Eclalbasaponin I causes mitophagy to repress oxidative stress-induced apoptosis via activation of p38 and ERK in SH-SY5Y cells

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Supporting Information

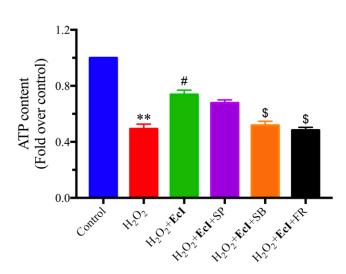


Figure S1. Cells were pre-treated with SP, SB or FR and cultured with **EcI** (50 μ M) prior to 200 μ M H₂O₂. The ATP levels were quantified in SH-SY5Y cells. Data are presented as means \pm S.D. (n=3). **p<0.01 vs. control group, *p<0.05 vs. H₂O₂ treated group. \$p<0.05 vs. (**EcI**, 50 μ M + H₂O₂, 200 μ M) group.

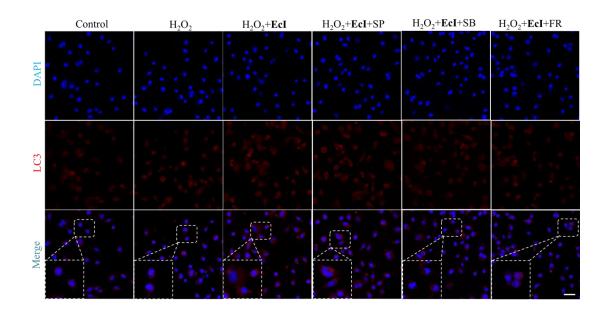


Figure S2. Cells were pre-treated with SP, SB or FR and cultured with **EcI** (50 μ M) prior to 200 μ M H₂O₂. The distribution of LC3 was investigated by immunofluorescence analysis. The scale bar represents 50 μ m.

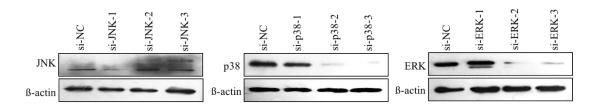


Figure S3. Cells were transfected with negative control siRNA (si-NC) or siRNA targeting JNK (si-JNK), p38 (si-p38), or ERK (si-ERK) followed by Western blot analysis. The specific sequences si-JNK-1, si-p38-3 and si-ERK-2 were used in following assays. NC means negative