

Matrine Datasheet

4th Edition (Revised in July, 2016)

[Product Information]

Name: Matrine

Catalog No.: CFN98835

Cas No.: 519-02-8

Purity: > 98%

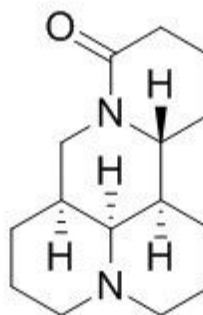
M.F: C₁₅H₂₄N₂O

M.W: 248.4

Physical Description: Powder

Synonyms: Alpha-Matrine; (+)-Matrine; MatrineAS; Matridin-15-one;

(5beta)-matridin-15-one; (5beta,6beta,7beta,11alpha)-matridin-15-one.



[Intended Use]

1. Reference standards;
2. Pharmacological research;
3. Cosmetic research;
4. Synthetic precursor compounds;
5. Intermediates & Fine Chemicals;
6. Others.

[Source]

The root of *Sophora japonica*.

[Biological Activity or Inhibitors]

Matrine, an alkaloid purified from the chinese herb *Sophora flavescens* Ait, is well known to possess activities including anti-inflammation, anti-fibrotic and anticancer, it could inhibit cell proliferation and induce apoptosis of SGC-7901 cells in vitro by up-regulating Fas/FasL expression and activating caspase-3 enzyme.^[1]

Matrine upregulates the cell cycle protein E2F-1 and triggers apoptosis via the mitochondrial pathway in K562 cells, it is a potential anti-drug.^[2]

Matrine suppresses PMA-induced MMP-1 expression through inhibition of the AP-1 signaling pathway and also may be beneficial for treatment of some inflammatory skin disorders.^[3]

Matrine can induce gastric cancer MKN45 cells apoptosis via increasing pro-apoptotic molecules of Bcl-2 family, also inhibits matrix metalloproteinase-9 expression and invasion of human hepatocellular carcinoma cells.^[4,5]

Matrine can improve 2,4,6-trinitrobenzene sulfonic acid (TNBS)-induced colitis in mice and the therapeutic mechanism might be related to the reduction of up-regulated colonic TNF- α production caused by TNBS.^[6]

Matrine seems to be a novel autophagy inhibitor that can modulate the maturation process of lysosomal proteases.^[7]

Matrine can reduce the mortality of acetaminophen overdosed mice more effectively, attenuate acetaminophen-induced hepatotoxicity, and reduces the number and area of γ -GT positive foci, thus protecting liver function and preventing HCC from occurring; it also has a protective effect on immunosuppression, a strong non-specific anti-inflammatory effect, and an effect of reducing the incidence of sodium and water retention.^[8]

[Solvent]

Chloroform, Dichloromethane, DMSO, Acetone.

[HPLC Method]^[9]

Mobile phase: Acetonitrile-Phosphate buffer solution(pH 5.5)=3:97 ;

Flow rate: 1.0 ml/min;

Column temperature: 30 °C;

The wave length of determination: 220 nm.

[Storage]

2-8°C, Protected from air and light, refrigerate or freeze.

[References]

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- [2] Hua J, Hou C H, Zhang S B, *et al.* *Eur. J. Pharmacol.*, 2007, 559(2-3):98-108.
- [3] Eunsun Jung , Jongsung Lee , Huh S, *et al.* *Biofactors*, 2008, 33(2):121–8.
- [4] Luo C, Zhu Y, Jiang T, *et al.* *Toxicol.*, 2007, 229(3):245-52.
- [5] Yu H B, Zhang H F, Li D Y, *et al.* *J. Asian Nat. Prod. Res.*, 2011, 13(3):242-50.
- [6] Hong C, Bing X, Lin Z, *et al.* *Pharmacol .Res.*, 2006, 53(3):202-8.
- [7] Wang Z, Zhang J, Wang Y, *et al.* *Carcinogenesis*, 2012, 34(1):128-38.
- [8] Wan X Y, Luo M, Li X D, *et al.* *Chem.-Biol. Interact.*, 2009, 181(1):15-9.
- [9] Jiang Y L, Zhang S Y. *Lishizhen Medicine & Materia Medica Research*, 2006, 17(11): 2204-5.

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