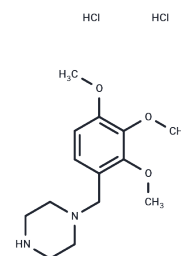


Trimetazidine dihydrochloride

Chemical Properties

| | |
|-------------------|---|
| CAS No. : | 13171-25-0 |
| Formula: | C ₁₄ H ₂₄ Cl ₂ N ₂ O ₃ |
| Molecular Weight: | 339.258 |
| Appearance: | no data available |
| Storage: | Powder: -20°C for 3 years In solvent: -80°C for 1 year |



Biological Description

| | |
|---------------|---|
| Description | Trimetazidine dihydrochloride (Vastarel F) can improve myocardial glucose utilization by inhibiting fatty acid metabolism. Trimetazidine is the first cytoprotective anti-ischemic agent, also used as a vasodilator in ischemic heart disease or angina. |
| Targets(IC50) | Autophagy,Fatty Acid Synthase |
| In vitro | Trimetazidine up-regulates miR-21 expression, then miR-21 targets PTEN increasing the PI3K pathway and finally the activation of this pathway counteracts the apoptotic effect of hypoxia/reperfusion[3]. |
| In vivo | The administration of TMZ reduces myocardial infarction size in WT C57BL/6J hearts. Both AMPK and ERK signaling pathways mediate the cardioprotection of TMZ against ischemic injury. Trimetazidine Shifts Metabolism from Fatty Acid Oxidation to Glucose Oxidation and improves Contractile Functions of Cardiomyocytes during Hypoxia[2]. |
| Cell Research | The H9C2 cells are randomly assigned into three groups: Sham group, in which the cells are treated with 0μM TMZ for 48 h and then cultured under normal oxygenation conditions (5% CO ₂ , and 95% air); H/R group, in which cells are treated with 0 μM TMZ for 48 h and then cultured under H/R conditions; and H/R+TMZ group, in which cells are treated with 10 μM TMZ for 48 h and then subjected to H/R treatment. (Only for Reference) |

Solubility Information

| | |
|------------|--|
| Solubility | DMSO: 65 mg/mL (191.59 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble) |
|------------|--|

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|-----------|------------|
| 1 mM | 2.9476 mL | 14.738 mL | 29.4759 mL |
| 5 mM | 0.5895 mL | 2.9476 mL | 5.8952 mL |
| 10 mM | 0.2948 mL | 1.4738 mL | 2.9476 mL |
| 50 mM | 0.059 mL | 0.2948 mL | 0.5895 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

Kantor PF, et al. Circ Res. 2000, 86(5):580-588.

Chen B, Huang Y, He S, et al. N6-methyladenosine modification in 18S rRNA promotes tumorigenesis and chemoresistance via HSF4b/HSP90B1/mutant p53 axis. Cell Chemical Biology. 2023, 30(2): 144-158. e10.

Liu Z, et al. Metabolism. 2016, 65(3):122-130.

Yang Q, et al. Int J Clin Exp Pathol. 2015, 8(4):3735-3741.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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