

Table 1S

Inhibitor	IC₅₀ (kinase assay)	IC₅₀ in cells/tissue	Citation
CAL-101	p110 α = 820 nM	pAKT following PDGF stimulation EC ₂₅ = 10 μ M; pAKT following LPA stimulation EC ₅₀ = 1.9 μ M in fibroblasts	Lannutti BJ <i>et al.</i> (2011) Blood, 117:591-594.
	p110 β = 565 nM		
	p110 γ = 2.5 nM		
	p110 δ = 89 nM		
LY294002	PI3K = 1.4 μ M	BrdU uptake in aortic smooth muscle = 32 μ M	Vlahos CJ <i>et al.</i> (1994) JBC, 269:5241-5248.
Y27632	ROCK1 Ki = 0.22 μ M	G1-S phase cell cycle progression in 3T3 cells = between 10 and 100 μ M	Ishizaki <i>et al.</i> (2000) Mol Pharmacol, 57:976-983.
	ROCK2 Ki = 0.30 μ M		