

Selisistat

Catalog No: tcsc0960



Available Sizes

Size: 5mg

Size: 10mg

Size: 25mg



Specifications

CAS No:

49843-98-3

Formula:

$C_{13}H_{13}ClN_2O$

Pathway:

Epigenetics;Cell Cycle/DNA Damage

Target:

Sirtuin;Sirtuin

Purity / Grade:

>98%

Solubility:

DMSO : ≥ 32 mg/mL (128.66 mM)

Alternative Names:

EX-527

Observed Molecular Weight:

248.71

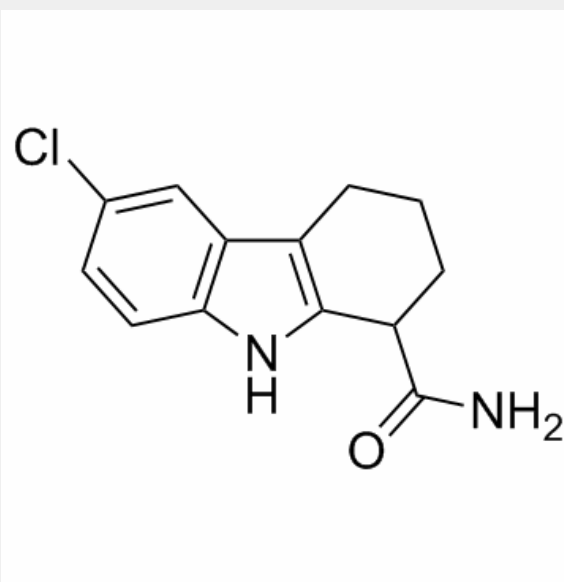
Product Description

Selisistat (EX-527) is a potent and selective **SIRT1** inhibitor with **IC₅₀** of 98 nM.

IC50 & Target: IC50: 98 nM (SIRT1), 19.6 μ M (SIRT2), 48.7 μ M (SIRT3)^[4]

In Vitro: Selisistat (EX-527) is an inhibitor of SIRT1 enzymatic activity (IC₅₀, 98 nM), identified in a high-throughput screen using bacterially expressed human SIRT1. Selisistat (EX-527) inhibits SIRT1 in a concentration-dependent manner with an IC₅₀ of 38 nM, in agreement with the activity on bacterially expressed SIRT1. Selisistat (EX-527) has much lower potency against SIRT2 (IC₅₀, 19.6 μ M) or SIRT3 (IC₅₀, 48.7 μ M) but does not inhibit class I/II HDAC activity at concentrations up to 100 μ M^[1]. Selisistat (EX-527) exerts an inhibitory effect on SIRT1 activity without affecting SIRT1 expression on both mRNA and protein levels^[2].

In Vivo: Selisistat (EX-527) abolishes Resveratrol (RSV)-induced attenuation of microvascular inflammation in *ob/ob* septic mice. Finally, *ob/ob* mice in Sepsis+RSV group has significantly increased 7-day survival vs. Sepsis+Vehicle group^[3].



All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!