## 4,5-Dimethoxycanthin-6-one

## Chemical Properties

CAS No.:
18110-87-7
Formula: $\quad \mathrm{C} 16 \mathrm{H} 12 \mathrm{~N} 2 \mathrm{O} 3$
Molecular Weight: 280.3
Appearance: N/A
Storage: $\quad 0-4^{\circ} \mathrm{C}$ for short term (days to weeks), or $-20^{\circ} \mathrm{C}$ for long term (months).

## Biological Description

| Description | 4,5-Dimethoxycanthin-6-one has antibacterial actiyity, it exhibits inhibition against Staphylococcus aureus and <br> its drug-resistant strains. 4,5-Dimethoxycanthin-6-one has a strong inhibitory effect on cyclic adenosine <br> monophosphate (cAMP) phosphodiesterase. It also shows cytotoxicity against the tumor cell lines, U937 and <br> HepG2. |
| :--- | :--- |
| Targets(IC50) | Antifection: None <br> cAMP: None |
| In vitro | Six alkaloid compounds were isolated from the chloroform soluble fraction of the methanolic extract of the <br> wood of Picrasma quassioides Benn (Simarobaceae) as the cytotoxic components against the tumor cell lines, <br> U937 and HepG2.METHODS AND RESULTS: The compounds were identified as 4,9-dimethoxy-1-vinyl- $\gamma$ - <br> carboline (1), 1-carbomethoxy- $\beta$-carboline (2), 4,5-Dimethoxycanthin-6-one (3), 5-hydroxy-4-methoxycanthin- <br> $6-$ 6-one (4), 3-methoxycanthin-5,6-dione (5) and 4,8-dimethoxy-1-vinyl- $\beta$-carboline (6) by spectroscopic analysis. <br> Among them, compound 5 showed the most potent cytotoxicity in a dose-dependent manner against the two <br> tumor cell lines. CONCLUSIONS:This is the first report of the cytotoxic effect of those isolated compounds 1-5 on on <br> U937 and HepG2 cell lines. |

## Solubility Information

$\square$
Preparing Stock Solutions

|  | $\mathbf{1 m g}$ | $\mathbf{5 m g}$ | $\mathbf{1 0 m g}$ |
| :--- | :---: | :---: | :---: |
| 1 mM | 3.568 mL | 17.838 mL | 35.676 mL |
| 5 mM | 0.714 mL | 3.568 mL | 7.135 mL |
| 10 mM | 0.357 mL | 1.784 mL | 3.568 mL |
| 50 mM | 0.071 mL | 0.357 mL | 0.714 mL |

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. The storage conditions and period of the stock solution: - $80^{\circ} \mathrm{C}$ for 6 months; - $20^{\circ} \mathrm{C}$ for 1 month. Please use it as soon as possible.

Reference

1. Cytotoxic alkaloids from the wood of Picrasma quassioides. J. Korean Soc. Appl. Biol. Chem.,2009, 52(6):663-7.
2. Inhibitors of cyclic AMP phosphodiesterase in Picrasma quassioides Bennet, and inhibitory activities of related .BETA.-carboline alkaloids. Chem Pharm Bull (Tokyo). 1984 May;32(5):1872-7.
3. Chemical investigation of the alkaloids of Ku-Mu [Picrasma quassioides (D. Don) Benn. Yao Xue Xue Bao,1979, 14(3):167-77.

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