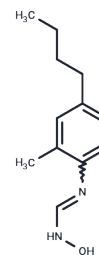


HET0016

Chemical Properties

CAS No. : 339068-25-6
Formula: C₁₂H₁₈N₂O
Molecular Weight: 206.28
Appearance: no data available
Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year



Biological Description

| | |
|----------------------------|---|
| Description | HET0016 is a potent and selective 20-HETE synthase inhibitor (IC ₅₀ s: 17.7 nM, 12.1 nM, and 20.6 nM for recombinant CYP4A1-, CYP4A2-, and CYP4A3-catalyzed 20-HETE synthesis) and a selective CYP450 inhibitor, shown to inhibit angiogenesis and tumor growth. |
| Targets(IC ₅₀) | Others |
| In vitro | HET0016 (100 μM; 24 hours, 48 hours) decreases migration and invasion of breast cancer metastatic cells. HET0016 is a selective, non-competitive and irreversible inhibitor of CYP4A. |
| In vivo | HET0016 decreases expression of pro-inflammatory and growth factors and granulocytic MDSCs population in lung microenvironment. HET0016 protects BBB dysfunction after I/R by regulating the expression of MMP-9 and tight junction proteins. HET0016 (10 mg/kg/day; i.v.; for 3 weeks) reduces tumor volume and lung metastasis in an immunocompetent breast cancer mouse model. HET0016 reduces the metalloproteinases' levels in the lungs via PI3K/AKT pathway in mice. |

Solubility Information

| | |
|------------|---|
| Solubility | DCM: 12.5 mg/mL (60.60 mM), Sonication is recommended. DMSO: 5 mg/mL (24.24 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble) |
|------------|---|

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|------------|------------|
| 1 mM | 4.8478 mL | 24.2389 mL | 48.4778 mL |
| 5 mM | 0.9696 mL | 4.8478 mL | 9.6956 mL |
| 10 mM | 0.4848 mL | 2.4239 mL | 4.8478 mL |
| 50 mM | 0.097 mL | 0.4848 mL | 0.9696 mL |

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Reference

- Seki T, et al. Cytochrome P450 4A isoform inhibitory profile of N-hydroxy-N'-(4-butyl-2-methylphenyl)-formamidine (HET0016), a selective inhibitor of 20-HETE synthesis. *Biol Pharm Bull.* 2005 Sep;28(9):1651-4.
- Borin TF, et al. HET0016 decreases lung metastasis from breast cancer in immune-competent mouse model. *PLoS One.* 2017 Jun 13;12(6):e0178830.
- Liu Y, et al. The protective effect of HET0016 on brain edema and blood-brain barrier dysfunction after cerebral ischemia/reperfusion. *Brain Res.* 2014 Jan 28;1544:45-53.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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