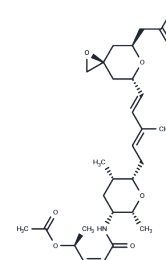


Thailanstatin D

Chemical Properties

CAS No. :	1609105-89-6
Formula:	C ₂₈ H ₄₁ N ₀ O ₈
Molecular Weight:	519.635
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year Actual storage temperature shall be subject to the COA.



Biological Description

Description	Thailanstatin D, an analogue of Thailanstatin A, inhibits AR-V7 gene splicing by disrupting the interaction between U2AF65 and SAP155, hindering their binding to the polypyrimidine tract situated between the branch point and the 3' splice site. This compound displays potent tumor inhibitory properties in human castration-resistant prostate cancer (CRPC) xenografts, resulting in cellular apoptosis.
Targets(IC50)	Apoptosis,Others,DNA/RNA Synthesis
In vitro	Thailanstatin D (0-50 nM; 4 hours) significantly reduces AR-V7 and other AR splice variants (AR-Vs)[1] protein levels and dose-dependently suppresses mRNA levels of AR-V7 and AR-FL in VCaP, 22RV1, and LN95 cell lines, as observed in a Cell Viability Assay[1].
In vivo	Thailanstatin D, administered at a dosage of 300 µg/kg through ALZET osmotic pumps daily for four days, significantly inhibited tumor growth in an animal model of NOD-SCID mice bearing 22RV1 xenografts.

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	1.9244 mL	9.622 mL	19.2441 mL
5 mM	0.3849 mL	1.9244 mL	3.8488 mL
10 mM	0.1924 mL	0.9622 mL	1.9244 mL
50 mM	0.0385 mL	0.1924 mL	0.3849 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

Wang B, et al. Developing new targeting strategy for androgen receptor variants in castration resistant prostate cancer. Int J Cancer. 2017;141(10):2121-2130.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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