
Product Data Sheet

Product Name: PNU-176798

Cat. No.: GC32378

Chemical Properties

Cas. No. 428861-91-0

SMILES CC(NC[C@H]1CN(C2=CC=C(C3=NC=C(C#N)S3)C(F)=C2)C(O1)=O)=OFormula C₁₆H₁₃FN₄O₃S M.Wt 360.36

Solubility Soluble in DMSO Storage Store at -20°C

General tips For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure **Protocol****Kinase experiment:**

Binding of the f[35S]Met-tRNA to E. coli 70S ribosomes is conducted and the initiation reaction mixtures are prepared without initiation factors and contain 6 mM magnesium acetate [Mg(Ac)₂], 0.08 μM AUG, 30 mM NH₄Cl, 10 mM Tris (pH 7.4), and 20 pmol of 70S ribosomes in a final volume of 60 μL. The reaction mixtures are incubated for 15 min at 35°C, and the reactions are terminated by addition of cold buffer A, washed with buffer A, and counted by liquid scintillation[1].

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

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References:

[1]. Aoki H, et al.
Oxazolidinone
antibiotics target
the P site on
Escherichia coli
ribosomes.
Antimicrob Agents
Chemother. 2002
Apr;46(4):1080-5.

Background

PNU-176798 is an antimicrobial agent, targeting protein synthesis in a wide spectrum of gram-positive and anaerobic bacteria.

PNU-176798 is an antimicrobial agent, with a minimum inhibitory concentration (MIC) of 1.4 μM for E. coli. PNU-176798 inhibits fMet-tRNA binding to the 70S ribosomes, with an IC₅₀ of 32 μM . PNU-176798 also blocks translation, 70S initiation with IC₅₀s of 0.53, and 32 μM , respectively. PNU-176798 inhibits peptidyl transferase (IC₅₀, 40 μM), and the inhibition is more pronounced in the presence of elongation factor P (EF-P). PNU-176798 markedly inhibits the EF-G-mediated translocation of fMet-tRNA (IC₅₀, 8 μM)[1].

[1]. Aoki H, et al. Oxazolidinone antibiotics target the P site on Escherichia coli ribosomes. Antimicrob Agents Chemother. 2002 Apr;46(4):1080-5.

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