Scriptaid

Cat. No.: HY-15489
CAS No.: 287383-59-9
Molecular Formula: C₁₈H₁₈N₂O₄
Molecular Weight: 326.35
Target: HDAC; Autophagy
Pathway: Cell Cycle/DNA Damage; Epigenetics; Autophagy
Storage: Powder -20°C 3 years
        4°C  2 years
        In solvent -80°C 6 months
        -20°C 1 month

Solvent & Solubility

In Vitro

DMSO : ≥ 150 mg/mL (459.63 mM)
* “≥” means soluble, but saturation unknown.

Preparing Stock Solutions

<table>
<thead>
<tr>
<th>Solvent Concentration</th>
<th>Mass 1 mg</th>
<th>Mass 5 mg</th>
<th>Mass 10 mg</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 mM</td>
<td>3.0642 mL</td>
<td>15.3210 mL</td>
<td>30.6419 mL</td>
</tr>
<tr>
<td>5 mM</td>
<td>0.6128 mL</td>
<td>3.0642 mL</td>
<td>6.1284 mL</td>
</tr>
<tr>
<td>10 mM</td>
<td>0.3064 mL</td>
<td>1.5321 mL</td>
<td>3.0642 mL</td>
</tr>
</tbody>
</table>

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description
Scriptaid is a potent histone deacetylase (HDAC) inhibitor, used in cancer research.

IC₅₀ & Target
HDAC

In Vitro
Scriptaid (1 µg/mL) treatment inhibits cell growth in breast cancer cell lines, results in increased accumulation of both acetyl H3 and acetyl H4 proteins in MDA-MB-231, MDA-MB-435, and Hs578t cells. Scriptaid also inhibits cell growth of MDA-MB-231, MDA-MB-435, and Hs578t cell lines, with IC₅₀ values of 0.5-1.0 µg/mL. Scriptaid (0.1-1.0 µg/mL) induces ER and PR mRNA expression in a dose dependent manner; when it is combined with AZA, they enhance ER expression and induce a functional ER protein[1]. Scriptaid and SAHA preferentially inhibit the Class I histone deacetylases, hda1, 2, and 3. Scriptaid is a potent anti-T. gondii compound with low cytotoxicity, and the IC₅₀ is 39 nM. Scriptaid has atypical effects in T. gondiiinfected H568 cells[2]. Scriptaid inhibits the growth of HeLa cells with IC₅₀ of 2 µM at 48 h in a dose-dependent manner. Scriptaid also affects cell-cycle and apoptosis[3].
In Vivo

Scriptaid (3.5 µg/g mouse, i.p.) clearly inhibits tumor growth in a xenograft mouse model[1].

PROTOCOL

Cell Assay [1]

IC₅₀ concentrations of Scriptaid are determined in MDA-MB-231, MDA-MB-435 and Hs578t cells by MTT assay. For cell growth assays, MDA-MB-231, MDA-MB-435, and Hs578t cells are plated at a cell density of 5000 cells/well in 12 well plates and treated with 1.0 µg/mL Scriptaid for up to 3 days. Cells are counted daily using a Coulter counter. Percent growth inhibition is determined by comparison of treated and untreated cells[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Administration [1]

Four to six week old athymic female nude mice are housed under laminar flow hoods in an environmentally controlled pathogen free animal facility for the duration of experiments. Mice are injected with 2×10⁶ MDA-MB-231 human breast cancer cells into each flank. Tumors are allowed to grow to approximately 0.1 cm³ in diameter before treatment. Mice are then treated with Scriptaid (3.5 µg/g mouse), TSA (0.5 µg/g mouse), or DMSO vehicle intraperitoneally for five consecutive days with 2 days rest each week for a total of 4 weeks. Individual tumor measurements are recorded from each flank weekly[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES


Caution: Product has not been fully validated for medical applications. For research use only.

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