

## **MS-275 (Entinostat)**

Catalog #: 27011 Lot #: 150715

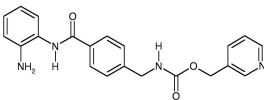
Size: 25 mg Structure:

CAS Registry #: 209783-80-2

Purity: ≥99%

Chemical Formula: C<sub>21</sub>H<sub>20</sub>N<sub>4</sub>O<sub>3</sub>

Molecular Weight: 376.41



**Description:** MS-275, also known as Entinostat, is an inhibitor of histone deacetylases (HDACs) and preferentially inhibits HDAC1 over HDAC3. It does not inhibit HDAC8 (IC $_{50} > 100$   $\mu$ M). It induces cyclin-dependent kinase inhibitor 1A (p21/CIP1/WAF1), slowing cell growth, differentiation, and tumor development *in vivo*. Recent studies suggest that MS-275 may be particularly useful as an antineoplastic agent when combined with other drugs like adriamycin, inhibitors of PARPs, or inhibitors of Hsp90.

Appearance: White to off-white crystalline solid

**Solubility:** Soluble in DMSO at 25 mg/ml with slight warming. MS-275 is poorly soluble in ethanol and water.

**Biological Activity:** MS-275 inhibits HDAC1 with an IC<sub>50</sub> of 300 nM and HDAC3 with an IC<sub>50</sub> value of 8  $\mu$ M.

Storage/Stability: Store at or below -20 °C.

**Quality Control:** The purity was determined by HPLC analysis.

## References:

- 1. Hu., E. et al., J. Pharmacol. Exp. Ther. 2003; **307**: 720-728.
- 2. Saito, A., et al., Proc. Natl. Acad. Sci. USA 1999; 96: 4592-4597.
- 3. Jaboin, J., et al., Cancer Res. 2002; 62: 6108-6115.
- 4. Xu, J. et al., Cancer Res. 2008; (68)16: 6718 6726.
- 5. Gaymes, T.J., et al., Haematologica 2009; **94**: 638-646.