## **Product** Data Sheet

## Aha1/Hsp90-IN-1

 Cat. No.:
 HY-151337

 CAS No.:
 2768265-58-1

 Molecular Formula:
  $C_{22}H_{17}F_3N_4O_2$ 

Molecular Weight: 426.39

Target: Microtubule/Tubulin

Pathway: Cell Cycle/DNA Damage; Cytoskeleton

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

## **BIOLOGICAL ACTIVITY**

Description	Aha1/Hsp90-IN-1 (Compound 17) is an Aha1/Hsp90 complex inhibitor. Aha1/Hsp90-IN-1 disrupts Aha1/Hsp90 interactions with an IC $_{50}$ of 3.32 $\mu$ M. Aha1/Hsp90-IN-1 inhibits tau aggregation [1].	
IC <sub>50</sub> & Target	IC $_{50}$ : 3.32 $\mu$ M (Aha1/Hsp90 interactions) $^{[1]}$	
In Vitro	Aha1/Hsp90-IN-1 (Compound 17) (10 $\mu$ M; 24 h) disrupts the Aha1/Hsp90 complex without direct inhibition of Hsp90 protein folding activity <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Western Blot Analysis <sup>[1]</sup>	
	Cell Line:	SH-SY5Y neuroblastoma cells and Her2 overexpressing SK-BR-3 breast cancer cells
	Concentration:	10 μΜ
	Incubation Time:	24 h
	Result:	Exhibited the ability to disrupt interactions between Aha1 and Hsp90. Did not induce the degradation of Hsp90 client proteins Her2 (in SK-BR-3 cells), Cdk6, or pAkt <sup>S473</sup> (in SH SY5Y cells), nor do they induce the expression of Hsp70, a marker of the heat shock response.

## **REFERENCES**

[1]. Keegan BM, et al. Synthesis and Evaluation of Small Molecule Disruptors of the Aha1/Hsp90 Complex for the Reduction of Tau Aggregation. ACS Med Chem Lett. 2022 Apr 15;13(5):827-832.

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

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