Supplemental Figures

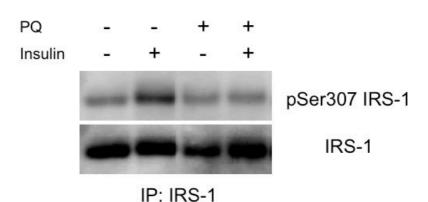
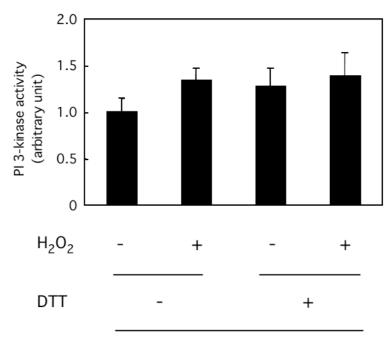


Fig. S1 Effect of paraquat on phosphorylation of IRS-1.

3T3-L1 adipocytes were serum-starved for 18 h and were then treated with 0 mM or 10 mM paraquat (PQ) for 3 h followed by incubation with 10 nM insulin for 5 min. Cells were then solubilized. Equal amounts of protein were subjected to SDS-PAGE and immunoblotting was performed with indicated antibody. Representative results of three independent experiments are shown.



myc-p85 α , FLAG-p110 α

Fig. S2 Effect of H₂O₂-treatment on PI 3-kinase activity of p110a in cell-free system.

p110 PI 3-kinase was immunoprecipitated with anti-FLAG antibody from HEK293T cells co-transfected with myc-p85 α and FLAG-p110 α . The immunoprecipitates were treated with 0.5 mM H₂O₂ in lysis buffer for 1.5 h at 4°C. After washing the immunoprecipitates with lysis buffer, LiCl, distilled water, TNE buffer, and reaction buffer, PI 3-kinase assay were performed with or without DTT. The experiments were performed in triplicate, and the results are shown as the mean \pm SEM.