Cycloastragenol

| Cat. No.: | HY-N1485 |
|--------------------|--|
| CAS No.: | 78574-94-4 |
| Molecular Formula: | $C_{30}H_{50}O_{5}$ |
| Molecular Weight: | 490.72 |
| Target: | Telomerase |
| Pathway: | Cell Cycle/DNA Damage |
| Storage: | 4°C, protect from light * In solvent : -80°C, 6 months: -20°C, 1 month (protect from light) |

SOLVENT & SOLUBILITY

| In Vitro | DMSO : 66.67 mg/mL (135.86 mM; Need ultrasonic) | | | | | | |
|----------|--|--|-----------|------------|------------|--|--|
| | Preparing Stock Solutions | Solvent Mass Concentration | 1 mg | 5 mg | 10 mg | | |
| | | 1 mM | 2.0378 mL | 10.1891 mL | 20.3782 mL | | |
| | | 5 mM | 0.4076 mL | 2.0378 mL | 4.0756 mL | | |
| | | 10 mM | 0.2038 mL | 1.0189 mL | 2.0378 mL | | |
| | Please refer to the solubility information to select the appropriate solvent. | | | | | | |
| In Vivo | 1. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (5.09 mM); Clear solution | | | | | | |
| | 2. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (4.24 mM); Clear solution | | | | | | |
| | 3. Add each solvent o Solubility: ≥ 2.08 n | one by one: 10% DMSO >> 90% cor ng/mL (4.24 mM); Clear solution | n oil | | | | |

| DIOLOGICAL ACTIV | |
|------------------|--|
| Description | Cycloastragenol (Astramembrangenin), the active form of astragaloside IV, has anti-oxidant, anti-inflammatory, anti-aging, anti-apoptotic, and cardiovascular protective effects. Cycloastragenol is a potent telomerase activator and can lengthen telomeres. Cycloastragenol alleviates age-related bone loss and improves bone microstructure and biomechanical properties ^{[1][2][3]} . |
| In Vitro | Cycloastragenol (0.03-3 μM; 24-72 hours) promotes viability, osteoblastic differentiation, and mineralization in MC3T3-E1 cells ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. |

HO

Ĥ I

ЮH

ОН



In Vivo

Cycloastragenol (5, 10, 20 mg/kg) is injected intraperitoneally at the onset of reperfusion, 12 h later and then twice daily for up to three days. Cycloastragenol dose-dependently reduces brain infarct volume, significantly ameliorated functional deficits, and prevents neuronal cell loss in middle cerebral artery occlusion (MCAO) mice. Cycloastragenol suppresses the mRNA expression of pro-inflammatory cytokines, including TNF- α and IL-1 β , and inhibits the activation of microglia and astrocytes in the ischemic brain^[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Int J Mol Sci. 2023, Mar 31;24(7), 6554.
- Int J Mol Sci. 2023 Feb 6;24(4):3179.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Yu Y, et al. Cycloastragenol: An exciting novel candidate for age-associated diseases. Exp Ther Med. 2018;16(3):2175-2182.

[2]. Li M, et al. Cycloastragenol upregulates SIRT1 expression, attenuates apoptosis and suppresses neuroinflammation after brain ischemia. Acta Pharmacol Sin. 2020;41(8):1025-1032.

[3]. Yu Y, et al. Cycloastragenol prevents age-related bone loss: Evidence in d-galactose-treated and aged rats. Biomed Pharmacother. 2020;128:110304.

Caution: Product has not been fully validated for medical applications. For research use only.