

LY294002 2-(4-Morpholinyl)-8-phenyl-4H-1benzopyran-4-one PRODUCT ANALYSIS SHEET

Catalog Number:	PHZ1144
Lot Number:	See product label
Quantity:	5.0 mg
Appearance:	Off-white to pale yellow solid
Molecular Formula:	C ₁₉ H ₁₇ NO ₃
Molecular Weight:	307.4
Purity:	99%, as assessed by TLC
Summary:	LY294002 is a potent and specific cell-permeable inhibitor of phosphatidylinositol 3-kinase (PI3-kinase). LY294002 at a concentration of 50 μ M completely abolishes the PI3-K activity of intact neutrophils stimulated with fMet-Leu-Phe, without apparent cell toxicity. While a concentration or 50 μ M is observed to be efficacious for inhibiting PI3-K activity in stimulated neutrophils, this concentration produces no significant inhibition of other kinases including PKC, PKA, MAP kinase, S6 kinase, EGF receptor tyrosine kinase, Src, PI4-kinase, diacylglycerol kinase, or rabbit kidney ATPase. This compound is a useful tool for identifying cellular events that are regulated by the PI3-kinase/Akt axis, and is observed to induce apoptosis in many cell types by blocking the PI3-kinase/Akt anti-apoptotic pathway.
Biological Activity:	$IC_{50} = 1.4 \ \mu M$
Solubility:	Soluble in warm DMSO at a concentration of 25 mg/mL; soluble in warm ethanol at a concentration of 25 mg/mL.
Sterility:	This product is not sterile.
Storage:	Store, as supplied, at -20° C. Upon solubilization, apportion into working aliquots and store at -20° C. Avoid repeated freeze/thaw cycles. Solutions are stable at -20° C for up to three months.
Expiration Date:	Expires one year from date of receipt when stored as instructed.
Related Products:	Akt/PKB [pS ⁴⁷³] antibody, Cat. # 44-622
References:	Vlahos, C.J., et al. (1994) A specific inhibitor of phosphatidylinositol 3-kinase, 2-(4-morpholinyl)-8-phenyl-4H-1- benzopyran-4-one (LY294002). J. Biol. Chem. 269(7):5241-5248.
	Sanchez-Margalet, V., et al. (1994) Role of phosphatidylinositol 3-kinase in insulin receptor signaling: studies with inhibitor, LY294002. Biochem. Biophys. Res. Commun. 204(2):446-452.
	Bancroft, C.C., et al. (2002) Effects of pharmacologic antagonists of epidermal growth factor receptor, P13K and MEK signal kinases on NF-kappaB and AP-1 activation and IL-8 and VEGF expression in human head and neck squamous cell carcinoma lines. Int. J. Cancer 99(4):538-548.
	Williamson, R., et al. (2002) Rapid tyrosisne phosphorylation of neuronal proteins including tau and focal adhesion kinase in response to amyloid-beta peptide exposure: involvement of Src family protein kinases. J. Neurosci. 22(1):10-20.
	Shoba, L.N., et al. (2001) LY294002, an inhibitor of phosphatidylinositol 3-kinase, inhibits GH-mediated expression of the IGF-I gene in rat hepatocoytes. Endocrinology 142(9):3980-3986.
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Caution:

Avoid contact with eyes, skin, and mucous membranes. Wear protective clothing when handling this product. Not for human use.

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