

FIGURE S1: (a) HepG2 and LO2 cells were treated with different concentration of H_2O_2 for 3h, cell viability was measured by CCK8 assay. (b) Level of GSH in HepG2 cells after treatment with various concentrations of OA. (c, d, e) Relative TNF- α , IL-1 β and IL-6 expression of HepG2 cells treated with various concentrations of OA. Values are expressed as the mean \pm s d from three independent experiments. **P<0.01, *P<0.05. Bars indicate the standard deviation of the mean.



FIGURE S2: H_2O_2 induced the Phosphorylation of the AKT/mTOR pathway. (a) Western blot analysis of (p)PI3K/(p)AKT/(p)mTOR protein levels with or without H_2O_2 treatment. (b, c, d) Densitometry analysis of (p)PI3K/(p)AKT/(p)mTOR protein levels. **P<0.01, *P<0.05, compared with control group. Data are plotted as the mean \pm s d from three independent experiments. Bars indicate the standard deviation of the mean.



FIGURE S3: (a, b) Relative PA and OA content of mouse liver after surgical procedure. (c, d) Relative PA and OA content of HepG2 cells after treated with OA. **P<0.01, *P<0.05, compared with control group. Data are plotted as the mean \pm s d from three independent experiments. Bars indicate the standard deviation of the mean.



FIGURE S4: (a) Western blot analysis of (p)PI3K and LC3- || protein levels after oxidative stress in HepG2 cells with or without OA treatment. Values are expressed as the mean \pm s d from three independent experiments. **P<0.01, *P<0.05. Bars indicate the standard deviation of the mean.

