Relative cell growth to the ETOH-treated control cells with Negative siRNA control

(A)

PC-3 – ERβ mRNA

(B)

PC-3 – Cell growth

(C)

PC-3 – miR-765

(D)

DU145 – ERβ mRNA

(E)

DU145 – Cell growth

(F)

DU145 – miR-765
Figure S5. Blocking effects of another ERβ siRNA on prostate cancer cell growth and up-regulation of hsa-miR-765 expression. (A and D) Effectiveness of siRNA#2 on knockdown of ERβ in PC-3 (A) and DU145 (D) cells. PC-3 and DU145 cells were treated with fulvestrant or ethanol (Control) in the presence of ERβ siRNA or negative-control siRNA for 4 days. The levels of ERβ in the cells were quantified by real-time RT-PCR analysis. Fulvestrant inhibits PC-3 and DU145 cell growth and up-regulated has-miR-765 expression via an ERβ-dependent mechanism. (B and E) Growth of the fulvestrant-treated PC-3 (B) and DU145 (E) cells with or without ERβ siRNA knockdown for 4 days relative to the ethanol-treated control cells with negative-control siRNA are presented and compared (n=8) (C and F) Hsa-miR-765 is induced by fulvestrant in PC-3 (C) and DU145 (F) cells in an ERβ-dependent mechanism. The hsa-miR-765 in the fulvestrant- and ethanol-treated control cells was quantified by miRNA qRT-PCR analysis. Relative fold changes between the expression of hsa-miR-765 in the fulvestrant-treated and control cells are presented. Columns=means; bars =S.D.; n=3.