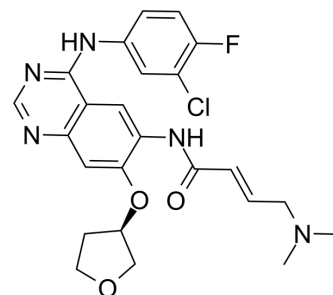


## (R)-Afatinib

|                    |   |
|--------------------|---|
| Cat. No.:          | HY-10261E   |
| CAS No.:           | 439081-17-1   |
| Molecular Formula: | C <sub>24</sub> H <sub>25</sub> ClFN <sub>5</sub> O <sub>3</sub>                          |
| Molecular Weight:  | 485.94  |
| Target:            | EGFR; c-Met/HGFR; p38 MAPK  |
| Pathway:           | JAK/STAT Signaling; Protein Tyrosine Kinase/RTK; MAPK/ERK Pathway                         |
| Storage:           | Please store the product under the recommended conditions in the Certificate of Analysis. |



### BIOLOGICAL ACTIVITY

#### Description

(R)-Afatinib ((R)-BIBW 2992) is the Afatinib isomer. [Afatinib](#) (HY-10261) is an orally active, potent and irreversible dual specificity inhibitor of ErbB family (EGFR and HER2), with IC<sub>50</sub> values of 0.5 nM, 0.4 nM, 10 nM and 14 nM for EGFR<sup>wt</sup>, EGFR<sup>L858R</sup>, EGFR<sup>L858R/T790M</sup> and HER2, respectively. Afatinib can be used for the research of esophageal squamous cell carcinoma (ESCC), non-small cell lung cancer (NSCLC) and gastric cancer<sup>[1]</sup>.

### REFERENCES

[1]. Li D, et al. BIBW2992, an irreversible EGFR/HER2 inhibitor highly effective in preclinical lung cancer models. *Oncogene*. 2008 Aug 7;27(34):4702-11.

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

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