Gefitinib (GMP)

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Cat. No.:	HY-50895G		6
CAS No.:	184475-35-2		
Molecular Formula:	C ₂₂ H ₂₄ ClFN ₄ O ₃		
Molecular Weight:	446.9	N O N	2
Target:	EGFR; Autophagy; Apoptosis	Ó CI ŃH	5
Pathway:	JAK/STAT Signaling; Protein Tyrosine Kinase/RTK; Autophagy; Apoptosis	F	•
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.		

 BIOLOGICAL ACTIVITY

 Description
 Gefitinib (ZD1839) (GMP) is <u>Gefitinib</u> (HY-50895) produced by using GMP guidelines. GMP small molecules work appropriately as an auxiliary reagent for cell therapy manufacture. Gefitinib is a potent, selective and orally active EGFR tyrosine kinase inhibitor^{[1][2]}.

 IC₅₀ & Target
 IC50: 37 nM (Tyr1173 site, in NR6wtEGFR cells), 37 nM (Tyr992 site, in NR6wtEGFR cells)^[1].

 In Vitro
 Gefitinib (GMP) abolishes the effect of EGF-induced dedifferentiation of astrocytes into astrocyte precursor cells (APCs)^[2]. Gefitinib (3 μM) can produce a subgroup of EGFR-mutant NSCLC cell lines (Gefitinib-resistant cells) that undergo cellular reprogramming, such as HCC827 cells^[3]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[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]. Liu X, Li C, et al. EGF signaling promotes the lineage conversion of astrocytes into oligodendrocytes. Mol Med. 2022 May 4;28(1):50.

[3]. Ware KE, et al. A mechanism of resistance to gefitinib mediated by cellular reprogramming and the acquisition of an FGF2-FGFR1 autocrine growth loop. Oncogenesis. 2013 Mar 25;2(3):e39.

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

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Product Data Sheet



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