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Product Information

Product ID A9601

CAS No. 1143532-39-1

Chemical Name

Synonym AZD5363

Formula C₂₁H₂₅ClN₆O₂

Formula Wt. 428.92

Melting Point

Purity ≥99%, ≥99%ee

Solubility In DMSO(86mg/ml), water

with 2 eq. of acid, insoulble

on ethanol

OH

Bulk quanitites available upon request

Product ID Size A9601 1 mg A9601 5 mg A9601 25 mg

Store Temp -20°C Ship Temp Ambient

Description AZD-5363 is an oral pan-AKT inhibitor that causes hyperphosphorylation of AKT, locking it in a catalytically inactive state,

unable to phosphorylate downstream signaling substrates such as PRAS40 and GSK-8. This compound displays anticancer chemotherapeutic activity, inhibiting proliferation and inducing tumor regression in in vitro and in vivo models of HER2+ breast cancer and prostate cancer. AZD-5363 may also exhibit inhibitory activity against Rho-associated protein kinase (ROCK),

p70S6K, PKA, MKK1, MSK1, MSK2, PKC, PKG, PRKX, and RSK2/3.

References Addie M, Ballard P, Buttar D, et al. Discovery of 4-amino-N-[(1S)-1-(4-chlorophenyl)-3-hydroxypropyl]-1-(7H-pyrrolo[2,3-d]pyrimidin -4-yl)piperidine-4-carboxamide (AZD5363), an orally bioavailable, potent inhibitor of Akt kinases. J Med Chem. 2013 Mar 14:56 (5):2059-73. PMID: 23394218.

> Lamoureux F, Thomas C, Crafter C, et al. Blocked autophagy using lysosomotropic agents sensitizes resistant prostate tumor cells to the novel Akt inhibitor AZD5363. Clin Cancer Res. 2013 Feb 15;19(4):833-44. PMID: 23258740.

> Davies BR, Greenwood H, Dudley P, et al. Preclinical pharmacology of AZD5363, an inhibitor of AKT: pharmacodynamics, antitumor activity, and correlation of monotherapy activity with genetic background. Mol Cancer Ther. 2012 Apr;11(4):873-87. PMID: 22294718.

Caution: This product is intended for laboratory and research use only. It is not for human or drug use.