PRODUCT INFORMATION



GSK-F1

Item No. 41366

CAS Registry No.: 1402345-92-9

Formal Name: 5-[2-amino-3,4-dihydro-4-oxo-3-[2-

(trifluoromethyl)phenyl]-6-quinazolinyl]-N-(2,4-

difluorophenyl)-2-methoxy-3-pyridinesulfonamide

MF: $C_{27}H_{18}F_5N_5O_4S$

603.5 FW: ≥98% **Purity:** Supplied as: A solid Storage: -20°C Stability: ≥4 years

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

Laboratory Procedures

GSK-F1 is supplied as a solid. A stock solution may be made by dissolving the GSK-F1 in the solvent of choice, which should be purged with an inert gas. GSK-F1 is soluble (≥10 mg/ml) in DMSO and slightly soluble (0.1-1 mg/ml) in ethanol.

Description

GSK-F1 is a PI4KIII α inhibitor (IC $_{50}$ = 5.01 nM). 1 It is selective for PI4KIII α over PI4K β , PI3K α , PI3K β , PI3K δ , and PI4K γ (IC $_{50}$ s = 1,000, 2,512, 7,943, 2,512, and 2,512 nM, respectively). GSK-F1 inhibits the replication of hepatitis C virus (HCV) genotypes 1a, 1b, and 2a in a cell-based replicon assay (IC_{50} s = 12.6, 2.51, and 12.6 nM, respectively).

Reference

1. Leivers, A.L., Tallant, M., Shotwell, J.B., et al. Discovery of selective small molecule type III phosphatidylinositol 4-kinase alpha (PI4KIIIα) inhibitors as anti hepatitis C (HCV) agents. J. Med. Chem. 57(5), 2091-2106 (2014).

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.

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