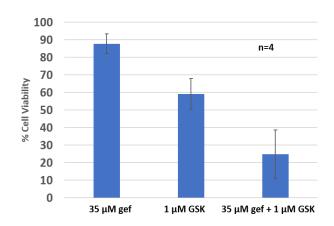
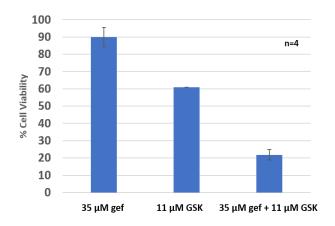
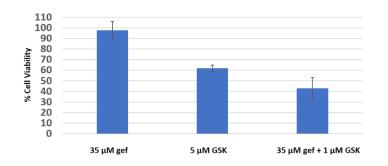
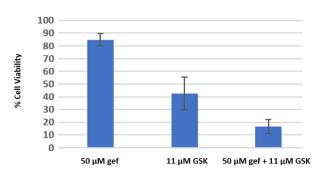
Supplementary Materials:



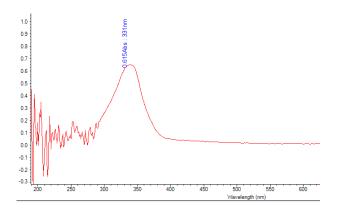




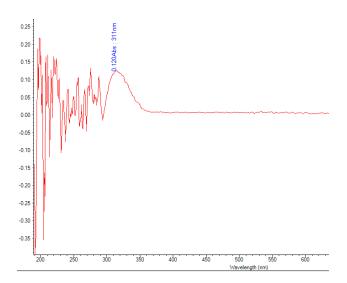


Supplementary Figure S1: Evaluation of cell viability post treatment with different sets of combination of free gefitinib and free GSK461364A

(2A)



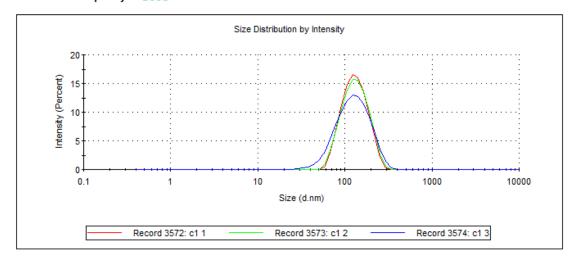
(2B)



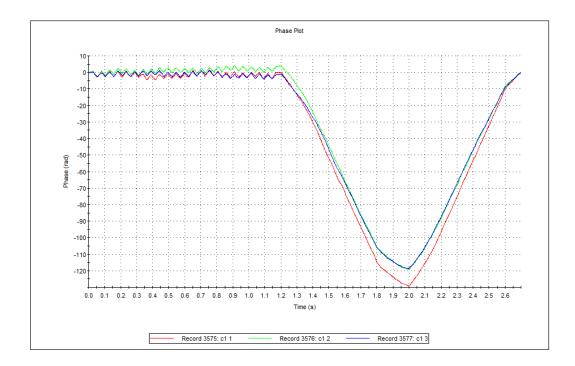
Supplementary Figure S2: Peak absorbance of **(2A)** gefitinib at 331nm dissolved in 1% Triton-x, with GSK461364A as blank **(2B)** GSK461364A at 311nm with gefitinib as blank

(3A)

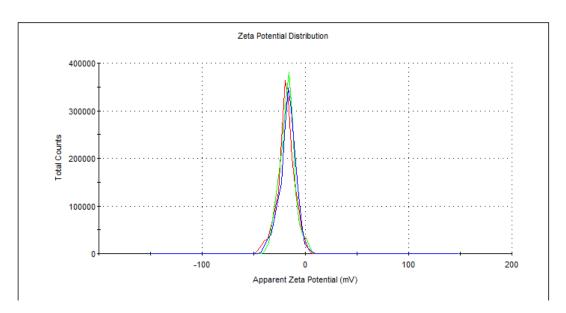
			Size (a.iiii):	70 intensity.	St Dev (a.iiii):
Z-Average (d.nm):	122.1	Peak 1:	135.5	100.0	43.99
Pdl:	0.091	Peak 2:	0.000	0.0	0.000
Intercept:	0.966	Peak 3:	0.000	0.0	0.000
Result quality :	Good				



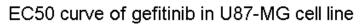
(3B)

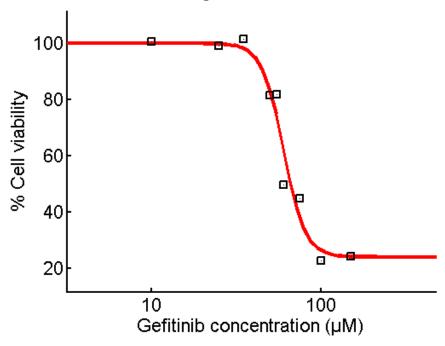


(3C)



Supplementary Figure S3: (A) Size distribution by intensity with the average size being \sim 122 nm. (B) Phase plot of zeta potential plotted against time (C) Zeta potential distribution with total counts plotted against apparent zeta potential. Average zeta potential was \sim 20 mV.





Supplementary Figure S4: Effective concentration (EC₅₀) curve of gefitinib in U87-MG cell line plotted using Dr.Fit[®] software. These data points follow monophasic model (standard Hill equation).