## Supplementary data

Figure S1. (A)The breast cancer cell lines MDA-MB-231 and Hs-578T were transiently transfected with control or RECK siRNA for 12h. RECK mRNA was analyzed by qRT-PCR (mean  $\pm$  SD, n=3) analysis. \*\* P<0.01 compared with si-con transfected MDA-MB-231 cells and ## P<0.01 compared with si-con transfected Hs-578T cells.(B)The breast cancer cell lines MDA-MB-231 and Hs-578T were transiently transfected with si-con or RECK siRNA for 12h, then exposed to 0.0 or10.0µM of ISL for 24h. RECK mRNA was analyzed by qRT-PCR (mean±SD, n=3) analysis. \*\* P<0.01 compared with si-con transfected MDA-MB-231 cells treated with  $10.0 \ \mu$  M of ISL and ## P<0.01 compared with si-con transfected Hs-578T cells treated with 10.0 µ M of ISL.(C)Breast cancer cell lines Hs-578T and MDA-MB-231 cells were transiently transfected with anti-con or anti-miR-21 for 12h. Relative expression level of miR-21 was determined by qRT-PCR (Fold, mean  $\pm$  SD, n=3). \*\* P<0.01 compared with anti-con transfected MDA-MB-231 cells and ##  $P \le 0.01$  compared with anti-con transfected Hs-578T cells. (D)MDA-MB-231 and Hs-578T cells were transiently transfected with mimic-con or mimic-miR-21 for 12h, then exposed to 10.0µM of ISL for 24h. Relative expression level of miR-21 was determined by qRT-PCR (Fold, mean±SD, n=3). \*\* P<0.01 compared with mimiccon transfected cells and # P $\leq$ 0.01 compared with mimic-con-transfected cells treated with 10.0 µ M of ISL.

