Punicalagin Attenuates LPS-Induced Inflammation and ROS Production in Microglia by Inhibiting the MAPK/NF-κB Signaling Pathway and NLRP3 Inflammasome Activation

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Purpose: Neurodegenerative diseases are associated with neuroinflammation along with activation of microglia and oxidative stress, but currently lack effective treatments. Punicalagin is a natural bio-sourced product that exhibits anti-inflammatory effects on several chronic diseases; however, the anti-inflammatory and anti-oxidative effects on microglia have not been well examined. This study aimed to investigate the effects of punicalagin on LPS-induced inflammatory responses, NLRP3 inflammasome activation, and the production of ROS using murine microglia BV2 cells.

Methods: BV2 cells were pre-treated with punicalagin following LPS treatment to induce inflammation. The secretion of NO and PGE2 was analyzed by Griess reagent and ELISA respectively, while the expressions of iNOS, COX-2, STAT3, ERK, JNK, and p38 were analyzed using Western blotting, the production of IL-6 was measured by ELISA, and the activity of NF-κB was detected using promoter reporter assay. To examine whether punicalagin affects NLRP3 inflammasome activation, BV2 cells were stimulated with LPS and then treated with ATP or nigericin. The secretion of IL-1β was measured by ELISA. The expressions of NLRP3 inflammasome-related proteins and phospho IκBα/IκBα were analyzed using Western blotting. The production of intracellular and mitochondrial ROS was analyzed by flow cytometry.

Results: Our results showed that punicalagin attenuated inflammation with reduction of pro-inflammatory mediators and cytokines including iNOS, COX-2, IL-1β, and reduction of IL-6 led to inhibition of STAT3 phosphorylation by LPS-induced BV2 cells. Punicalagin also suppressed the ERK, JNK, and p38 phosphorylation, attenuated NF-κB activity, inhibited the activation of the NLRP3 inflammasome, and reduced the production of intracellular and mitochondrial ROS by LPS-induced BV2 cells.

Conclusion: Our results demonstrated that punicalagin attenuated LPS-induced inflammation through suppressing the expression of iNOS and COX-2, inhibited the activation of MAPK/NF-κB signaling pathway and NLRP3 inflammasome, and reduced the production of ROS in microglia, suggesting that punicalagin might have the potential in treating neurodegenerative diseases.

Keywords: punicalagin, microglia, NLRP3 inflammasome, MAPK, NF-κB
**Introduction**

Neurodegenerative disease, also known as the silent epidemic, is characterized by a progressive loss of neuronal cells in the brain. As a consequence of increase in the aging population, the range of medical resources used to treat and prevent senile-related diseases are also increasing year by year. Alzheimer’s disease is the most common type of neurodegenerative disease, and at present, there are ample studies to demonstrate that Alzheimer’s disease is characterized by the deposition of extracellular amyloid-beta and abnormal tau phosphorylation expression around the affected area of the brain; additionally, these cause dementia, progressive memory loss and the impairment of cognitive function. Oxidative stress and neuroinflammation are important factors in the pathogenesis of neurodegenerative diseases. Chronic oxidative stress is associated with neuroinflammation and neurodegeneration by activating signaling pathways of proinflammatory activities. As the major resident macrophage-like immune cells in the central nervous system, microglia have a pivotal role in neuroinflammation and furnish multiple beneficial functions to neuron cells, including maintenance of cellular homeostasis and innate immunity.

Neuroinflammation is a crucial defense mechanism that competes with infection or trauma in the central nervous system. During neuroinflammation, microglia recognize pathogen-associated molecular patterns (PAMPs) (eg lipopolysaccharide (LPS) through Toll-like receptor-4 (TLR-4) and induce the robust activation of innate immune responses leading to an increase in the production of inflammatory mediators and cytokines such as nitric oxide (NO), prostaglandin E₂ (PGE₂) and cyclooxygenase (COX)-2, interleukin (IL)-6, tumor necrosis factor (TNF)-α and IL-1β as well as promoting the generation of reactive oxygen species (ROS) through nicotinamide-adenine dinucleotide phosphate oxidase. Additionally, microglial cells are stimulated by damage-associated molecular patterns (DAMPs) like adenosine triphosphate (ATP) and bind to P2X7 receptors then further induce the assembly of the nucleotide-binding oligomerization domain (NOD)-like receptor family.
pyrin domain containing 3 (NLRP3) inflammasome.\textsuperscript{3} NLRP3 inflammasome is a cytosolic protein complex composed of NLRP3, an apoptotic speck-containing protein with a CARD (ASC) and caspase-1 that is formed in response to both PAMPs and DAMPs stimuli consequently leading to cleavage of pro-caspase-1 while promoting maturation of IL-1β and inducing pyroptosis.\textsuperscript{8} Inhibition of NLRP3 inflammasome activation has been considered as a therapeutic strategy for ameliorating the progression of Alzheimer’s disease.\textsuperscript{9–11}

Polyphenols are found in various kinds of food sources such as tea, cocoa, fruits, berries, and vegetables, which have been demonstrated to exhibit antioxidative and anti-inflammatory properties\textsuperscript{12,13} and represent a protective role in many chronic diseases including cardiovascular disease, neurodegenerative diseases and diabetes.\textsuperscript{14} Punicalagin is a large polyphenol compound with a molecular mass of 1084.7 and is the main compound of pomegranate (\textit{Punica granatum}).\textsuperscript{15} Although previous studies have demonstrated that punicalagin has benefits in inflammation-associated chronic diseases,\textsuperscript{16–18} the antioxidative and anti-inflammatory effects of punicalagin on microglia have not been well examined; consequently, we aimed to investigate the anti-oxidative and anti-inflammatory effects of punicalagin on microglia while concurrently examining these effects on the activation of NLRP3 inflammasome in LPS-induced BV2 cells.

\textbf{Materials and Methods}

\textbf{Materials and Reagents}

Punicalagin (purity $\geq$ 98\%) was purchased from ChemFaces (catalog number: CFN99938, Wuhan, Hunan, China). Lipopolysaccharide (LPS from \textit{E. coli}, O111:B4, catalog number: L3024), Griess reagent, and protease inhibitor cocktails were obtained from Sigma Aldrich (St. Louis, MO, USA). MTT reagent (3-(4, 5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) was supported by MDBio, Inc. (catalog number: 101-298-93-1, Taipei, Taiwan, R.O.C). IL-1β (catalog number: 88-7013-86) and IL-6 (catalog number: 88-7064-86) enzyme-linked immunosorbent assay (ELISA) kits, reactive oxygen species (ROS) detection reagent (catalog number: D399), and mitosox red mitochondrial superoxide indicator (catalog number: M36008) were purchased from Invitrogen (Carlsbad, CA, USA). PGE\textsubscript{2} ELISA kit (catalog number: 514010), nigericin (catalog number: 11437), and ATP (catalog number: 14498) were purchased from Cayman Chemical (Ann Arbor, MI, USA). Phosphatase inhibitor cocktails and Radioimmunoprecipitation assay (RIPA) buffer (catalog number: RIPA-50) were obtained from FIVEphoton Biochemicals (San Diego, CA, USA). BCA protein assay kit (catalog number: 2325) was purchased from Thermo Scientific (Waltham, MA, USA). Zeocin (catalog number: ant-zn-lp) was purchased from Invivogen (San Diego, CA, USA). Antibodies for phospho-signal transducer and activator of transcription 3 (STAT3) (catalog number: 9145), STAT3 (catalog number: 9139), phospho-c-Jun N-terminal kinase (JNK) (catalog number: 9255), JNK (catalog number: 9258), phospho-p38 MAPK (catalog number: 4511), p38 MAPK (catalog number: 8690), phospho-extracellular signal-regulated kinase (ERK)1/2 (catalog number: 4370), ERK1/2 (catalog number: 4695), ASC (catalog number: 67824), cleaved-IL-1β (catalog number: 52718), IL-1β (catalog number: 12242), cleaved-caspase-1 (catalog number: 67314), caspase-1 (catalog number: 24232), phospho-IκBα (catalog number: 2859), and IκBα (catalog number: 4814) were purchased from Cell Signaling Technology (Danvers, MA, USA). COX-2 (catalog number: sc-514489) and iNOS (catalog number: sc-7271) antibodies were obtained from Santa Cruz Biotechnology (Dallas, TX, USA). NLRP3 (catalog number: GTX00763) and β-actin (catalog number: TA328070) antibodies were purchased from GeneTex (Irvine, CA, USA) and OriGene Technologies (Rockville, MD, USA) respectively.

\textbf{Cell Culture}

Mouse microglial BV2 cell line was obtained from the Food Industry Research and Development Institute (Hsinchu, Taiwan, R.O.C). Cells were cultured in a humidified atmosphere at 37°C under 5\% CO\textsubscript{2} in high-glucose Dulbecco’s modified Eagle’s medium supplemented with 10\% fetal bovine serum and 100 U/mL penicillin and 100 U/mL streptomycin (Gibco, Waltham, MA, USA).

\textbf{MTT Assay for Cell Viability}

Cell viability of BV2 cells was measured by MTT assay. Briefly, BV2 cells ($1 \times 10^5$ cells/well) were cultured in 96-well culture plates and allowed to attach overnight. Afterward, cells were pre-treated with different concentrations of
punicalagin for 30 min following 1 μg/mL LPS treatment for 24 h. Then, 10 μL of 5 mg/mL MTT solution was added to each well and further incubated at 37°C for 4 h. Subsequently, 100 μL acidic isopropanol/HCl (isopropanol with 0.04 N HCl) was added to dissolve the formazan crystals. The absorbance was determined by spectrophotometry at 570 nm, and the percentage of viable cells was normalized to the untreated control.

### Nitric Oxide Assay
The concentration of nitric oxide in the medium was measured by Griess reagent. Briefly, BV2 cells (1 × 10^5 cells/well) were cultured in 96-well culture plates and allowed to attach overnight. Afterward, cells were pre-treated with various concentrations of punicalagin for 30 min and then treated with LPS (1 μg/mL) for 24 h. Cell culture supernatant was collected and the concentration of nitric oxide was assessed by Griess reagent. The absorbance was determined at 540 nm by spectrophotometrically and the standard curve of NaNO_2 was used to calculate the nitric oxide concentration.

### Western Blot Analysis
Cells were washed with PBS and lysed by RIPA buffer supplemented with protease inhibitor cocktails and phosphatase inhibitor cocktails. The protein concentration was determined by the BCA protein assay kit. Equal amounts of protein were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis and transferred to polyvinylidene fluoride membranes. After blocking with 5% non-fat milk or BSA in TBST for 1 h, the membranes were incubated overnight with primary antibodies at 4°C overnight. Afterward, blots were washed three times with TBST, blocked with anti-rabbit or anti-mouse horseradish peroxidase-conjugated immunoglobulin G secondary antibodies diluted in TBST (1:5000) for 1 h at room temperature, and then the blots were washed three times using TBST. The intensities of protein bands were determined by Molecular Imager® Gel Doc™ XR System (Bio-Rad Laboratories, Hercules, California, USA) with Image Lab™ Software using an ECL chemiluminescence substrate (Thermo Scientific, Waltham, MA, USA). The result of β-actin was used to normalize the quantity of the protein bands.

### Enzyme-Linked Immunosorbent Assay (ELISA)
The levels of IL-6, IL-1β, and PGE_2 in the medium were measured by ELISA. Briefly, BV2 cells (1 × 10^5 cells/well) were cultured in 96-well culture plates and allowed to attach overnight. Cells were pre-treated with different concentrations of punicalagin for 30 min following 1 μg/mL LPS treatment for 24 h or 48 h. Cell culture supernatant was collected for analysis according to the manufacturer’s protocol.

### NF-κB Promoter Reporter Assay
BV2-Blue cells were derived from BV2 cells that were stably expressing a secreted embryonic alkaline phosphatase (SEAP) gene inducible by NF-κB as described in the previous study and maintained in Dulbecco’s modified Eagle’s medium supplemented with Zeocin (15 μg/mL) (InvivoGen, San Diego, CA, USA). Cells (1 × 10^5 cells/well) were seeded in a 96-well plate and allowed to attach overnight. Afterward, cells were pre-treated with different doses of punicalagin for 30 min and then treated with LPS (1 μg/mL) for 24 h. The medium then was harvested and mixed with QUANTI-Blue medium (100 μL cell culture supernatant to 100 μL QUANTI-Blue medium) (InvivoGen, San Diego, CA, USA) in 96-well plates and incubated at 37 °C for 45 min. SEAP activity was assessed by measuring the optical density at 655 nm using a microplate reader.

### Flow Cytometry
The productions of intracellular ROS and mitochondrial ROS were measured by flow cytometry. BV2 cells (5 × 10^5 cells/well) were cultured in 6-well plates and allowed to attach overnight. Afterward, cells were pre-treated with different concentrations of punicalagin for 30 min following 1 μg/mL LPS treatment for 24 h. For the detection of intracellular ROS, cells were stained by the fluorescent probe dichloro-dihydro-fluorescein diacetate (DCFH-DA) reagent (Invitrogen, Carlsbad, California, USA) with 10 μM in HBSS buffer. For the detection of mitochondrial ROS, cells were stained by the MitoSOX™ Red reagent with 5 μM in HBSS buffer. After staining for 40 or 45 min, cells were washed with PBS and
then analyzed using flow cytometry (Beckman Coulter Cytomics FC500 MCL Flow Cytometer System with CXP cytometer software, Elkin, NC, USA).

**Immunofluorescence Staining**
BV2 cells were seeded on the coverslips at a density of $2 \times 10^5$ cells/well, placed in 6-well plates and allowed to attach overnight. The cells were pre-treated with different doses of punicalagin for 30 min following 1 μg/mL LPS treatment for 24 h, then stained with DCFH-DA or MitoSOX™ Red following the above experimental conditions. Afterward, cells were washed with PBS and then stained with DAPI. Images were taken by fluorescence microscope (Nikon, Tokyo, Japan).

**Statistical Analysis**
The experimental results are presented as mean ± standard deviation (SD). All data are shown from three independent experiments. Statistical analysis was performed by one-way ANOVA followed by Tukey post-hoc test using GraphPad Prism 6 (San Diego, CA, USA). The significant difference between the groups was defined as $^*p < 0.05$; $^{**}p < 0.01$; $^{***}p < 0.001$.

**Results**

**Punicalagin Reduces the Production of NO and PGE$_2$ and Inhibits the Expression of COX-2 and NOS2 in LPS-Induced BV2 Cells**
To determine the toxic effect of punicalagin, the effect of punicalagin on the cell viability of BV2 cells was examined. Cells were pre-treated with various doses (0, 25, 50, 75, 100 μM) of punicalagin for 30 min following 1 μg/mL LPS treatment for 24 h. The results showed that the LPS-alone group slightly reduced cell viability, whereas punicalagin had no cytotoxic effect on BV2 cells when cells were treated with 0 to 100 μM punicalagin (Figure 1A). In addition, at higher doses of punicalagin groups (75 and 100 μM), punicalagin rescued LPS-induced cell death compared with the LPS-alone group (Figure 1A). To examine whether punicalagin affected LPS-induced NO and PGE$_2$ productions, BV2 cells were pre-treated with various doses (0 to 100 μM) of punicalagin for 30 min and then treated with LPS (1 μg/mL) for 24 h. The levels of NO and PGE$_2$ in cell culture supernatants were examined by Griess reagent and ELISA respectively.

As shown in Figure 1B, punicalagin significantly attenuated NO production by LPS-induced BV2 cells in a concentration-dependent manner. In addition, a high dosage (100 μM) of punicalagin significantly decreased the secretion of PGE$_2$ by LPS-induced BV2 cells (Figure 1C). Inducible nitric oxide synthase (iNOS) is a key enzyme generating nitric oxide (NO) from the amino acid L-arginine,
while cyclooxygenase-2 (COX-2) is a key enzyme converting arachidonic acid into PGE$_2$. We further examined whether punicalagin affected the expression of iNOS and COX-2. BV2 cells were pre-treated with various doses of punicalagin (0 to 100 μM) for 30 min following 1 μg/mL LPS treatment for 24 h. The expression of iNOS and COX-2 was detected by Western blot. As shown in Figure 1D–F, punicalagin significantly suppressed both iNOS and COX-2 expressions in a dose-dependent manner as compared with LPS alone.

**Punicalagin Reduces the Secretion of IL-6 and Inhibits the Phosphorylation of STAT3 by LPS-Induced BV2 Cells**
The pro-inflammatory cytokine IL-6 is demonstrated as being involved in the etiopathology of Alzheimer’s disease. To examine whether punicalagin affects the secretion of IL-6 by LPS-induced BV2 cells, cells were pre-treated with various doses of punicalagin (0 to 100 μM) for 30 min following 1 μg/mL LPS treatment for 24 h. The level of IL-6 in the cell culture medium was detected by ELISA. As shown in Figure 2A, punicalagin significantly decreased the secretion of IL-6 by LPS-induced BV2 cells in a dose-dependent manner. IL-6 is known to regulate the phosphorylation of the signal transducer and activator of transcription 3 (STAT3) during LPS/TLR4-driven inflammation. To further examine whether punicalagin affects STAT3 activation by LPS-induced BV2 cells, cells were pre-treated with various doses of punicalagin (0 to 100 μM) for 30 min following 1 μg/mL LPS treatment for 2 h. The expression of phospho-STAT3 and...
STAT3 was determined by Western blot. As shown in Figures 2B and C, the experimental results showed that punicalagin significantly suppressed the phosphorylation of STAT3 in a concentration-dependent manner.

Punicalagin Attenuates the Activation of Both MAPK and NF-κB Signaling Pathways in LPS-Induced BV2 Cells

Both MAPK and NF-κB signaling pathways are known to drive inflammation-associated gene expressions during LPS-induced inflammation. To assess whether punicalagin regulated the activation of MAPK and NF-κB signaling pathways by LPS-induced BV2 cells, cells were pre-treated with various doses of punicalagin (0 to 100 μM) for 30 min following 1 μg/mL LPS stimulation for 24 h.

Figure 1 The effect of punicalagin on the production of pro-inflammatory mediators (NO and PGE2) by LPS-induced BV2 cells. Cells were pre-treated with various concentrations (0, 25, 50, 75, 100 μM) of punicalagin for 30 min, and then treated with LPS (1 μg/mL) for 24 h. (A) The cell viability was determined by MTT assay. (B and C) The secretion of NO and PGE2 was measured by Griess reagent assay and ELISA respectively. Expressions of iNOS and COX-2 were analyzed using Western blot. The representative images are shown in (D) and the quantitative results of three independent experiments shown in (E and F). β-actin was used as a loading control. Statistical significance was indicated as *p < 0.05; **p < 0.01; ***p <0.001.

Figure 2 The effect of punicalagin on the production of IL-6 and the activation of STAT3 in LPS-induced BV2 cells. Cells were pre-treated with punicalagin (0, 25, 50, 75, 100 μM) for 30 mins and then treated with LPS (1 μg/mL) for 24 h. (A) The level of IL-6 was measured by ELISA. The data are presented as the means ± SD of three independent experiments. Statistical significance was assessed by one-way ANOVA represented as follows: ***p < 0.001 vs LPS alone. Cells were pre-treated with punicalagin (0, 25, 50, 75, 100 μM) for 30 min and then treated with LPS (1 μg/mL) for 2 h. The expression of phospho-STAT3 and STAT3 was determined by Western blot. The representative images are shown in (B) and the quantitative results of three independent experiments shown in (C). β-actin was used as a loading control. Statistical significance was indicated as ***p <0.001.

STAT3 was determined by Western blot. As shown in Figures 2B and C, the experimental results showed that punicalagin significantly suppressed the phosphorylation of STAT3 in a concentration-dependent manner.
mL LPS treatment for 6 h. The expression of phospho-ERK, ERK, phospho-JNK, JNK, phospho-p38, p38 was determined by Western blot. The representative images are shown in (A) and the quantitative results of three independent experiments shown in (B–D). β-actin was used as a loading control. BV2-Blue cells were pre-treated with punicalagin (0, 25, 50, 75, 100 μM) for 30 min following 1 μg/mL LPS treatment for 24 h. The activation of NF-κB was measured by detected SEAP activity. BV2 cells were pretreated with punicalagin (0, 25, 50, 100 μM) for 30 min following with 1 μg/mL LPS treatment for 30 min. The expressions of phospho-IκBα and IκBα by LPS-activated BV2 cells were determined by Western blot. The representative images are shown in (F) and the quantitative results of three independent experiments shown in (G). β-actin was used as a loading control. Statistical significance was indicated as *p < 0.05, **p < 0.01 and ***p < 0.001 vs LPS alone.

**Punicalagin Attenuates the Secretion of IL-1β and Inhibits the Cleavage of IL-1β and Caspase-1 in LPS/ATP-Induced and LPS/Nigericin-Induced BV2 Cells**

The activation of NLRP3 inflammasome has been demonstrated to associate with neuroinflammatory diseases including Alzheimer’s disease, Parkinson’s disease and amyotrophic lateral sclerosis.26,27 To examine whether punicalagin affected NLRP3 inflammasome activation, BV2 cells were pre-treated with various doses of punicalagin for 30 min, primed with LPS for 47.5 h, and then stimulated with ATP or nigericin for 30 min. The secretion of IL-1β was analyzed by ELISA and the expression of inflammasome-associated proteins was examined using Western blot. As shown in Figures 4A and B, the experimental results revealed that punicalagin significantly reduced the secretion of IL-1β by both LPS/ATP-induced and LPS/nigericin-induced BV2 cells in a dose-dependent manner. Furthermore, our results also showed that punicalagin significantly decreased the expressions of cleave-IL-1β and cleave-caspase-1 by both LPS/ATP-induced and LPS/nigericin-induced BV2 cells (Figures 4C–H).
Punicalagin Attenuates the Production of Both Intracellular and Mitochondrial ROS in LPS-Induced BV2 Cells

ROS plays a vital role in immunity by enhancing immunological defense and causing oxidative damage. Next, we further investigated the anti-oxidation potential of punicalagin by determining the levels of reactive oxygen species (ROS) in intracellular and mitochondrial using DCFH-DA and MitoSOX red staining respectively. The results were examined by flow cytometry and fluorescent microscopy. As shown in Figure 5, punicalagin significantly inhibited intracellular ROS production in LPS-induced BV2. Moreover, punicalagin also attenuated the mitochondrial ROS levels by LPS-induced BV2 cells (Figure 6).

Discussion

Neurodegenerative diseases are common in aging adults globally and cause serious health problems by progressive morbidity, memory and cognitive impairment. By 2020, about 50 million people worldwide were suffering from dementia, and as the most common type of neurodegenerative disease, Alzheimer’s disease accounts for 60% to 80% of causes of dementia. Neuroinflammation and oxidative damage are the key features of neurodegenerative diseases, and the hallmark of neuroinflammation is the activation of microglia in the central nervous system. Activated microglia in neurodegenerative processes induce the release of proinflammatory cytokines and mediators including IL-1β, IL-6, TNF-α and ROS, thereby causing neuronal cell degeneration. Currently, there is no effective treatment to slow or stop the progression of neurodegenerative diseases other than relying on supportive and symptomatic care; therefore, suppressing activated microglia-induced inflammatory responses have been considered as a therapeutic strategy for treating these diseases. In this study, our experimental results revealed that punicalagin effectively inhibited LPS-activated murine microglial BV2 cells by attenuating the secretion of inflammatory mediators and cytokines, inhibiting the activation of the NLRP3 inflammasome, and suppressing the production of intracellular and mitochondrial ROS.
Natural products are important sources for new drug development. Punicalagin is a major active component mainly found in pomegranate, a plant widely distributed in the tropics and subtropical regions. Previous studies have reported that punicalagin alleviates macrophage-mediated acute inflammation, acute lung injury and acute respiratory distress syndrome, acute kidney injury, and chronic diseases including arthritis, diabetes, obesity, cardiovascular and neurodegenerative diseases. Neuroinflammation is mediated by several proinflammatory molecules. As an inflammatory mediator, iNOS is one of the three different isoforms of NOS that expresses in glial cells, macrophages and neutrophils, and is only generated after induction by inflammatory mediators like cytokines or endotoxins. It has a neurotoxic effect on neurodegenerative diseases when generating higher concentrations of NO. Furthermore, inhibiting COX-2 and subsequent synthesis of PGE\textsubscript{2} leads to a decrease in neuronal degeneration. Proinflammatory cytokine IL-6 is reported to be associated with the pathogenesis of Alzheimer’s disease, and it may cause neuroinflammation by recruiting leukocytes across the blood-brain barrier and promoting the LPS-driven inflammatory responses by regulating the phosphorylation of STAT3. In addition, the MAPK cascade and NK-κB signaling pathway both regulate the expression and production of LPS-induced proinflammatory cytokines. NF-κB has been demonstrated to play an integral role in the progression of Alzheimer’s disease and ischemic stroke. A previous murine model in vivo and in vitro study revealed the anti-neuroinflammatory effect of punicalagin on microglia and astrocytes by inhibiting the production of proinflammatory cytokines including IL-1β, IL-6 and TNF-α and interfering with NF-κB signaling via binding to its subunit p50 directly. Similarly, our results demonstrated that punicalagin suppressed LPS-induced inflammatory responses by reducing the expression of iNOS, COX-2, inhibiting the secretion of NO, PGE\textsubscript{2} and IL-6 while suppressing the phosphorylation of STAT3, IκBα, and MAPKs in BV2 cells, indicating that punicalagin has potential in attenuating microglia-induced inflammation.
Previous studies have indicated that the activation of either NF-κB and MAPK signaling pathways was partly responsible for inducing the expression and activation of NLRP3 inflammasome proteins in neurons and brain tissue. Activation of NLRP3 inflammasome is associated with neurodegenerative diseases including Alzheimer’s disease, Parkinson’s disease and amyotrophic lateral sclerosis. Neuroinflammation and activation of microglia around the degenerating neurons with subsequent secretion of proinflammatory molecule IL-1β lead to neuronal damage. The canonical activation of the NLRP3 inflammasome, a multiprotein complex composed of NLRP3, ASC and caspase-1, is responsible for the production of IL-1β from microglia upon cellular stress as well as induction of pyroptosis, a type of programmed cell death that causes rupture of the cell membrane resulting in the release of more pro-inflammatory cytokines thereby promoting the inflammatory response.

Previous studies have indicated that the levels of NLRP3, ASC, and caspase-1 proteins were upregulated in the brain and plasma of patients with neurodegenerative diseases including Alzheimer’s disease, Parkinson’s disease, and amyotrophic lateral sclerosis. Neuroinflammation and activation of microglia around the degenerating neurons with subsequent secretion of proinflammatory molecule IL-1β lead to neuronal damage. The canonical activation of the NLRP3 inflammasome, a multiprotein complex composed of NLRP3, ASC and caspase-1, is responsible for the production of IL-1β from microglia upon cellular stress as well as induction of pyroptosis, a type of programmed cell death that causes rupture of the cell membrane resulting in the release of more pro-inflammatory cytokines thereby promoting the inflammatory response.

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Oxidative stress is an imbalance state between prooxidant and antioxidant species and is characterized by an increasing level of reactive species including ROS and nitrogen reactive species (RNS). Previous evidence shows that oxidative stress and neuroinflammation play important roles in the development and progression of neurodegenerative diseases because reactive species can become injurious under chronic oxidative stress by oxidizing intracellular proteins and lipids, causing DNA damage, and mediating activation of microglia and astrocytes that promote inflammatory responses.
disease is the most common type of neurodegenerative disease and its pathogenesis is associated with oxidative stress and amyloid-β, a major component of the senile plaques found in the pathology of patients with Alzheimer’s disease.\textsuperscript{1,2,53} Altogether, the increase of ROS production and intracellular Ca\textsuperscript{2+} cause excessive Ca\textsuperscript{2+} influx into mitochondria resulting in mitochondrial impairment and subsequently releasing pro-apoptotic molecules leading to neuronal degeneration and damage.\textsuperscript{53} Mitochondria are important cellular organelles and as being considered the powerhouse in cells, mitochondrial impairment leads to defective energy metabolism and excessive production of ROS.\textsuperscript{53}

Furthermore, damaged mitochondria might promote the activation of caspase-1, release of proinflammatory cytokines and activation of inflammasome formation.\textsuperscript{8} Recent studies have suggested antioxidants counteract the oxidative damage conferred by ROS as a therapeutic target in treating neurodegenerative diseases including Alzheimer’s disease.\textsuperscript{12,53} Punicalagin has been reported to have the potential of being an effective antioxidant in attenuating cardiac mitochondrial impairment in an obesity rat model\textsuperscript{15} while reducing the pro-inflammatory cytokines IL-1β, IL-6, and TNF-α as well as reactive species hydrogen peroxide (H\textsubscript{2}O\textsubscript{2}) in brain tissue of an LPS-stimulated mouse model.\textsuperscript{46} In line with our study, our results indicate that punicalagin presented an anti-oxidative effect by attenuating the production of intracellular ROS and mitochondrial ROS in LPS-activated microglia.

Conclusion
Our study demonstrated the anti-inflammatory effects of punicalagin through alleviating LPS-induced inflammation through MAPK and NF-κB signaling pathways and suppressing the activation of NLRP3 inflammasome in microglia. Furthermore, punicalagin also presented anti-oxidative effects in LPS-activated microglia by attenuating the production of both intracellular and mitochondrial ROS. Our results shed light on the molecular mechanism of anti-inflammatory and anti-oxidative effects of punicalagin as an agent possessing potential for treating neurodegenerative diseases.

Abbreviations
ASC, apoptotic speck containing protein with a CARD; ATP, adenosine triphosphate; COX-2, cyclooxygenase-2; DAMPs, damage-associated molecules patterns; ELISA, enzyme-linked immunosorbent assay; ERK, extracellular signal–regulated kinase; IL, interleukin; JNK, c-Jun N-terminal kinase; LPS, lipopolysaccharides; MAPK, mitogen-activated protein kinase; NF-κB, nuclear factor-κB; NLRP3, NOD-like receptor family pyrin domain containing 3; NO, nitric oxide; NOD, nucleotide binding oligomerization domain; PAMPs, pathogen-associated molecular patterns; PGE\textsubscript{2}, prostaglandin E\textsubscript{2}; ROS, reactive oxygen species; SEAP, secreted embryonic alkaline phosphatase; TLR-4, Toll-like receptor-4; TNF-α, tumor necrosis factor-α.

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Disclosure
The authors declare no conflicts of interest.

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