

## Pyrazofurin

## Chemical Properties

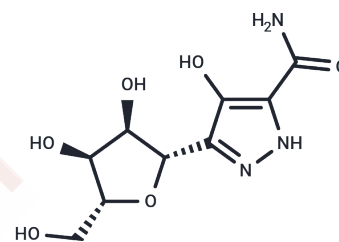
CAS No. : 30868-30-5

Formula: C<sub>9</sub>H<sub>13</sub>N<sub>3</sub>O<sub>6</sub>

Molecular Weight: 259.22

Storage: Keep away from moisture, Store at low temperature  
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	Pyrazofurin is a pyrimidine nucleoside analogue that potently inhibits cell proliferation and DNA synthesis by targeting UMP synthase, while also acting as a highly sensitive orotate-phosphoribosyltransferase inhibitor with IC <sub>50</sub> values ranging from 0.06 to 0.37 μM in Hep-2, HNSCC-14B, and HNSCC-14C cell lines, supporting the use of Pyrazofurin as a mechanistic probe for nucleotide biosynthesis inhibition and antiproliferative research.
Targets(IC <sub>50</sub> )	Proton pump, Dehydrogenase
In vitro	In head and neck cancer cell lines, specifically HEP-2, UMSCC-14B, and UMSCC-14C, Pyrazofurin exhibited potent antiproliferative activity with IC <sub>50</sub> values ranging from 0.06 to 0.37 μM[2]. In Vero cells, Pyrazofurin displayed broad-spectrum antiviral activity against Parainfluenza 3, Measles, Vaccinia, and Herpes Simplex Virus type 2 (HSV-2) with ED <sub>50</sub> values ranging from 2.8 to 20 μM following a 72-hour incubation [3].

## Solubility Information

Solubility	H <sub>2</sub> O: 4 mg/mL (15.43 mM), Sonication is recommended. DMSO: 40 mg/mL (154.31 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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### Preparing Stock Solutions

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	1mg	5mg	10mg
1 mM	3.8577 mL	19.2886 mL	38.5773 mL
5 mM	0.7715 mL	3.8577 mL	7.7155 mL
10 mM	0.3858 mL	1.9289 mL	3.8577 mL
50 mM	0.0772 mL	0.3858 mL	0.7715 mL

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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

- Ringer DP, et al. Alteration in de novo pyrimidine biosynthesis during uridine reversal of pyrazofurin-inhibited DNA synthesis. *Neuropsychopharmacology. J Biochem Toxicol.* 1991 Spring;6(1):19-27.
- Peters GJ, et al. Antiprimidine effects of five different pyrimidine de novo synthesis inhibitors in three head and neck cancer cell lines. *Nucleosides Nucleotides Nucleic Acids.* 2018;37(6):329-339.
- Petrie CR 3rd, et al. Synthesis and biological activity of certain nucleoside and nucleotide derivatives of pyrazofurin. *J Med Chem.* 1986 Feb;29(2):268-78.
- Canonico PG, et al. Antiviral efficacy of pyrazofurin against selected RNA viruses. *Antiviral Res.* 1982 Dec;2(6):331-7.

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