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Supplementary Information



Figure. S1 Cell growth rate of (A) HCT116, (B) RKO, (C) SW480 and (D) HT29 was determined by IncuCyteTM Zoom live cell imaging system after indicative dose of α -MG treatment ($\mu g/ml$).

Figure. S2





Figure. S2 Clonogenic assay of (A) RKO, (B) SW480 and (C) HT29 cells after indicative dose of α -MG ($\mu g/ml$) was determined one week after treatment (left panel). The quantification of number of colony was presented as a graph (right panel).

Figure. S3



Figure. S3 Flow cytometry analysis of (A) RKO, (B) SW480 and (C) HT29 cells treated with $5\mu g/mL$ of α -MG at the indicated times was shown (left panel). Cell cycle distribution was calculated as the percentage of cells in G1, S and G2/M phase and was graphically presented as a bar graph (right panel).

Figure. S4



Figure. S4 Total cell lysates of HCT116WT cells after 5µg/ml of α -MG treatment was harvested at the indicated time. Protein expressions of phospho-p38 (pp38), phospho-ERK (pERK), and phospho-JNK (pJNK) MAPK and α/β -tubulin (control) were determined by immunoblotting analysis.