

# Neuroinflammation

Mutations in the [Complement factor I](#) (CFI) gene have previously been identified as causes of recurrent CNS inflammation

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[Cerebral autoregulation](#) is the ability of the brain to maintain a constant [blood flow](#) despite changes in [blood pressure](#). Dysfunction in cerebral [autoregulation](#) can occur in acute brain injury, such as [traumatic brain injury](#) or [stroke](#), and can lead to poor outcomes. [Neuroinflammation](#) has been suggested as a potential mechanism underlying cerebral autoregulation dysfunction in these conditions.

## Neuroinflammatory biomarkers

[Neuroinflammatory biomarkers](#).

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## Etiology

[Alcohol induced neuroinflammation](#).

SARS-CoV-2-mediated neuroinflammation

## Neuroinflammation treatment

[Microglia](#)-related [neuroinflammation](#) is associated with a variety of [neurodegenerative diseases](#). [Flavonoids](#) have demonstrated different pharmacological effects, such as antioxidation, [neuroprotection](#) and anti-inflammation. However, the effect of flavonoid 6-methoxyflavone (6-MeOF) on microglia-mediated neuroinflammation remain unknown.

A study aimed to study the antineuroinflammatory effects of 6-MeOF in lipopolysaccharide- (LPS-) induced microglia in vitro and in vivo.

Pretreatment of BV2 microglia cells with 6-MeOF for 1 h then stimulated with LPS (100 ng/ml) for 24 h. The expression levels of pro-inflammatory factors, NO and reactive oxygen species (ROS) were performed by the enzyme-linked immunosorbent assay (ELISA), Griess assay and flow cytometry. Western blotting was used to assess MAPK, NF-κB signal transducer and antioxidant enzymes-related proteins. Analysis of ROS and microglial morphology was confirmed in the zebrafish and mice brain, respectively.

The results demonstrated that 6-MeOF dose-dependently prevent cell death and decreased the levels of pro-inflammatory mediators in LPS-stimulated BV2 microglia cells. Phosphorylated NF-κB/IκB and TLR4/MyD88/p38 MAPK/JNK proteins after exposure to 6-MeOF was suppressed in LPS-activated BV-2

microglial cells. 6-MeOF also presented antioxidant activity by reduction of NO, ROS, iNOS and COX-2 and the induction of the level of HO-1 and NQO1 expressions in LPS-activated BV2 microglial cells. Furthermore, we demonstrated that 6-MeOF inhibited LPS-induced NO generation in an experimental zebrafish model and prevent the LPS-induced microgliosis in the prefrontal cortex and substantia nigra of mice.

These results explored that 6-MeOF possesses potential as anti-inflammatory and anti-oxidant agents against microglia-associated neuroinflammatory disorders <sup>1)</sup>.

<sup>1)</sup>

Chen WF, Shih YH, Liu HC, Cheng CI, Chang CI, Chen CY, Lin IP, Lin MY, Lee CH. 6-methoxyflavone suppresses neuroinflammation in lipopolysaccharide- stimulated microglia through the inhibition of TLR4/MyD88/p38 MAPK/NF-κB dependent pathways and the activation of HO-1/NQO-1 signaling. *Phytomedicine*. 2022 Mar 1;99:154025. doi: 10.1016/j.phymed.2022.154025. Epub ahead of print. PMID: 35272244.

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