaling pathway were direct or indirect must be identified.

Conclusion

To sum up, the study revealed for the first time the effects of specific concentrations of kirenol on the proliferation, migration and inflammation of TNF-α-stimulated HaCaT cells. Even more profound is the revelation of possible mechanisms of signaling pathways in the efficacy of kirenol. The findings of the study suggest that kirenol is conducive to improving the clinical symptoms of psoriasis, and could be a new therapeutic strategy to treat the scaly skin disease.

Competing Interests

The authors state that there are no conflicts of interest to disclose.

Contribution of Authors

Jin Li and Fang Ren designed the study, supervised the data collection, Wenliang Yan analyzed the data, interpreted the data, Hong Sang prepare the manuscript for publication and reviewed the draft of the manuscript. All authors have read and approved the manuscript.

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Molecular mechanism of kirenol in psoriasis


