



Figure S2. Depletion of Sec61 β causes ER stress.

(A) COS-7 cells infected with shRNA-expressing viruses were immunostained for Calreticulin, Tubulin, GM130 and TOM20 respectively (with DAPI staining), and visualized by confocal microscopy. Scale bar: 10 μ m.

(B) COS-7 cells were infected with shRNA-expressing viruses, and the level of Sec61 β determined by Western blotting.

(C) The levels of Climp63 in indicated cells was determined by Western blot, and the unspliced (U) *XBP1* and spliced (S) *XBP1* by RT-PCR of *XBP1* mRNA were resolved by agarose gel.

(D) COS-7 cells were transfected with siRNAs of REEP1 for 48 hours or 96 hours. The levels of phosphorylated eIF2 α were determined by Western blotting, and the *REEP1*, *GAPDH*, unspliced *XBP1* and spliced *XBP1* by RT-PCR were resolved by agarose gel. Asterisk (*) indicates a nonspecific band.

(E) MEF cells were transfected with siRNAs of REEP1 for 48 hours or 96 hours. The levels of phosphorylated eIF2 α and REEP1 were determined by Western blotting, and the unspliced *XBP1* and spliced *XBP1* by RT-PCR of *XBP1* mRNA were resolved by agarose gel. TG treated group was the positive control. Asterisk (*) indicates a nonspecific band.

(F) Lysates of generated Flp-In-293 cell lines were analyzed with Western blotting. The band with asterisk (*) may be degraded RFP-Sec61 β -HA.