PRODUCT INFORMATION



Alflutinib (mesylate)

Item No. 31473

CAS Registry No.: 2130958-55-1

Formal Name: N-[2-[[2-(dimethylamino)ethyl]

methylamino]-5-[[4-(1-methyl-1H-indol-3-yl)-2-pyrimidinyl] amino]-6-(2,2,2-trifluoroethoxy)-3-pyridinyl]-2-propenamide, monomethanesulfonate

Synonym:

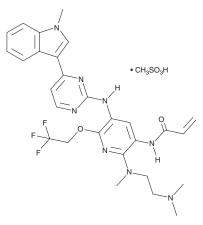
MF: $C_{28}H_{31}F_3N_8O_2 \bullet CH_3SO_3H$

FW: **Purity:**

UV/Vis.: λ_{max} : 267, 332 nm A crystalline solid Supplied as:

-20°C Storage: Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.



Laboratory Procedures

Alflutinib (mesylate) is supplied as a crystalline solid. A stock solution may be made by dissolving the alflutinib (mesylate) in the solvent of choice, which should be purged with an inert gas. Alflutinib (mesylate) is soluble in the organic solvent chloroform at a concentration of approximately 10 mg/ml. Alflutinib (mesylate) is also slightly soluble in DMSO and dimethyl formamide.

Description

Aflutinib is an inhibitor of mutant EGFRs.¹ It is selective for EGFR^{L858R}, EGFR^{G719X}, EGFR^{L858R}. EGFR^{L861A}. and EGFR^{T790M} mutant EGFRs over wild-type EGFR. Aflutinib (10 and 30 mg/kg) reduces tumor growth in an EGFR^{L858R} and EGFR^{T790M}-expressing LU1868 non-small cell lung cancer (NSCLC) patient-derived xenograft (PDX) mouse model.

Reference

1. Shi, Y., Zhang, S., Hu, X., et al. Safety, clinical activity, and pharmacokinetics of alflutinib (AST2818) in patients with advanced NSCLC with EGFR T790M mutation. J. Thorac. Oncol. 15(6), 1015-1026 (2020).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

WARRANTY AND LIMITATION OF REMEDY

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