Product Specification Sheet

Product Name: GDC-0980 (RG7422)

Catalog Number: C4098

Technical information:

Chemical Formula: C₂₃H₃₀N₈O₃S

CAS #: 1032754-93-0

Molecular Weight: 498.6

Purity: > 98%

Appearance: White solid

Solubility: Soluble in DMSO up to 40 mM

Chemical Name: (S)-1-(4-((2-(2-aminopyrimidin-5-yl)-7-methyl-4-morpholinothieno[3,2-d]pyrimidin-6-

yl)methyl)piperazin-1-yl)-2-hydroxypropan-1-one

Storage: Store solid powder at 4°C desiccated; Store DMSO solution at -20°C.

Shelf Life: In the unopened package, powder is stable for 1 year and DMSO solution is stable for 6 months

under proper storage condition.

Handling: • To make 10 mM stock solution, add 0.201mL of DMSO for each mg of GDC-0980 (RG7422).

• For DMSO solution, briefly spin the vial at 500 rpm in a 50 mL conical tube to ensure maximum

FK866 (APO866)

sample recovery.

Biological Activity: GDC0980, a orally-available, thienopyrimidine-based, ATP-competitive dual inhibitor of PI3K class I

isoforms (IC50 of 5, 27, 7, and 14 nM for PI3Ka, b, d, and g) and mTOR (Ki of 17 nM). [1] GDC0980 potently inhibits signal transduction (Thr308, Ser473) downstream of both PI3K and mTOR and showed most potent antitumor activity in breast, prostate, and lung cancer (IC50 < 500 nM) cells.

Treatment of cancer cells with GDC0980 resulted in G1 cell-cycle arrest. [2]

Reference: 1. Sutherlin et al., Discovery of a Potent, Selective, and Orally Available Class I Phosphatidylinositol 3-Kinase

(PI3K)/Mammalian Target of Rapamycin (mTOR) Kinase Inhibitor (GDC-0980) for the Treatment of Cancer. J.

Med. Chem. 2011, 54, 7579-7587. Pubmed ID: 21981714

2. Wallin et al., GDC-0980 Is a Novel Class I PI3K/mTOR Kinase Inhibitor with

Robust Activity in Cancer Models Driven by the PI3K Pathway, Mol. Cancer Ther. 2011, 10, 2426-2436.

Pubmed ID: 21998291

To reorder: http://www.cellagentech.com/GDC-0980-RG7422/

For Technical Support: technical@cellagentech.com

Chemicals are sold for research use only, not for clinical or diagnostic use.