

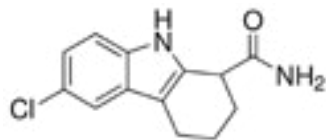
EX-527

Catalog Number P005-5MG

Catalog Number P005-25MG

FEATURES

- Selective SIRT1 inhibitor
- Does not inhibit HDACs
- Inhibitor of SIRT1 deacetylation of p53



ARBOR
ASSAYS

INTRODUCTION

EX-527 is a cell-permeable, selective inhibitor of SIRT1 ($IC_{50} = 98 \text{ nM}$ *in vivo*, 38 nM *in vitro*) that does not inhibit histone deacetylase (HDAC) or other sirtuin deacetylase family members (IC_{50} values are $20 \mu\text{M}$, $49 \mu\text{M}$, $> 100 \mu\text{M}$ and $> 100 \mu\text{M}$ for SIRT2, SIRT3, HDAC and NADase respectively). EX-527 has been used to investigate the relationship between SIRT1-mediated deacetylation of p53, p53 activity, and cell survival following DNA damage.

FORM: Light yellow powder

MOLECULAR WEIGHT: 284.71

STORAGE: 4°C , desiccated

FORMULA: $\text{C}_{13}\text{H}_{13}\text{ClN}_2\text{O}$

CAS NUMBER: 49843-98-3

OTHER NAMES: 6-Chloro-2,3,4,9-tetrahydro-1*H*-carbazole-1-carboxamide, racemate

USES: Soluble to 18 mg/mL in DMSO and 10 mg/mL in Ethanol

REFERENCES:

Napper, AD, et al. Discovery of indoles as potent and selective inhibitors of the deacetylase SIRT1. *J. Med. Chem.* 48:8045. (2005)

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Nayagam, V.M., Wang, X., Tan, Y.C., et al. SIRT1 modulating compounds from high-throughput screening as anti-inflammatory and insulin-sensitizing agents. *J. Biomol. Screen* 11:959-967 (2006)

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