## Supplemental Material to:

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**Figure S1.** D261 inhibits MK production in vitro. NCI-H460 or A549 cells were treated with various concentrations of D261. (**A**) MK RNA levels (D261 treated for 24 h) were detected by qRT-PCR. Data are shown as mean  $\pm$  SEM (n = 3) of one representative experiment. Similar results were obtained in at least three independent experiments. \*P < 0.05, \*\*\*P < 0.001 compared to DMSO control. (**B**) MK protein levels (D261 treated for 48 h) were detected by western blot.



**Figure S2.** Effects of D261 on cell apoptosis. (**A**) NCI-H460 or A549 cells were treated with various concentrations of D261 for 48 h, cell apoptosis was determined by Annexin V-FITC/PI staining. (**B**) Data are shown as mean  $\pm$  SEM of three independent experiments. \**P* < 0.05, \*\**P* < 0.01, \*\*\**P* < 0.001 compared to each DMSO control.



**Figure S3.** Overexpression of activated mTOR rescued D261-induced inhibition of cell growth. (**A**) A549 cells were transfected with pcDNA3-Flag mTOR or vehicle plasmid pcDNA3 for 72 h. Expression of Flag was detected by western blot. Tubulin was performed as a loading control. (**B**) A549 cells were transfected with pcDNA3-Flag mTOR or vehicle plasmid pcDNA3 24 h before treated with D261 (4  $\mu$ M) for another 48 h. Expression of mTOR was detected by western blot. Cell viability was assessed using CCK-8 assay. The bars indicate mean ± SEM (n = 5). \*P < 0.05, \*\*\*P < 0.001.