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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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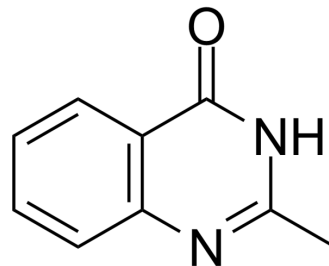
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2-Methylquinazolin-4-ol

Cat. No.:	HY-W051513		
CAS No.:	1769-24-0		
Molecular Formula:	C ₉ H ₈ N ₂ O		
Molecular Weight:	160.17		
Target:	PARP		
Pathway:	Cell Cycle/DNA Damage; Epigenetics		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro

DMSO : 50 mg/mL (312.17 mM; Need ultrasonic)
 H₂O : < 0.1 mg/mL (insoluble)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	6.2434 mL	31.2168 mL	62.4337 mL
5 mM	1.2487 mL	6.2434 mL	12.4867 mL
10 mM	0.6243 mL	3.1217 mL	6.2434 mL

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description

2-Methylquinazolin-4-ol is a potent competitive poly(ADP-ribose) synthetase inhibitor, with a K_i of 1.1 μM. 2-Methylquinazolin-4-ol mammalian aspartate transcarbamylase (ATCase) inhibitor, with 0.20 mM^{[1][2]}.

In Vitro

2-Methylquinazolin-4-ol inhibits ATCase in a concentration-dependent manner^[1].
 MCE has not independently confirmed the accuracy of these methods. They are for reference only.

In Vivo

2-Methylquinazolin-4-ol (0.2-1 mg; i.g.; for two days) inhibits ATCase in a dosedependent manner in vivo^[2].
 MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Mice ^[2]
Dosage:	0.2 mg, 0.4 mg, 0.6 mg, 0.8 mg, 1 mg
Administration:	Oral gavage, for two days

Result:	Shown an inhibitory effect upon ATCase.
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REFERENCES

- [1]. S Yoshida, et al. Production of 2-methyl-4[3H]-quinazolinone, an inhibitor of poly(ADP-ribose) synthetase, by bacterium. J Antibiot (Tokyo). 1991 Jan;44(1):111-2.
- [2]. Mahmoud Balbaa, et al. Inhibition of mammalian aspartate transcarbamylase by quinazolinone derivatives. J Enzyme Inhib Med Chem. 2008 Aug;23(4):483-92.
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Caution: Product has not been fully validated for medical applications. For research use only.

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