Gefitinib impurity 2

Cat. No.:	HY-100663			
CAS No.:	246512-44-7			
Molecular Formula:	C ₁₅ H ₂₃ N ₃ O ₄			
Molecular Weight:	309.36			
Target:	EGFR			
Pathway:	JAK/STAT Signaling; Protein Tyrosine Kinase/RTK			
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 : 100 mg/mL (323.25 mM; Need ultrasonic)						
Preparing Stock Solutions	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg		
		1 mM	3.2325 mL	16.1624 mL	32.3248 mL		
	5 mM	0.6465 mL	3.2325 mL	6.4650 mL			
		10 mM	0.3232 mL	1.6162 mL	3.2325 mL		
	Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (8.08 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (8.08 mM); Clear solution						
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (8.08 mM); Clear solution						

DIOLOGICALACITY	
Description	Gefitinib impurity 2 is the impurity of Gefitinib. Gefitinib (ZD1839; HY-50895) is a potent, selective and orally active EGFR tyrosine kinase inhibitor with an IC ₅₀ of 33 nM. Gefitinib selectively inhibits EGF-stimulated tumor cell growth (IC ₅₀ of 54 nM) and that blocks EGF-stimulated EGFR autophosphorylation in tumor cells. Gefitinib also induces autophagy. Gefitinib has antitumour activity ^{[1][2]} .

REFERENCES

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[1]. Wakeling AE, et al. ZD1839: an orally active inhibitor of epidermal growth factor signaling with potential for cancer therapy. Cancer Res. 2002 Oct 15;62(20):5749-54.

[2]. Pedersen MW, et al. Differential response to gefitinib of cells expressing normal EGFR and the mutant EGFRvIII. Br J Cancer. 2005 Oct 17;93(8):915-23.

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

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