Molecular Pharmacology

Obatoclax and Lapatinib interact to induce toxic autophagy through NOXA

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Supplemental Figures S1-S3. Expression / phosphorylation of proteins whose activity or expression has been targeted. The panels show proteins whose expression/phosphorylation has been: (Figure S1) knocked down by siRNA (ATM, Beclin1, ATG5); (Figure S2) where dominant negative p38 blocks HSP27 phosphorylation and dominant negative MEK1 blocks ERK1/2 phosphorylation; (Figure S3) where ca-p70 and ca-mTOR both increase the levels of P-S6 and maintain S6 phosphorylation after lapatinib + obatoclax treatment.



	BT474	MCF7	
siSCR	+	+	
siBeclin-1	+	+	
Beclin-1	-	1000	
β-actin			

	MCF7	BT474
siSCR	+	+
siATM	+	+
ATM		Print party
β-actin		



Figure S2



		MCF7	
Figure S3		CMV	ca-p70
	Vehicle	+	+
	GX+LAP	+	+
	P-p70	100.000	-
	P-S6	-	-
	β-actin	-	

