## TC-E 5002

| ALTERNATE NAME: | 3-[9-cyclopropylnonanoyl(hydroxy)amino]propanoic acid; N -(9-Cyclopropyl-1-oxononyl)-N-hydroxy-beta- <br> alanine; 3 -(9-cyclopropyl-N-hydroxynonanamido)propanoic acid |
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| CATALOG \#: | B2856-1 $\quad 1 \mathrm{mg}$ <br>  <br> B2856-5 $\quad 5 \mathrm{mg}$ |

## STRUCTURE:

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| MOLECULAR FORMULA: | $\mathrm{C}_{15} \mathrm{H}_{27} \mathrm{NO}_{4}$ |
| MOLECULAR WEIGHT: | 285.38 |
| CAS NUMBER: | 1453071-47-0 |
| APPEARANCE: | A crystalline solid |
| PURITY: | $\geq 98 \%$ |
| SOLUBILITY: | $\sim 30 \mathrm{mg} / \mathrm{ml}$ in DMSO <br> $\sim 50 \mathrm{mg} / \mathrm{ml}$ in Ethanol <br> $\sim 10 \mathrm{mg} / \mathrm{ml}$ in DMF <br> $\sim 0.25 \mathrm{mg} / \mathrm{ml}$ in PBS, pH 7.2 |

DESCRIPTION: TC-E 5002 is an inhibitor of the histone lysine demethylase (KDM) subfamily KDM2/7. It potently inhibits KDM2A, KDM7A and KDM7B with an $\mathrm{IC}_{50}$ of $6.8 \mu \mathrm{M}, 0.2 \mu \mathrm{M}$ and $1.2 \mu \mathrm{M}$ respectively. It inhibits the growth of KYSE150 and HeLa cells with $\mathrm{GI}_{50}$ values of $16 \mu \mathrm{M}$ and $40 \mu \mathrm{M}$ respectively and arrests cells in the $\mathrm{G}_{0} / \mathrm{G}_{1}$ phase of the cell cycle. It inhibits demethylation of H 3 K 27 me 2 at similar concentrations.

STORAGE TEMPERATURE: $-20^{\circ} \mathrm{C}$

| HANDLING: | Do not take internally. Wear gloves and mask when handling the product! Avoid contact by all modes of <br> exposure. |
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| REFERENCE: | Suzuki, T., Ozasa, H., Itoh, Y., et al. Identification of the KDM2/7 histone lysine demethylase subfamily <br> inhibitor and its antiproliferative activity. J. Med. Chem. 56(18), 7222-7231 (2013). |

## RELATED PRODUCTS:

HDM Inhibitor, 2,4-PDCA (Cat. No. 2304)
RN 1 dihydrochloride (Cat. No. B2080)
Lysine-specific Demethylase Inhibitor (1C) (Hydrochloride) (Cat. No. B2814)
DDP-38003 dihydrochloride (Cat. No. B1958)
2,4-Pyridinedicarboxylic Acid (Cat. No. B2809)

DISCLAIMER:
FOR RESEARCH USE ONLY! Not to be used on humans.

