



## **Echinacoside**

**Catalog No: tcsc3400** 

Available Sizes	
ize: 5mg	
ize: 10mg	
ize: 50mg	
Specifications	
<b>AS No:</b> 2854-37-3	
ormula:	
35 <sup>H</sup> 46 <sup>O</sup> 20	
athway: others	
<b>arget:</b> thers	
urity / Grade: 98%	
<b>olubility:</b> 0 mM in DMSO	
bserved Molecular Weight: 86.73	

## **Product Description**

Echinacoside is a natural polyphenolic compound, has various kinds of pharmacological activities, such as antioxidative, antiinflammatory, neuroprotective, hepatoprotective, nitric oxide radical-scavenging and vasodilative ones.





IC50 value:

Target:

in vitro: Echinacoside(ECH) dose dependently inhibited HEWL aggregation, and this inhibition occurred in different fiber-forming stages. ECH could also scavenge the DPPH and OH free radicals in a concentration-dependent manner. ECH could increase viability of rat pheochromocytoma PC12 cells injured by A $\beta$  and suppress the increase in intracellular reactive oxygen species (ROS) triggered by A $\beta$  [1]. Transient treatment with echinacoside inhibits cytochrome c release and caspase-3 activation caused by ensuing rotenone exposure via activating Trk-extracellular signal-regulated kinase (ERK) pathway in neuronal cells [2]. ECH caused a significant increase in cell proliferation, ALP activity, COL I contents, OCN levels and an enhancement of mineralization in osteoblasts at the concentration range from 0.01 to 10nmol·L(-1) (p

in vivo: In OVX rats, the increases of body weight, serum hydroxyproline (HOP) levels, and the decreases of uterus wet weight and BMD were significantly reversed by ECH treatment [3]. Echinacoside (60 mg/kg) was given intraperitoneally to mice at 1 h prior to GalN/LPS exposure. Pretreatment with echinacoside remarkably improved the survival rate of GalN/LPS-treated mice and attenuated acute hepatotoxicity, as demonstrated by decreased ALT levels and improved histological signs. Echinacoside shows both anti-apoptotic and anti-inflammatory properties, characterized by a substantial inhibition of hepatocyte apoptosis and a significant reduction in the inflammatory markers, including myeloperoxidase, extracellular nucleosomes, high-mobility group box 1, and inflammatory cytokines in the plasma of mice, which may be important mechanisms related to its protective effect [5].

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!