Research Article

Inhibition of the PERK/TXNIP/NLRP3 Axis by Baicalin Reduces NLRP3 Inflammasome-Mediated Pyroptosis in Macrophages Infected with *Mycobacterium tuberculosis*

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*Mycobacterium tuberculosis* (Mtb) remains a significant threat to global health as it induces granuloma and systemic inflammatory responses during active tuberculosis. Mtb can induce macrophage pyroptosis, leading to the release of IL-1β and tissue damage, promoting its spread. Here, we established an in vitro Mtb-infected macrophage model to seek an effective antipyroptosis agent. Baicalin, isolated from Radix Scutellariae, was found to reduce pyroptosis in Mtb-infected macrophages. Baicalin could inhibit activation of the PERK/eIF2α pathway and thus downregulates TXNIP expression and subsequently reduces activation of the NLRP3 inflammasome, resulting in reduced pyroptosis in Mtb-infected macrophages. In conclusion, baicalin reduced pyroptosis by inhibiting the PERK/TXNIP/NLRP3 axis and might thus be a new adjuvant host-directed therapy (HDT) drug.

1. **Introduction**

As one of the top 10 causes of death worldwide and the leading cause of death caused by a single infectious agent, tuberculosis (TB) is a communicable disease caused by *Mycobacterium tuberculosis* (Mtb). Generally, Mtb can induce excessive inflammatory responses by infecting macrophages in the host, exacerbating tissue damage and patient symptoms [1]. Over the past two decades, research on TB and the development of drugs against TB have been a concern. However, owing to antimicrobial resistance and a lack of targeting drugs, these methods have not been satisfactory [2]. Therefore, a new approach to treat TB is urgently needed.

Host-directed therapy (HDT) is an emerging treatment concept that utilizes drugs to enhance protective immune responses against the pathogen or reduce exacerbated inflammatory responses, to finally balance immune reactivity and protect the host [3]. Since the excessive inflammatory response of macrophages induced by Mtb infection is the key factor in the pathological damage to a TB patient, the most important measure to alleviate the symptoms of patients is suppressing the excessive inflammatory response when the bacteria cannot be cleared in time [4]. Thus, finding drugs based on HDT might help the host immunity system “correctly” respond to Mtb infection and indirectly enhance the effectiveness of anti-TB drugs.

Baicalin is a constituent of Radix Scutellariae and exerts strong biological effects, such as anti-inflammatory [5], blood pressure-reducing [6], and glucose and lipid metabolism regulatory activities [7]. Moreover, baicalin can attenuate H$_2$O$_2$-induced cytotoxicity by reducing ER stress and activating Nrf2 signaling [8]. Our previous study demonstrated that baicalin can suppress inflammatory responses in Mtb-infected macrophages by inducing autophagy through the PI3K/Akt/NF-κB signaling pathway [9]. Here, we found a new mechanism through which baicalin
alleviates TB, in which it could inhibit pyroptosis in Mtb-infected macrophages to reduce excessive inflammatory responses.

2. Material and Methods

2.1. Chemicals and Reagents. RIPA lysis buffer, BCA protein assay kit, Protein A/G agarose/sepharose beads, and Lactate Dehydrogenase (LDH) Cytotoxicity Assay Kit were purchased from the Beyotime Institute of Biotechnology (Shanghai, China). The following antibodies were used: anti-BIP (3177S), anti-CHOP (289ST), anti-NLRP3 (15101S), IRE1a (3294T), and p-eIF2α (3398T) were purchased from Cell Signaling Technology, Inc.; anti-GSDMD (A18281) was obtained from ABclonal Technology (Wuhan, China); anti-TXNIP (sc-166234) was bought from Santa Cruz Biotechnology, Inc.; anti-β-actin monoclonal antibody was purchased from Proteintech Group (Chicago, IL); goat anti-NLRP3 (49012) was purchased from SAB Laboratories, Ltd. (Beijing, China); anti-NLRP3 (49012) was purchased from BD Biosciences (BD, Sparks, MD, USA). Middlebrook 7H9 and 7H10 media were received from Difco (Detroit, MI, USA), and oleic acid-albumin-dextrose-catalase (OADC) supplements were from BD Biosciences (BD, Sparks, MD, USA). RibofLect CP Transfection Kit was purchased from Ribo Biotechnology (Guangzhou, China). IL-1β ELISA Kit was purchased from R&D (USA).

2.2. Animals. Male C57BL/6J mice (6-8 weeks, 20 ± 3 g) were obtained from Vital River Laboratory Animal Technology Co., Ltd. (Beijing, China). All mice were acclimatized for at least 1 week before the experiments and housed in a pathogen-free facility under standard condition (4 per cage, bedding replacement once a week), with free access to food and water. Intraperitoneal injection of 2 ml of 3% sodium thioglycolate and the primary peritoneal macrophages were collected after three days. For animal sacrifice, all mice were anesthetized using isoflurane and then killed by cervical dislocation. Animal experiments strictly follow the National Institutes of Health Guide for the Care and Use of Laboratory Animals, with the approval of the Scientific Investigation Board of Shanghai University of Traditional Chinese Medicine (Shanghai, China).

2.3. Drugs. Baicalin (molecular weight: 446.37, purity ≥ 98%, the chemical structure shown in Fig. S1) is extracted from the Radix Scutellariae (the full names have been checked according to http://www.theplantlist.org). Baicalin was purchased from Shanghai Tauto Biotech Co., LTD. (CAS: 21967-41-9) (Shanghai, China). Firstly, it was dissolved with DMSO to a storage concentration of 100 mM and stored at -20°C, and then, it was diluted to 100 μM with DMEM containing 10% FBS for the following experiments.

2.4. Cell Culture. Mouse macrophage-like cell line Raw264.7 was purchased from ATCC (Manassas, VA) and cultured in DMEM supplemented with 10% fetal bovine serum (FBS) in 5% CO₂ at 37°C. Thioglycolate-elicited mouse primary peritoneal macrophages were prepared from male C57BL/6J mice (6-8 weeks of age) as described previously [10]. After 2 h, nonadherent cells were removed and the adherent cells were used as primary peritoneal macrophages.

2.5. Bacterial Strains. The Mtb H37Ra was used in this study. H37Ra strain was grown in Middlebrook 7H9 or 7H10 broth supplemented with 0.2% glycerol, 0.05% Tween-80, and 10% Middlebrook OADC supplement.

2.6. Mtb Infection. The Raw264.7 cells or primary peritoneal macrophage cells were seeded at various specifications of the cell culture plates and grown at 37°C overnight. The cells (the density of Raw264.7 cells was 3 × 10⁶, 4 × 10⁵, or 8 × 10⁵; the density of primary peritoneal macrophage cells was 6 × 10⁶, 8 × 10⁵, or 1.5 × 10⁶) were infected with Mtb H37Ra at a MOI (multiplicity of infection) of 10:1. After 4 h of coincubation at 37°C, cells were washed three times with sterile phosphate-buffered saline (PBS) and cultured with DMEM containing 10% FBS in the presence and absence of baicalin (100 μM) for different times (6, 12, and 24 hours). The experimental groups are control, Mtb 6 h, Mtb+baicalin 6 h, Mtb 12 h, Mtb+baicalin 12 h, Mtb 24 h, and Mtb+baicalin 24 h.

2.7. Western Blotting. The Raw264.7 cells or primary peritoneal macrophage cells were seeded at 6-well cell culture plates (Raw264.7 cells: 8 × 10⁵ cells/well; primary peritoneal macrophage cells: 1.5 × 10⁶ cells/well) and grown at 37°C overnight. The cells were infected with Mtb H37Ra (MOI = 10 : 1). Then, cells were washed three times with sterile PBS and cultured with DMEM containing 10% FBS in the presence and absence of baicalin (100 μM) for different times (6, 12, and 24 hours). Cells were collected and lysed in RIPA lysis buffer (Beyotime Institute of Biotechnology, China) after different times, and then, the total protein concentration of each sample was detected by a BCA protein assay kit. Then, the whole cell lysate was added 5x SDS and boiled 10 minutes in 100°C and separated by SDS-PAGE and transferred onto nitrocellulose membranes. The membranes were incubated with different primary antibodies at 4°C overnight after blocking with 5% (w/v) bovine serum albumin dissolved in TBST (1% Tween-20); the dilution ratio of each primary antibody was 1:1000. The membranes were incubated with HRP-conjugated secondary antibodies (1:10000) at room temperature for 2 hours after being washed by TBST three times. Chemiluminescence was tested with ECL-chemiluminescent kit (Thermo Scientific). The quantification of each protein was performed by ImageJ (1.51 (100), 2015).

2.8. Coimmunoprecipitation. The Raw264.7 cells or primary peritoneal macrophage cells were seeded at 30 mm dish culture plates (Raw264.7 cells: 3 × 10⁶ cells/well; primary peritoneal macrophage cells: 6 × 10⁶ cells/well) with three groups (control, Mtb 12 h, and Mtb+baicalin 12 h). The cells were infected with Mtb H37Ra (MOI = 10 : 1). Then, cells were washed three times with sterile PBS after 4 hours and cultured with DMEM containing 10% FBS in the presence...
and absence of baicalin (100 μM) for 12 hours. Raw264.7 cells or primary peritoneal macrophages were lysed by ice-cold cell lysis buffer, and cell lysates were cleared by centrifugation (12000 rpm, 10 min). Before immunoprecipitation, samples containing equal amounts of protein were pre-cleared with specific antibodies (1-2 mg/ml) overnight at 4°C with gentle rotation and then incubated with Protein A/G agarose/sepahrose beads at 4°C with gentle rotation for four hours. Then, agarose/sepahrose beads were washed extensively with lysis buffer four times, and proteins were eluted by boiling in 1x SDS buffer before SDS-PAGE electrophoresis. The steps after SDS-PAGE electrophoresis are the same as western blotting measurement.

2.9. Immunofluorescence. The Raw264.7 cells were seeded at culture plates (7 × 10^4 cells/well) with three groups (control, Mtb 12 h, and Mtb+baicalin 12 h) and grown at 37°C overnight. The cells were infected with Mtb H37Ra (MOI = 10 : 1). Then, cells were washed three times with sterile PBS after 4 hours and cultured with DMEM containing 10% FBS in the presence and absence of baicalin (100 μM) for 12 hours. The cells were washed three times with 37°C prewarmed PBS and fixed with 37°C prewarmed 4% paraformaldehyde at room temperature for 15 min, and then, the cells were washed three times with 37°C prewarmed PBS again. The cells were treated with penetrating reagents (0.2% of BAS, 2% of Triton X-100) for 15 min at room temperature and washed three times with 37°C prewarmed PBS, and then, the cells were blocked with 5% bovine serum albumin for 0.5 hours at room temperature. Rabbit anti-NLRP3 and mouse anti-TXNIP antibodies were used for immunofluorescence, primary antibodies were dissolved in 5% bovine serum albumin, and the dilution ratio of each primary antibody was 1 : 200. The cells were washed three times with 37°C prewarmed PBS after being incubated with primary antibodies at 4°C overnight. Donkey anti-mouse IgG and goat anti-rabbit IgG antibodies were used as secondary antibodies; the dilution ratio of each secondary antibody was 1 : 200. The cells were incubated with secondary antibodies for 2 hours. Then, the cells were washed three times with 37°C prewarmed PBS. The nuclei were stained with DAPI at the concentration of 2 μg/ml for 10 min. Then, the cells were washed three times with 37°C prewarmed PBS. All operations should be protected from light from the step of adding the secondary antibodies. Confocal microscopy (LSM 880, Zeiss Optics International Trading Co., LTD.) was used for examination. The quantification of each protein was performed by ImageJ (1.51 (100), 2015).

2.10. IL-1β ELISA. The Raw264.7 cells were seeded at 24-well plates (2 × 10^5 cells/well) with three groups (control, Mtb 12 h, and Mtb+baicalin 12 h) and grown at 37°C overnight. The cells were infected with Mtb H37Ra (MOI = 10 : 1). The cells were washed three times with sterilized PBS 4 hours later and then were added with or without baicalin (100 μM) medium continually cultivated for 24 hours. To measure the production of IL-1β, the supernatants of the cells were collected and measured using enzyme-linked immunosorbent assay (ELISA) kits purchased from R&D according to the manufacturer’s protocol.

2.11. LDH Release Assay. Cell death was evaluated applying the Lactate Dehydrogenase (LDH) Cytotoxicity Assay Kit. The Raw264.7 cells or primary peritoneal macrophage cells were seeded at 24-well plates (2 × 10^5 cells/well) with fifteen groups (control 6 h, Mtb 6 h, Mtb+baicalin 6 h, control 12 h, Mtb 12 h, Mtb+baicalin 12 h, control 24 h, Mtb 24 h, Mtb+baicalin 24 h, control 48 h, Mtb 48 h, Mtb+baicalin 48 h, control 72 h, Mtb 72 h, and Mtb+baicalin 72 h) and grown at 37°C overnight. The cells were infected with Mtb H37Ra (MOI = 10 : 1). The cells were washed three times with sterilized PBS 4 hours later and then were added with or without baicalin (100 μM) medium continually cultured. After baicalin (100 μM) treatment for 6, 12, 24, 48, and 72 hours, the chromogenic reagent provided in the LDH assay kit was then added, and the luminescence signals were measured using a microplate spectrophotometer. The relative amount of LDH was calculated according to the following formula: LDH relative release amount (%) = (absorbance of treated sample – absorbance of control hole of sample)/ (absorbance of maximum enzyme activity of cells – absorbance of control hole of sample) × 100.

2.12. siRNA Transfection. RAW264.7 cells were seeded at six-well tissue culture plates at a density of 3 × 10^4 cells per well with 2 ml of antibiotic-free DMEM (10% FBS) medium with six groups (control, Mtb, Mtb+siIRE1α, Mtb+siPERK, Mtb+siATF6, and Mtb+NC). The cells were incubated at 37°C in a CO2 incubator until about 80% of cells were confluent. This took approximately 24 hours. The following solutions were prepared: 1.858 ml antibiotic-free DMEM (10% FBS) medium, 120 μl buffer, 12 μl reagent, and 10 μl siRNA (20 μM). The solution was mixed gently by pipetting and incubated for 0-15 min at room temperature. The mixture was added to the 6-well plate. The cells were incubated for 48 hours. Then, the transfection efficiency was evaluated by western blotting to detect the IRE1α, p-eIF2α, ATF6, and GSDMD protein expressions. Higher interference efficiency siRNAs were selected for following researches, and Mtb was added after being transfected for 48 hours (MOI = 10 : 1). Cells were washed three times with sterilized PBS 4 hours later and then were added with or without baicalin (100 μM) medium continually cultivated for 12 hours. The cells were harvested and dissociated in ice-cold cell lysis buffer; subsequently, the proteins of whole cell were extracted and the target proteins are tested by western blotting eventually. The siRNA sequences are as follows: IRE1α-mus-963 (KD1) sense 5′-GCUAAGCCCUACUCUGUAAU TT-3′, antisense 5′-AUACAGAGUAGCGGUUAGCTT-3′; IRE1α-mus-1463 (KD2) sense 5′-CCUGAG CACCUUUTT-3′, antisense 5′-AAAGUGCCUGAUAU UGGTT-3′; IRE1α-mus-2523 (KD3) sense 5′-GGAGCU CAUUGCCUGUAATT-3′, antisense 5′-UCACGAGC AAUGACGUCCTT-3′; EIF2AK3-mus-525 (PERK-KD1) sense 5′-GCACUUAGAUAGCAUAAUUTT-3′, antisense 5′-AUUCGUCUAUCUAAAGUGCTT-5′; EIF2AK3-mus-
Figure 1: Continued.
3.1. Baicalin Ameliorates Pyroptosis in Mtb-Infected Macrophages. Pyroptosis is a form of programmed cell death characterized by upregulated expression of caspase-1-cleaved gasdermin D (GSDMD), which forms membrane pores and promotes the release of proinflammatory mediators, including IL-1β, in response to the adverse effects of certain bacteria [11]. In TB patients, Mtb can induce macrophage pyroptosis. During this progress, the nucleotide-binding domain leucine-rich repeat and pyrin domain-containing receptor 3 (NLRP3) inflammasome is activated, leading to the release of interleukin-1β (IL-1β) and HMGB1 and promoting the spread of Mtb to neighboring cells [11–13].

Our previous results suggested that baicalin can significantly inhibit NLRP3 inflammasome activation [9]. Thus, we determined whether baicalin could affect pyroptosis through the NLRP3 inflammasome. According to a previous MTT experiment, we used 100 μM baicalin to treat Mtb-infected macrophages and then investigated GSDMD expression by western blotting. As shown in Figure 1, GSDMD-N protein levels were increased in RAW264.7 cells (Figure 1(a)) and primary peritoneal macrophages (Figures 1(c) and 1(d)). Since LDH release not only occurs in pyroptosis but also occurs in other forms of cell death, the LDH release was assessed after GSDMD siRNA transfection. The results showed that LDH release was significantly reduced in Mtb-infected macrophages after GSDMD siRNA transfection, and there is no significance between the Mtb-
infected group and the treatment of baicalin (Figures 1(e)–1(g)). This result demonstrates that LDH release in Mtb-infected macrophages partly depends on the upregulation of GSDMD-N. Moreover, our previous work also revealed that baicalin could reduce the release of IL-1β [9]. In addition, the level of HMGB1 in the supernatants was assessed. The results showed that baicalin treatment could reduce the expression of HMGB1 in 24 hours (Figure S2).

Propidium iodide (PI), whose uptake indicates cell damage or death, was widely used to assess the viability of cells in different experimental models. To analyze whether baicalin affects the viability of macrophages, we determined the viability of macrophages in different groups by PI staining flow cytometric assay. The results of PI staining flow cytometric assay demonstrated that Mtb infection can increase the death or damage rates of macrophages at different time points both in RAW264.7 cells and in primary peritoneal macrophages, while treatment with baicalin can significantly abolish this effect within 24 h (Figure S3). These results indicated that baicalin can reduce pyroptosis by inhibiting NLRP3 inflammasome activation.

3.2. Baicalin Reduces TXNIP-NLRP3 Interactions in Mtb-Infected Macrophages. Generally, the NLRP3 inflammasome can be activated by various pathogen-associated molecular patterns (PAMPs) and damage-associated molecular patterns (DAMPs) [14, 15]. Thioredoxin-interacting protein (TXNIP) is a member of the thioredoxin (TRX) system and an endogenous regulator of redox/glucose-induced

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**Figure 2:** Baicalin reduces TXNIP-NLRP3 interactions in Mtb-infected macrophages. (a, c) Levels of TXNIP protein detected by western blotting analysis in RAW264.7 cells. (b, d) Levels of TXNIP protein determined by western blotting in primary peritoneal macrophage cells. (e, f) TXNIP or NLRP3 immunoprecipitated from RAW264.7 cells and primary peritoneal macrophage cells were immunoblotted for TXNIP or NLRP3, respectively, and rebotted for TXNIP or NLRP3, respectively. Experiment performed three times. *p < 0.05; **p < 0.01; ***p < 0.001; NS: nonsignificant.
stress and inflammation; it is also essential for maintaining the cellular redox status balance and is involved in various disorders including sepsis, traumatic brain injury, diabetes, and Alzheimer’s disease (AD) [16–19]. TXNIP upregulates inflammatory responses via decreased expression of TRX, and TRX suppression leads to increased oxidative stress [20]. Inhibiting the dissociation of TRX and TXNIP and blocking the interactions between TXNIP and NLRP3 markedly suppresses NLRP3 inflammasome activation, IL-1β maturation, and pyroptosis [21, 22]. As a secondary signal, TXNIP is also a key factor that activates the NLRP3 inflammasome [23]. To examine whether baikalin affects TXNIP protein expression, we measured its expression by western blotting. As shown in Figure 2, Mtb infection increased TXNIP protein levels, whereas baikalin reduced the expression in RAW264.7 cells and primary peritoneal macrophages (Figures 2(a)–2(d)).

TXNIP binds to NLRP3 through the leucine-rich repeat domain after it dissociates from TRX, and then, the dissociative TXNIP activates the NLRP3 inflammasome in response to oxidative stress via an association with NLRP3 [24]. To assess whether NLRP3 inflammasome activation is due to increased TXNIP expression, coimmunoprecipitation and immunofluorescence were used to analyze the interactions

Figure 3: Baicalin prevents the colocalization of TXNIP and NLRP3 in Mtb-infected macrophages. (a, b) Confocal microscopy of RAW264.7 cells with different treatments immunostained with anti-TXNIP (red), anti-NLRP3 (green), and DAPI (blue). Mean ± SD of colocalization of TXNIP and NLRP3, TXNIP and NLRP3 florescence intensity in three independent pictures was shown as scatter plot and error bar (n = 3). Kruskal-Wallis H rank and inspection was used as the statistical analysis method. According to the statistical results of Kruskal-Wallis H rank sum test, p < 0.05. The difference between the three groups was statistically significant.
Figure 4: Continued.
between TXNIP and the NLRP3 inflammasome. These interactions were reduced by baicalin treatment in RAW264.7 cells (Figure 2(e)) and primary peritoneal macrophages (Figure 2(f)). Confocal imaging also showed that Mtb increased the colocalization of TXNIP and NLRP3, and baicalin treatment significantly reduced TXNIP and NLRP3 proteins levels and reduced their colocalization (Figures 3(a) and 3(b)). These results showed that baicalin can significantly reduce the interactions between TXNIP and NLRP3.

3.3. Baicalin Reduces TXNIP Expression by Inhibiting ER Stress in Mtb-Infected Macrophages. ER stress plays a key role in inducing NLRP3 inflammasome activation by promoting interactions between TXNIP and NLRP3, thereby increasing the production of various proinflammatory factors and inducing pyroptosis [23, 25, 26]. Recent studies also demonstrated that Mtb infection can induce ER stress [27]. Therefore, we determined whether baicalin could affect TXNIP expression through ER stress in the Mtb-infected model.

Increased BIP and CHOP protein levels are reliable biomarkers of ER stress. Here, we found that ER stress was strongly induced after Mtb infection, whereas baicalin treatment significantly reduced these effects at different time points (Figures 4(a)–4(d)). Moreover, activation of three transmembrane ER-resident stress sensors (IRE1α, PERK, and ATF6) is involved in the induction of ER stress. IRE1α, PERK, and ATF6 remain inactive after binding to BIP under unstressed conditions. When misfolded proteins accumulate, BIP dissociates from these three ER stress sensors, resulting in the activation of each branch. Previous studies showed that the increase in TXNIP levels depends on the activation of PERK and IRE1α [22]. Here, we found that the increase of TXNIP did not depend on the activation of ATF6 by measuring TXNIP protein expression after IRE1α, PERK, and ATF6 siRNA transfection (Figures 4(e)–4(g)). These results indicated that baicalin decreased TXNIP protein levels, which depended on the inhibition of IRE1α and PERK, rather than the ATF6 pathway. Western blotting was used to assess whether Mtb infection would upregulate IRE1α and p-eIF2α expression. The results showed that Mtb infection increased PERK/p-eIF2α and IRE1α expression, whereas baicalin treatment markedly decreased PERK/p-eIF2α expression at different time points. However, the increase in IRE1α protein levels was not affected by the baicalin treatment (Figures 5(a)–5(d)). These results showed that baicalin reduces TXNIP expression by inhibiting the PERK/eIF2α pathway in Mtb-infected macrophages.

3.4. Baicalin Reduces Pyroptosis in Mtb-Infected Macrophages by Inhibiting PERK/eIF2α Activation. A previous study showed that TXNIP expression depends on the activation of the IRE1α and PERK pathways, rather than the ATF6 pathway in Mtb-infected macrophages. We determined whether baicalin could inhibit pyroptosis through the IRE1α and PERK pathways. Here, we measured the respective effect on pyroptosis by IRE1α siRNA and PERK siRNA and assessed GSDMD-N and IL-1β levels by western blotting and ELISA. As shown in Figure 6, GSDMD protein expressions in Mtb-infected macrophages did not differ between the IRE1α siRNA and non-IRE1α siRNA groups (Figures 6(a) and 6(c)). A similar phenomenon was also observed regarding IL-1β (Figure 6(e)). These results indicated that pyroptosis is independent on the IRE1α pathway in Mtb-infected macrophages.

Similarly, GSDMD protein and IL-1β levels were measured after PERK siRNA transfection. GSDMD expression was markedly reduced in Mtb-infected macrophages after PERK siRNA transfection (Figures 6(b) and 6(d)). Similar results were observed for IL-1β (Figure 6(f)). Previously, we have revealed that baicalin could suppress the activation of the NLRP3 inflammasome by inhibiting the interaction between TXNIP and the NLRP3 inflammasome. Moreover,
Figure 5: Baicalin inhibits endoplasmic reticulum stress in Mtb-infected macrophages. (a, c) Western blotting analysis of IRE1α, p-IRE1α, and p-eIF2α expression in RAW264.7 cells. (b, d) Levels of IRE1α, p-IRE1α, and p-eIF2α protein assessed by western blotting in primary peritoneal macrophage cells. Data are shown as mean ± SD of at least three independent experiments. **p < 0.01; ***p < 0.001; NS: nonsignificant.
Figure 6: Continued.
In the past years, some anti-TB drugs have achieved certain effects. However, due to wide use of these drugs, the drug-resistant Mtb has emerged. Thus, new effective TB drugs are needed [28]. But the research and development of new TB drugs are very difficult and slow, and thus, there is an urgent need to explore a new method of treating TB.

During active TB, Mtb often induces granuloma and systemic inflammatory responses. Most previous studies predominantly focused on killing Mtb, but the excessive inflammatory response induced by Mtb was ignored. In fact, Mtb could cause excessive inflammatory response to damage tissue and promote the spread of itself. Therefore, reducing excessive inflammation in the host could slow the spread of Mtb and help anti-Mtb drugs work.

HDT is an effective new treatment concept for multidrug-resistant Mtb, and it is aimed at enhancing protective immune responses or reducing exacerbated inflammation, finally to balance host immunity [29]. Based on HDT, we found a new drug for TB—baicalin. It is an active ingredient extracted from Radix Scutellariae and exerts various effects, including anti-inflammatory and antihyperglycemic activities [5–7]. Our previous study showed that baicalin could suppress NLRP3 inflammasome activation by facilitating the degradative mechanism of autophagy [9]. Here, we revealed that baicalin could significantly reduce pyroptosis in Mtb-infected macrophages by inhibiting NLRP3 inflammasome activation.

Activation of the NLRP3 inflammasome requires two signals, a priming signal and activation signal. The priming signal is induced by microbial components or endogenous cytokines and results in the activation of NF-κB and upregulation of NLRP3, pro-caspase-1, and pro-IL-1β. The activation signal is elicited by a series of stimuli including numerous molecular or cellular processes. For example, TXNIP can activate the NLRP3 inflammasome [14, 23]. Previously, we found that baicalin can inhibit the NF-κB pathway, thereby contributing to the inhibitory effect on NLRP3 inflammasome activation [9]. Consistent with those results, here, we revealed that baicalin also inhibits activation of the PERK/elf2α pathway and thus downregulates TXNIP expression and subsequently reduces activation of the NLRP3 inflammasome, resulting in reduced pyroptosis in Mtb-infected macrophages (Figure 7).

Furthermore, recent studies have demonstrated that suppressing the dissociation of TRX and TXNIP and reducing the interactions between TXNIP and NLRP3 significantly inhibited NLRP3 inflammasome activation, IL-1β maturation, and pyroptosis [21, 22]. Here, we also revealed that reducing the interactions between TXNIP and NLRP3 with baicalin can contribute to inhibiting the activation of the NLRP3 inflammasome in Mtb-infected macrophages (as...
shown in Figures 2 and 3). Previous studies demonstrated that baicalin attenuated ER stress induced by tunicamycin through the CHOP/eNOS/NO pathway in cardiomyocytes and prevented lipotoxicity in AML-12 cells by reducing the interactions between TXNIP and the NLRP3 in flamasome [30–33]; here, we also showed this effect.

Moreover, inhibiting the dissociation of TRX and TXNIP and blocking the interactions between TXNIP and NLRP3 markedly suppress NLRP3 inflammasome activation, IL-1β maturation, and pyroptosis [21, 22]. The effect of baicalin on inhibiting the interaction between TXNIP and NLRP3 might be due to suppressing the dissociation of TXNIP and TRX, but the specific mechanism through which baicalin inhibits TXNIP and NLRP3 needs to be further explored. Moreover, baicalin can alleviate pyroptosis in hepatocytes by inhibiting NLRP3-GSDMD signaling in vitro and reduces age-related macular degeneration through NLRP3-regulated pyroptosis [33, 34]. However, the current study showed that baicalin ameliorated pyroptosis in Mtb-infected macrophages by inhibiting the interactions between TXNIP and NLRP3.

ER stress plays a key role in human diseases, including colitis and acute pancreatitis, by increasing the release of proinflammatory factors such as IL-1β, cleaved caspase-1, and TNF-α which is elicited by TXNIP during neuroinflammation. Other studies demonstrated that ER stress can result in NLRP3 inflammasome activation, and Mtb stimulates consistent IL-1β production and pyroptosis. When ER stress occurs, three transmembrane proteins (protein kinase-like ER kinase (PERK), activating transcription factor 6 (ATF6), and inositol-requiring enzyme 1α (IRE1α)) are separated from the chaperone binding immunoglobulin protein (BIP) in response to unfolded or misfolded proteins [34]. Once activated, PERK phosphorylates eukaryotic initiation factor 2α (eIF-2α), which would attenuate translational initiation and block new proteins in the ER [35]. Severe ER stress induces the activation of the CCAAT-enhancer-binding protein homologous protein (CHOP) [36]. Furthermore, upregulation of IRE1α can elicit dissociation of TRX and TXNIP, after which TXNIP binds to the NLRP3 inflammasome and activates it [37, 38]. Moreover, ER stress can increase TXNIP expression through activation of the PERK or IRE1α.

![Figure 7: Schematic summary of baicalin inhibiting pyroptosis in Mtb-infected macrophages.](image)
signaling pathway [39–41]. Consistent with these results, our study showed that baicalin suppressed the expression of PERK/p-eIF2α. Moreover, we also found that the upregulation of TXNIP was achieved by activation of the IRE1α and PERK pathways. However, p-eIF2α protein expression was reduced by baicalin at different time points, whereas the expression of IRE1α and p-IRE1α in Mtb-infected macrophages was not affected. Our results also showed that GSDMD-N protein expression decreased after PERK siRNA transfection, but there was no change after IRE1α siRNA transfection (Figure 5). This indicated that baicalin reduced pyroptosis by suppressing the activation of the PERK/p-eIF2α pathway. The results of IL-1β were similar to those of GSDMD-N. Thus, we proposed that baicalin reduced pyroptosis by inhibiting the PERK/TXNIP/NLRP3 axis. To prove that the effect of baicalin on pyroptosis is extensive and associated with stabilization, we chose two cell lines, RAW264.7 cells and primary peritoneal macrophage cells. These two cell lines are both macrophage cells; RAW264.7 is the most commonly used cell line to study the role of macrophages in bacterial infections and inflammatory responses, whereas primary peritoneal macrophage cells are more sensitive to the stimulation by bacterial infection. All of our results were confirmed in both cell lines.

In conclusion, baicalin markedly reduced pyroptosis in Mtb-infected macrophages, which suggested that it is a novel candidate adjuvant drug for HDT to anti-TB. Additionally, our study generated new research insights regarding TB that could help to identify more effective and specific HDT targets. Drugs such as baicalin can alter host cellular defense mechanisms and prevent host tissue damage during Mtb infection. Taken together, baicalin showed promising effects to prevent pyroptosis in Mtb-infected macrophages through the inhibition of the PERK/TXNIP/NLRP3 axis.

In this study, we used Mtb strain H37Ra as a source of infection. It is an attenuated strain isolated from the standard virulent strain H37Rv, to which it is very similar genetically. Because H37Ra is less virulent, more simple to use, and safer, we used RAW264.7 cells and primary peritoneal macrophages infected with H37Ra as an infection model system [42]. Further research would be necessary to examine the effect of baicalin in combination with anti-TB drugs on animal models of H37Rv infection.

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

Conflicts of Interest
The authors declare no conflict of interest.

Authors’ Contributions
XJ and YZ conceived and designed the experiments. YF, JS, YL, FL, and BN performed the experiments. YF and JS analyzed the data. YF and XJ wrote the paper. All authors have read and approved the final manuscript.

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Supplementary Materials
Figure S1: the chemical structure of baicalin. Figure S2: baicalin evidently inhibits the Mtb-induced pyroptosis. Figure S3: baicalin attenuates pyroptosis in Mtb-infected macrophages. (Supplementary Materials)

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