

Table S1. IC₅₀ values of OVCAR3 cells with different p53 status to EGFR and/or MDM2 inhibitors.

Cell Line	Cell Status	Drug Treatment	IC ₅₀ (μM)
OVCAR3	Pre-transfected with empty plasmid for 18 hours	Gefitinib	33.74
		JNJ	21.83
		Gefitinib + JNJ	8.57
	Pre-transfected with p53 ^{R248Q} for 18 hours	Gefitinib	26.47
		JNJ	14.35
		Gefitinib + JNJ	8.18

Table S2. Combination index (CI) values and synergism grading of combined gefitinib and JNJ treatment at different effective doses (ED₅₀, ED₇₅ and ED₉₀).

CI values of specified effective dose (ED) for OVCAR3 cells with different p53 statuses were summarized. The synergism grading of combined inhibition was based on the grading standard defined by Chou and Talalay [39]. CI values from 0.85 to 0.9, 0.7 to 0.85 and 0.3 to 0.7 indicated slight synergism, moderate synergism and synergism, expressed as "+", "++" and "+++", respectively.

Cell Line	Cell Status	CI Values and Synergism Grading		
		ED ₅₀	ED ₇₅	ED ₉₀
OVCAR3	Pre-transfected with			
	empty plasmid for 18	0.65	0.51	0.47
	hours	+++	+++	+++
	Pre-transfected with	0.87	0.72	0.67
	p53 ^{R248Q} for 18 hours	+	++	+++

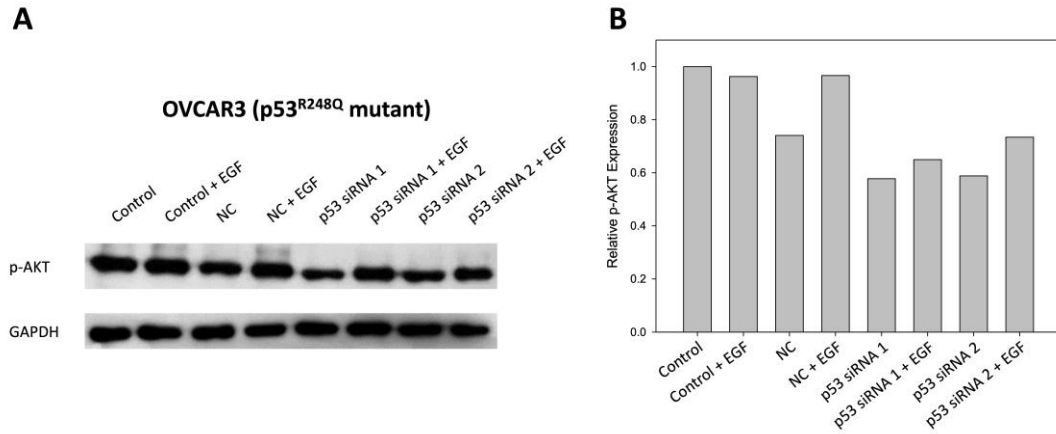


Figure S1. p53^{R248Q} resulted in regulation of p-AKT.

(A) The level of p-AKT was reduced after transfection with p53 siRNA in OVCAR3 cells. It should be noted that serum-starved OVCAR3 cells were transiently transfected with negative control siRNA (25 nM) or p53 siRNA (25 nM) for 24 hours and then stimulated with EGF for 15 minutes. GAPDH expression was measured to serve as a loading control.

(B) The band intensity of the immunoblots was quantified using ImageJ software. The relative intensities of p-AKT expression were normalized with the control group.

(NC: negative control siRNA)

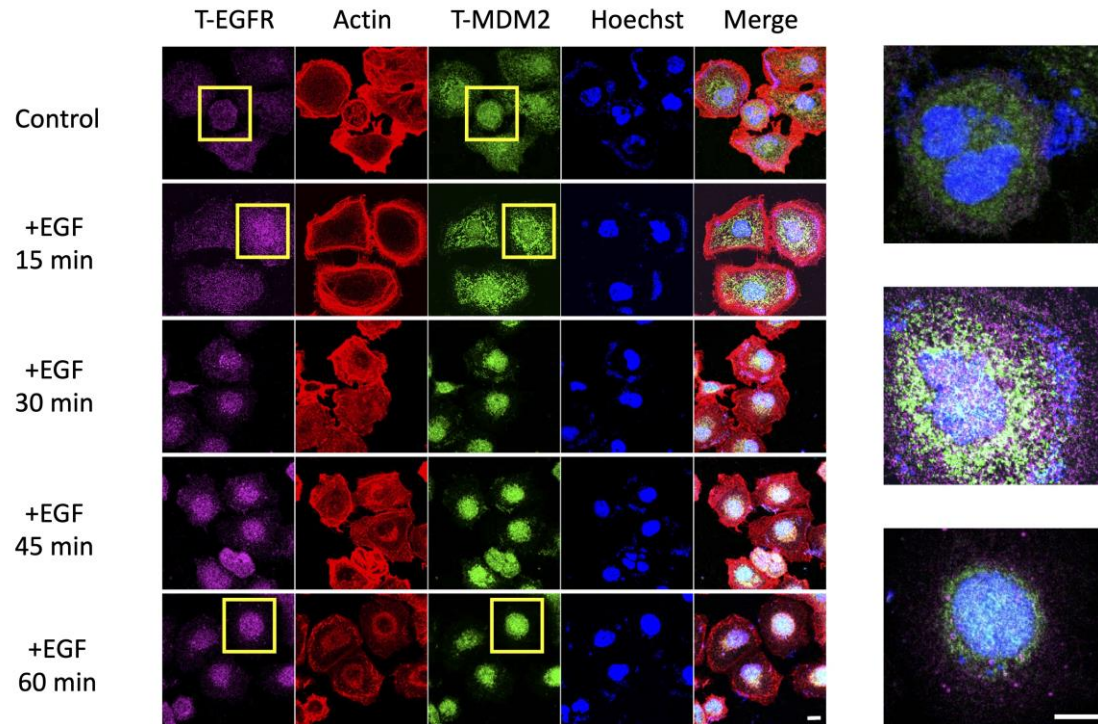


Figure S2. Time-dependent cytonuclear trafficking patterns of EGFR and MDM2. EGFR and MDM2 moved in a convergent manner from the cytoplasm to the nucleus over time after EGF stimulation. The right images were enlarged view of the yellow boxed region. The scale bar represented 10 μ m.

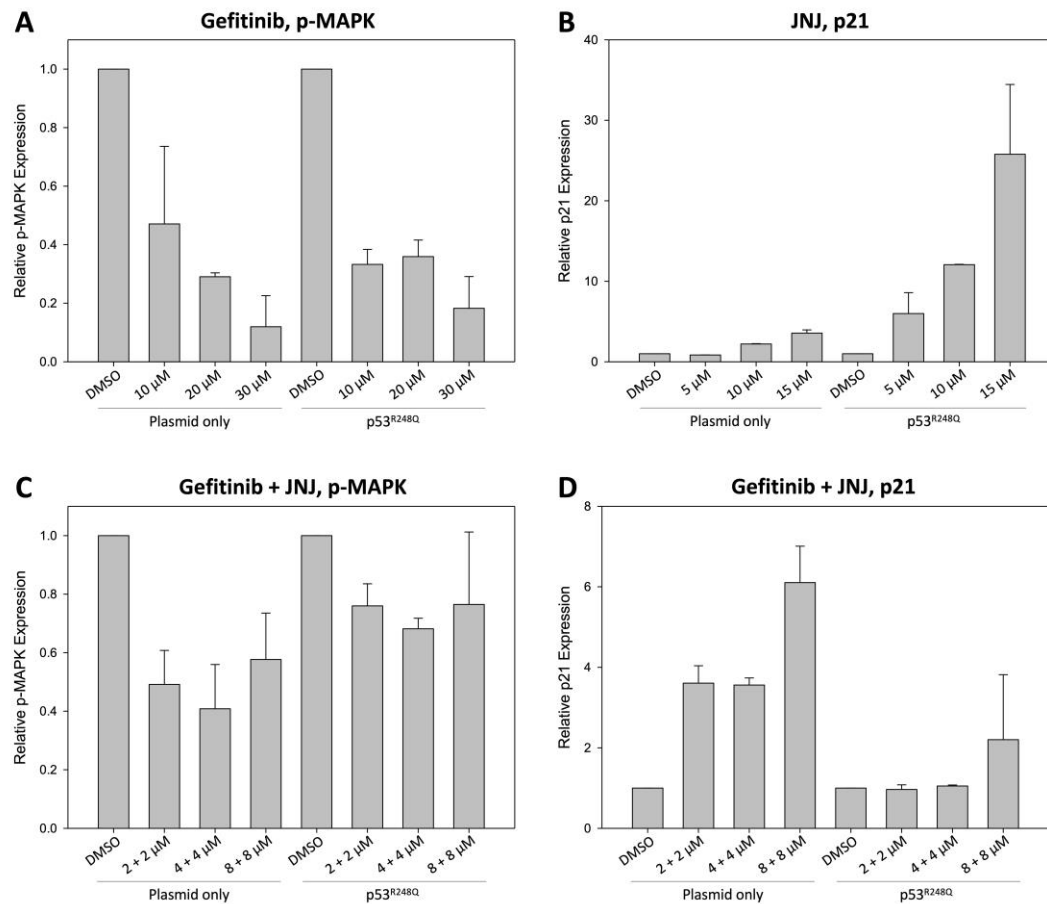


Figure S3. Quantification graphs of immunoblots in Figure 6.

The band intensity of the immunoblots was quantified using ImageJ software. The relative intensities of p-MAPK or p21 expression were normalized with the DMSO control group.