Figure S1. Propofol inhibits the invasion of bladder cancer cells by regulating miR-145-5p expression. (A) J82 and (B) T24 cells were transduced with anti-miR-145-5p, anti-miR-NC or miR-145-5p and exposed to $10 \mu g/ml$ propofol, then cells were used for Transwell cell invasion assays. Scale bar, $100 \mu m$. Each assay was performed in triplicate and data are presented as the mean \pm SD. *P<0.05. miR, microRNA; NC, negative control.

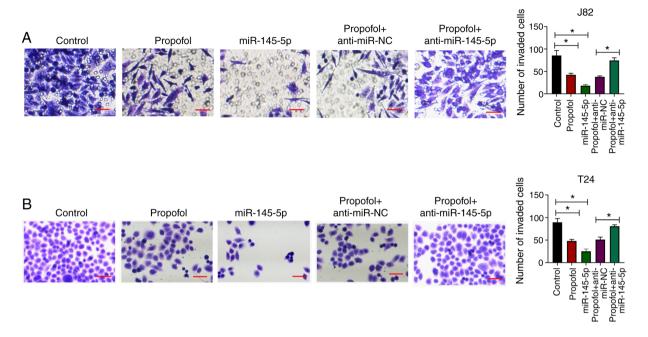


Figure S2. Propofol suppresses the migration and invasion of bladder cancer cells via the miR-145-5p/TOP2A axis. J82 and T24 cells were transduced with anti-miR-NC, anti-miR-145-5p, si-NC or si-TOP2A, then treated with 10 μ g/ml propofol or an equal volume of DMSO as control. (A) J82 and (B) T24 cells were used for wound-healing assays (scale bar, 500 μ m), or (C) J82 and (D) T24 cells were used for Transwell cell invasion assays (scale bar, 100 μ m). Each assay was performed in triplicate and data are presented as the mean \pm SD. *P<0.05. miR, microRNA; NC, negative control; si, small interfering RNA; TOP2A, topoisomerase II α .

