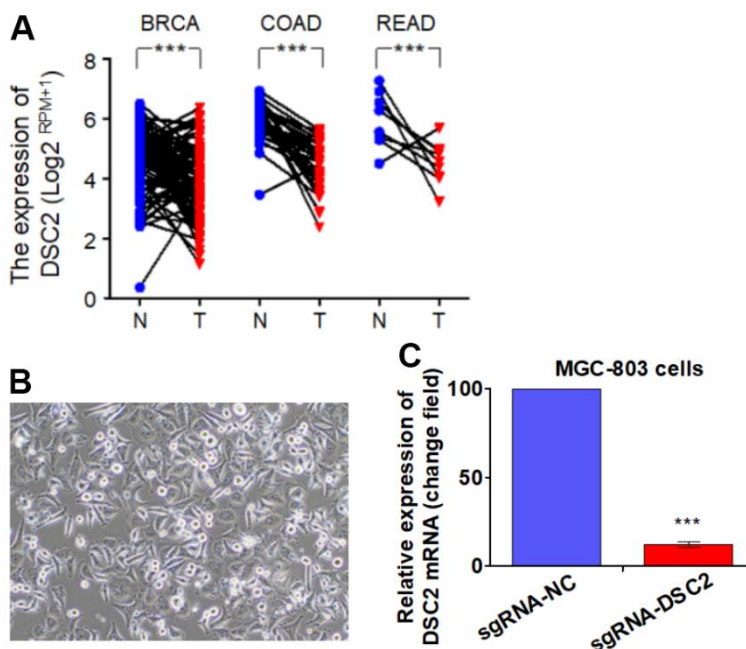
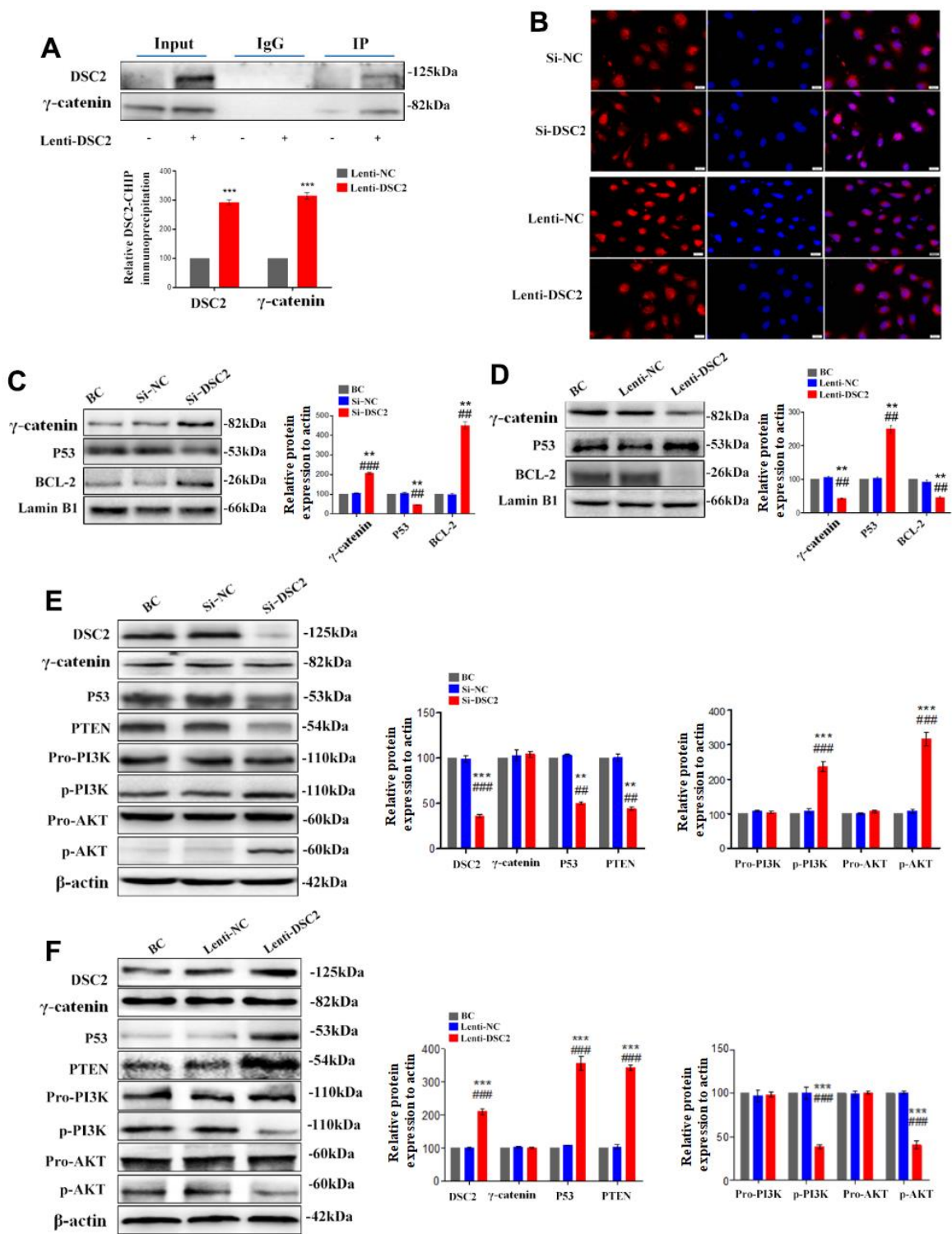


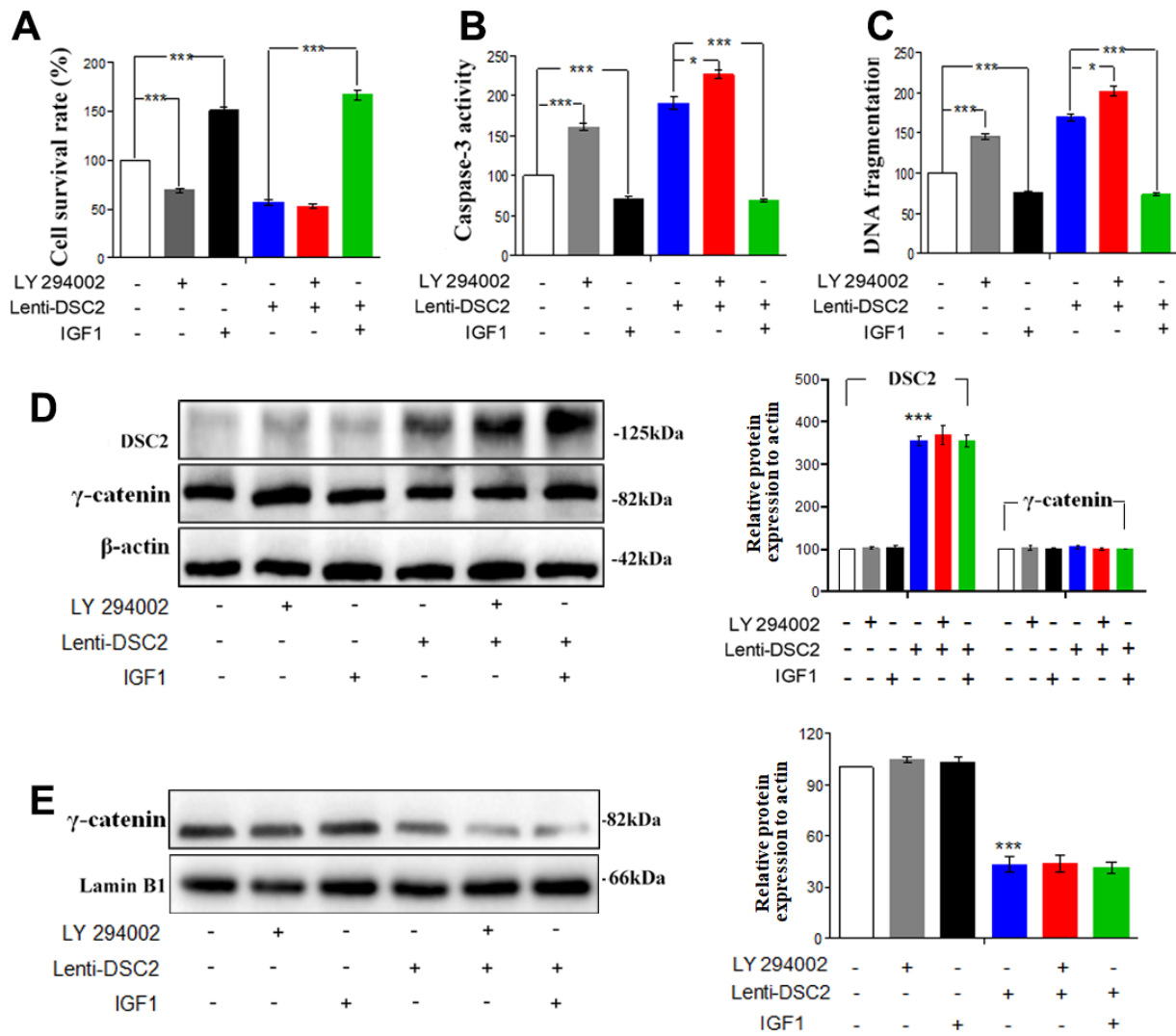
SUPPLEMENTARY FIGURES



Supplementary Figure 1. DSC2 was knocked down in MGC-803 cells significantly after transfecting with vectors that were inserted into the CRISPR EGFP plasmid. (A) Positive clone MGC-803 cells were photoed (100 ×). **(B)** DSC2 gene in sgRNA-NC and sgRNA-DSC2 were detected by qRT-PCR. Data are presented as mean ± SEM from three separate experiments. *** $p < 0.001$ vs. sgRNA-NC. **(C)** Comparison of DSC2 expression of paired samples in breast invasive carcinoma (BRCA), colon adenocarcinoma (COAD), rectum adenocarcinoma (READ), which was collected from the TCGA database. *** $p < 0.001$ vs. Tumor tissues.



Supplementary Figure 2. DSC2 inhibited the γ -catenin nuclear translocation and suppressed the PI3K/AKT signaling pathway of SGC-7901 cells. (A) Co-IP assay was performed to analyze the interaction of DSC2/ γ -catenin by DSC2. The data are represented as mean \pm SEM, $n=3$. *** $p<0.001$ vs. Lenti-NC. After being transfected with siDSC2 or stably expressing DSC2 gene of SGC-7901 cells, (B) the level of γ -catenin accumulated in the nucleus was detected by immunofluorescence assay. The scale bar = 20 μ m. (C) The expression of γ -catenin, BCL-2 and P53 in nucleus was determined by Western blot assay. (D) The expressions of γ -catenin, P53, PTEN, pro-PI3K, p-PI3K, pro-AKT, p-AKT were detected by Western blot assay. The data are represented as mean \pm SEM, $n=3$. ** $p<0.01$ and *** $p<0.001$ vs. BC. ## $p<0.01$ and ### $p<0.001$ vs. Si-NC or Lenti-NC group. (E, F) the expressions of PTEN, pro-PI3K, p-PI3K, pro-AKT, p-AKT were detected by Western blot assay. Data are presented as mean \pm SEM from three separate experiments. *** $p<0.001$ vs. BC. ### $p<0.001$ vs. Si-NC or Lenti-NC.



Supplementary Figure 3. DSC2 inhibited viability of SGC-7901 cells through suppressing PI3K/AKT signaling pathway. Effect of DSC2 on the viability of GC cells in the presence of LY294002 and IGF1 was determined by Caspase-3 activity assay (A), Sperm DNA fragmentation assay (B) and MTT assay (C). Data are presented as mean \pm SEM from three separate experiments. * $p < 0.05$ and *** $p < 0.001$ vs. Lenti-NC or Lenti-DSC2 group. (D, E) The levels of γ -catenin both in cells and in the nucleus among SGC-7901 cells that treated with LY294002 or IGF1, were tested by Western blot assay. The data are represented as mean \pm SEM, $n=3$. *** $p < 0.001$ vs. Lenti-NC.