## **Product Sheet**

## GDC-0941, PI3 Kinase Inhibitor

G941-005; G941-025; G941-100 Catalog #

Chemical Name: 2-(1H-Indazol-4-yl)-6-[[4-(methylsulfonyl)-1-piperazinyl]methyl]-4-Description

(4-morpholinyl)-thieno[3,2-d]pyrimidine

GDC-0941 is a potent inhibitor of PI3Kα, PI3Kβ, PI3Kδ and PI3Kγ with IC<sub>50</sub> of 3 nM, 33 nM, 3 nM and 75 nM, respectively. Administration of GDC-0941 in animals displays significant inhibitory effect against established human cancer xenografts. It is currently in clinical trials.

PI3K inhibitors have been used for differentiation of pluripotent stem cells.

Folkes AJ, et al. The identification of 2-(1H-indazol-4-yl)-6-(4-Ref: methanesulfonyl-piperazin-1-ylmethyl)-4-morpholin-4-yl-thieno[3,2-d]pyrimidine (GDC-0941) as a potent, selective, orally bioavailable inhibitor of class I PI3 kinase for the treatment of cancer. J Med Chem 51(18):5522-32, 2008

Touboul, et al. Generation of functional hepatocytes from human embryonic stem cells under chemically defined conditions that recapitulate liver development. Hepatology, 51: 1754–1765, 2010

Powder **Formulation** 

Reconstitution Before reconstitution, we recommend a brief spin to drive down any

> material dislodged from the bottom of the tube. The compound is soluble in DMSO. Poorly Soluble in water. Dilution into protein-containing

aqueous media is recommended.

**Stability** The powder is stable for at least 2 year if stored at -20 degree C. The

> dissolved compound is stable for at least 1 month at 4 degree C, but should be stored in aliquots at -20 degree C for longer term. Protect from light.

Greater than 99% as determined by LC/MS analysis. LC/MS and/or NMR Purity

data available upon request.

**Biological Activity** Not determined.

Structural Info

MW: 513.64

Formula: C23H27N7O3S2

**Solubility:** Soluble in DMSO up to 50 mM. Poorly soluble in water.

CAS: 957054-30-7

For Research Use Only. Not for Use in Humans.

