

Metformin (hydrochloride)

Catalog No: tcsc1851

Available Sizes

Size: 10g

Size: 50g

Specifications

CAS No:

1115-70-4

Formula:

 $C_4H_{12}CIN_5$

Pathway: Autophagy;Epigenetics;PI3K/Akt/mTOR;Autophagy

Target:

Autophagy;AMPK;AMPK;Mitophagy

Purity / Grade:

>98%

Solubility: DMSO : ≥ 1.7 mg/mL (10.26 mM); H2O : ≥ 32 mg/mL (193.21 mM)

Alternative Names:

1,1-Dimethylbiguanide hydrochloride

Observed Molecular Weight:

165.62

Product Description

Metformin (hydrochloride) is a first-line drug for the treatment of type 2 diabetes and there is increasing evidence of a potential efficacy of this agent as an anti-cancer drug.

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In Vitro: Metformin inhibits proliferation of ESCs in a concentration-dependent manner. The IC₅₀ is 2.45 mM for A-ESCs and 7.87 mM for N-ESCs. Metformin shows pronounced effects on activation of AMPK signaling in A-ESCs from secretory phase than in cells from proliferative phase^[3]. Metformin (0-500 μ M) decreases glycogen synthesis in a dose-dependent manner with an IC₅₀ value of 196.5 μ M in cultured rat hepatocytes^[4]. Metformin shows cell viability and cytotoxic effects on PC-3 cells with IC₅₀ of 5 mM^[5].

In Vivo: Metformin (100 mg/kg, p.o.) alone, and metformin (25, 50, 100 mg/kg) with isoproterenol groups attenuates myocyte necrosis through histopathological analysis^[1]. Metformin (> 900 mg/kg/day, p.o.) results in moribundity/mortality and clinical signs of toxicity in CrI:CD(SD) rats^[2].



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