Fostemsavir

Catalog No.C12531 Batch:C1253103

Product Datasheet

Technical Data

Formula	$C_{25}H_{26}N_7O_8P$					
Molecular Weight	583.49	**CAS No.**	864953-29-7			
Solubility (25°C)*	In vitro	DMSO	100 mg/mL (171.38 mM)			
		Water	Insoluble			
		Ethanol	Insoluble			
	In vivo (Add solvents to the product individually and in order)	Injection	Clear solution	5%DMSO [1]→ 40%PEG300 [2]→ 5%Tween80 [3]→ 50%ddH2O Validated by Capsid Bio labs. Should you need adjustments to this formulation, contact our sales team for custom testing.	5mg/ml (8.57mM)	Taking the 1 mL working solution as an example, add 50 μL of 100 mg/ml clarified DMSO stock solution to 400 μL of PEG300, mix evenly to clarify it; add 50 μL of Tween80 to the above system, mix evenly to clarify; then continue to add 500 μL of ddH2O to adjust the volume to 1 mL. The mixed solution should be used immediately for optimal results.
			Clear solution	5% DMSO [1]→ 95% Corn oil Validated by Capsid Bio labs. Should you need adjustments to this formulation, contact our sales team for custom testing.	0.25mg/ml (0.43mM)	Taking the 1 mL working solution as an example, add 50 μL of 5 mg/ml clear DMSO stock solution to 950 μL of corn oil and mix evenly. The mixed solution should be used immediately for optimal results.

* <1 mg/ml means slightly soluble or insoluble.
* Please note that Capsid Bio tests the solubility of all compounds in-house, and the actual solubility may differ slightly from published values. This is normal and is due to slight batch-to-batch variations.
* Room temperature shipping (Stability testing shows this product can be shipped at 2-8°C for storage purposes.)

Preparing Stock Solutions

Biological Activity

Description	Fostemsavir (BMS-663068, Rukobia), the phosphonooxymethyl prodrug of BMS626529, is an attachment inhibitor that targets **HIV-1 gp120** and prevents its binding to CD4+ T cells with **EC50** of < 10 nM.
Targets	HIV-1 gp120 [1] (Cell-free assay) 10 nM(EC50)
In vivo	BMS-663068 is the phosphonooxymethyl prodrug of BMS-626529 designed to increase the solubility of the compound in the gut. BMS-663068 is cleaved by alkaline phosphatase, located on the luminal surface of the small intestine brush border membranes, releasing BMS-626529. Due to its good membrane permeability, BMS-626529 is then rapidly absorbed.[1] BMS-663068 is an orally active inhibitor of HIV-1 attachment.[2]

Protocol (from reference)

Animal Study:[2]

- **Animal Models**
  C57BL/7 mice, NSG-SGM3-derived HSC-Hu mice
- **Dosages**
  20 mg/kg
- **Administration**
  Oral gavage

References

https://pubmed.ncbi.nlm.nih.gov/22547625/

https://pubmed.ncbi.nlm.nih.gov/32792548/

RETURN POLICY

Capsid Bio's Unconditional Return Policy ensures a smooth online shopping experience for our customers. If you are in any way unsatisfied with your purchase, you may return any item(s) within 7 days of receiving it. In the event of product quality issues, either protocol related or product related problems, you may return any item(s) within 365 days from the original purchase date. Please follow the instructions below when returning products.

SHIPPING & STORAGE

Capsid Bio products are transported at room temperature. If you receive the product at room temperature, please rest assured, the Capsid Bio Quality Inspection Department has conducted experiments to verify that the normal temperature placement of one month will not affect the biological activity of powder products. After collecting, please store the product according to the requirements described in the datasheet. Most Capsid Bio products are stable under the recommended conditions.

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