



Morphothiadin

Catalog No: tcsc0031547

Available Sizes
Size: 1mg
Size: 5mg
Size: 10mg
Size: 25mg
Size: 50mg
Size: 100mg
Specifications
CAS No: 1092970-12-1
Formula: C ₂₁ H ₂₂ BrFN ₄ O ₃ S
Pathway: Anti-infection
Target: HBV
Purity / Grade: >98%
Solubility: DMSO : 60 mg/mL (117.79 mM; Need ultrasonic and warming)
Alternative Names: GLS4





Observed Molecular Weight:

509.39

Product Description

Morphothiadin is a potent inhibitor on the replication of both wild-type and adefovir-resistant \mathbf{HBV} with an $\mathbf{IC_{50}}$ of 12 nM.

IC50 & Target: IC50: 12 nM (HBV)[1]

In Vitro: Morphothiadin is a potent inhibitor on the replication of both wild-type and adefovir-resistant HBV with an IC₅₀ of 12 nM. Morphothiadin (GLS4) shows no toxicity up to 25 μ M. The cytotoxic dose whereby 50% of cells die (CC₅₀) for primary hepatocytes is 115 μ M for Morphothiadin (P90 is 190 μ M for Morphothiadin (P[2].

In Vivo: The area under the concentration-time curve from 0 to 24 h (AUC₀₋₂₄) of Morphothiadin (GLS4) is 556 h•ng/mL. After intravenous administration of 10 mg/kg Morphothiadin, the total plasma clearance and apparent volume distribution are 4.2 liters/h/kg and 7.38 liters/kg, respectively. The bioavailability of Morphothiadin is 25.5%. It is found that virus titers have increased 83.5-fold in mice treated with 3.75 mg/kg per day of Morphothiadin, 28.3-fold among mice treated with 7.5 mg/kg per day, but only 3- to 6-fold among mice treated with the higher doses of Morphothiadin. There is generally an inverse relationship between Morphothiadin dose and virus titer, with the greatest rebound seen in mice treated with 3.75 mg/kg per day of Morphothiadin (540-fold) and the smallest rebound in mice treated with 60 mg/kg per day (23-fold) (P7.5 mg/kg per day significantly suppresses the virus replication cycle throughout the treatment period, while Morphothiadin doses of >15 mg/kg per day suppresses virus for up to 2 weeks after the end of treatment^[2].

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