PRODUCT INFORMATION



3-TYP

Item No. 29660

CAS Registry No.: 120241-79-4

Formal Name: 3-(1H-1,2,3-triazol-5-yl)-pyridine

MF: $C_7H_6N_4$ FW: 146.2 **Purity:** UV/Vis.: λ_{max} : 245 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

3-TYP is supplied as a crystalline solid. A stock solution may be made by dissolving the 3-TYP in the solvent of choice, which should be purged with an inert gas. 3-TYP is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of 3-TYP in ethanol is approximately 5 mg/ml and approximately 10 mg/ml in DMSO and DMF.

3-TYP is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, 3-TYP should first be dissolved in DMF and then diluted with the aqueous buffer of choice. 3-TYP has a solubility of approximately 0.16 mg/ml in a 1:5 solution of DMF:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

3-TYP is a sirtuin 3 (SIRT3) inhibitor (IC₅₀ = 38 μ M).¹ It is selective for SIRT3 over SIRT1 and SIRT2 at 1 mM. 3-TYP (1 mM) increases acetylation of lysine residues in mitochondrial proteins in HeLa cells. It inhibits the growth of CCRF-CEM, COLO 205, OVCAR-4, OVCAR-8, and SKOV3 cells by greater than 50% when used at a concentration of 10 μ M.

Reference

1. Galli, U., Mesenzani, O., Coppo, C., et al. Identification of a sirtuin 3 inhibitor that displays selectivity over sirtuin 1 and 2. Eur. J. Med. Chem. 55, 58-66 (2012).

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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