KPZ560

Cat. No.: CAS No.: Molecular Formula: Molecular Weight: Target: Pathway: Storage:	HY-156422 2397562-43-3 C ₂₆ H ₂₁ N ₅ O ₃ S ₂ 515.61 HDAC Cell Cycle/DNA Damage; Epigenetics Please store the product under the recommended conditions in the Certificate of	$H_{2}N$
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	

BIOLOGICAL ACTIVITY				
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Description	KPZ560 is a potent inhibitor of HDAC1 and HDAC2, with IC ₅₀ s of 12 nM and 68 nM, respectively. KPZ560 can increase in the spine density of granule neuron dendrites of mice and inhibitor cell growth of breast cancer cell line MCF ^[1] .			
IC₅₀ & Target	HDAC1 12 nM (IC ₅₀)	HDAC2 68 nM (IC ₅₀)		

REFERENCES

[1]. Yukihiro Itoh, et al. Discovery of Selective Histone Deacetylase 1 and 2 Inhibitors: Screening of a Focused Library Constructed by Click Chemistry, Kinetic Binding Analysis, and Biological Evaluation. J Med Chem. 2023 Oct 17. doi: 10.1021/acs.jmedchem.3c01095. Online ahead of print.

Product Data Sheet

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

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