Table S2. Pharmacological compounds used in this study

Drug Name	Function	Concentration	References
Cytochalasin D (CytD)	Inhibitor of actin polymerization (disruption of actin microfilaments and activation of p53-dependent pathways which results in the arrest of the cell cycle at the G1-S transition)	2.5 μg/ml	[1]
LY294002	PI3 kinase inhibitor (blocks PI3 kinase-dependent Akt phosphorylation and kinase activity)	30 μM	[2]
3-Methyladenine (3-MA)	Autophagy Inhibitor; PI3 kinase inhibitor	10 mM	[3]
Bafilomycin A1 (BAF)	Inhibitor of the vacuolar type H(+)-ATPase and endosomal acidification	100 nM	[4]

References

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