# PRODUCT INFORMATION



# SAHA

Item No. 10009929

CAS Registry No.: 149647-78-9

 $N^1$ -hydroxy- $N^8$ -phenyl-octanediamide Formal Name: Synonyms: Suberoylanilide Hydroxamic Acid,

Vorinostat

MF:  $C_{14}H_{20}N_2O_3$ 264.3 FW: ≥98% **Purity:** 

UV/Vis.:  $\lambda_{\text{max}}$ : 243 nm Supplied as: A crystalline solid

Storage: -20°C Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

## **Laboratory Procedures**

SAHA is supplied as a crystalline solid. A stock solution may be made by dissolving the SAHA in an organic solvent purged with an inert gas. SAHA is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of SAHA in ethanol is approximately 0.25 mg/ml and approximately 20 mg/ml in DMSO and DMF.

For maximum solubility in aqueous buffers, SAHA should first be dissolved in DMSO and then diluted with the aqueous buffer of choice. SAHA has the solubility of approximately 0.5 mg/ml in a 1:1 solution of DMSO:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

## Description

SAHA is a histone deacetylase (HDAC) inhibitor that binds directly to the catalytic site of the enzyme thereby blocking substrate access. It inhibits class I, II, and IV HDACs at 50-200 nM and arrests cell growth in a wide variety of transformed cells in culture at 2.5-5.0 µM.<sup>1</sup>

## Reference

1. Marks, P.A. and Breslow, R. Dimethyl sulfoxide to vorinostat: Development of this histone deacetylase inhibitor as an anticancer drug. Nature Biotechnology 25(1), 84-90 (2007).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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