Electronic Supplementary Material (ESI) for Food & Function. This journal is © The Royal Society of Chemistry 2021

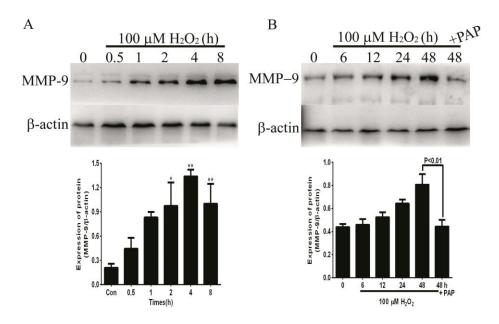


Figure S1 *Effect of H*<sub>2</sub> $O_2$  *on MMP-9 activation* in NCI-H1650 cells. (A) H<sub>2</sub> $O_2$  induced MMP-9 expression in a time-dependent manner. (B) Western blot analyses showed that 100  $\mu$ M H<sub>2</sub> $O_2$  induced MMP-9 secretion in a time-dependent manner, and pre-treatment of 400  $\mu$ g/mL PAP decreased MMP-9 protein. \*P<0.05, \*\*P<0.01 vs. Con.

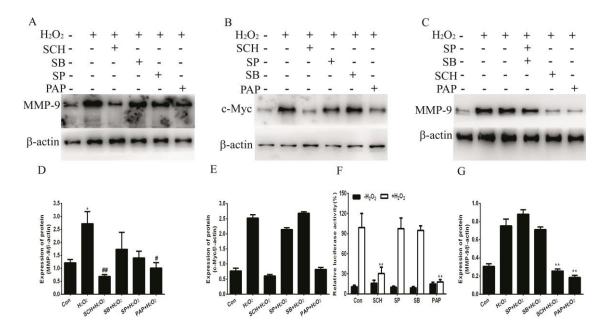


Figure S2 Regulation of MMP-9 transcriptions are dependent on ERK/c-Myc activities in NCI-H1650 cells. (A)  $H_2O_2$ -induced MMP-9 expression was associated with increased ERK signaling in NCI-H1650 cells. The results represent as mean  $\pm$  SD, and \*P<0.05 vs. Con; #P<0.05, ##P<0.01, vs.  $H_2O_2$  group. (B, E) ERK inhibitor SCH772984 and PAP effectively inhibited H2O2-induced c-Myc translocation. \*\*P<0.01, vs.  $H_2O_2$  group. (F) PAP suppressed  $H_2O_2$ -induced c-Myc transcriptional activity, the relative luciferase activities were normalized to pRL-TK Renilla luciferase. (C, G) Pre-treatment with PAP, SCH772984 prior to  $H_2O_2$  significantly reduced

the expression of MMP-9. The results represent as mean  $\pm$  SD, and \*\*P<0.01 vs. H<sub>2</sub>O<sub>2</sub> group.