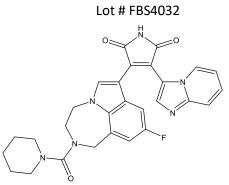


Catalog #10-4854 LY2090314

CAS# 603288-22-8

3-[6-Fluoro-10-(piperidine-1-carbonyl)-1,10-diazatricyclo[6.4.1.0^{4,13}]trideca-2,4,6,8(13)-tetraene-3-yl]-4-imidazo[1,2-a]pyridine-3-ylpyrrole-2,5-dione; 3-(9-Fluoro-2-(piperidine-1-carbonyl)-1,2,3,4-tetrahydro-[1,4]diazepino[6.7.1-hi]indol-7-yl)-4-(imidazo[1.2-a]pyridine-3-yl)-1H-pyrrole-2,5-dione



Wnt/ß-catenin pathway activator. LY2090314 is a highly potent ($IC_{50} = 1.5 \text{ nM GSK-3}\alpha$; 0.9 nM GSK-3ß) and selective inhibitor of glycogen synthase-3-kinase (GSK-3).^{1,2} It stabilized ß-catenin levels and elevated expression of Axin2 leading to tumor growth delays in A375 melanoma xenografts.2 LY2090314 significantly reduced growth in multiple neuroblastoma cell lines.³ It inhibited growth and induced apoptosis in osimertinibresistant lung cancer cells, specifically those with a mesenchymal phenotype.⁴

- 1) Engler et al. (2004), Substituted 3-imidazo[1,2-a]pyridine-3-yl-4-(1,2,3,4-tetrahydro-[1,4]diazepino-[6,7,1=hi]indol-7-yl)pyrrole-2,5-diones as and potent inhibitors of glycogen synthase-3-kinase; J. Med. Chem. **47** 3934
- 2) Atkinson et al. (2015), Activating the Wnt/ß-Catenin Pathway for the Treatment of Melanoma Application of LY2090314, a Novel Selective Inhibitor of Glycogen Synthase Kinase-3; PLoS One **10** e0125028
- 3) Kunnimalaiyaan et al. (2018), Antiproliferative and apoptotic effect of LY2090314, a GSK-3 inhibitor, in neuroblastoma in vitro BMC Cancer 18 560
- 4) Fukuda et al. (2020), Glycogen synthase kinase-3 inhibition overcomes epithelial-mesenchymal transitionassociated resistance to Osimertinib in EGFR-mutant lung cancer, Cancer Sci. **111** 2374

PHYSICAL DATA

Molecular Weight:	512.54
Molecular Formula:	C ₂₈ H ₂₅ FN ₆ O ₃
Purity:	>98% (HPLC)
	NMR: (Conforms)
Solubility:	DMSO (>20 mg/mL)
Physical Description:	Orange solid
Storage and Stability:	Store as supplied at -20°C for up to 2 years from the date of purchase. Solutions in
	DMSO may be stored at -20°C for up to 3 months.

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