BIBW-2992

REAGENTS DIRECT

5 mg

For research purposes only

BIBW-2992 is a irreversible and dual inhibitor of epidermal growth factor receptor 1 (EGFR) and 2 (HER2) kinases. The potency of BIBW-2992 on the EGFR and HER2 kinases revealed IC₅₀ values of 0.5 nM and 14 nM, respectively. BIBW-2992 is highly selective for these kinases and no additional inhibition of other kinases has been observed. BIBW-2992 has also been shown to suppress EGF-induced EGFR phosphorylation and cellular proliferation in various cell lines.

TECHNICAL INFORMATION

H₃C N N N HN N HN CI

Other Names: Afatinib

Chemical Formula: C₂₄H₂₅CIFN₅O₃

CAS Number: 850140-72-6

Molecular Weight: 485.94

Purity: >98%

Appearance: a crystalline solid

Solubility: DMSO

STORAGE AND HANDLING

Storage: Store at 4°C and protected from light. Following

reconstitution, store aliquots at -20°C.

Stability: Stock solutions stable at -20°C for up to 2 years.

Shipping Conditions: Shipped at room temperature.

PRODUCT USE

Soluble in DMSO at 200 mg/ml. Soluble in ethanol at 25mg/ml. If precipitate is observed, vortex for 5 minutes. For most cells, the maximum tolerance to DMSO is less than 0.5%.

REFERENCES

- 1. Li et al. (2008) BIBW2992, an irreversible EGFR/HER2 inhibitor highly effective in preclinical lung cancer models. Oncogene. 27(34):4702-11.
- Takezawa et al. (2010) Enhanced anticancer effect of the combination of BIBW2992 and thymidylate synthase-targeted agents in non-small cell lung cancer with the T790M mutation of epidermal growth factor receptor. Mol Cancer Ther. 9(6):1647-56.
- 3. Minkovsky et al. (2008) BIBW-2992, a dual receptor tyrosine kinase inhibitor for the treatment of solid tumors. Curr Opin Investig Drugs. 9(12): 1336-46.

