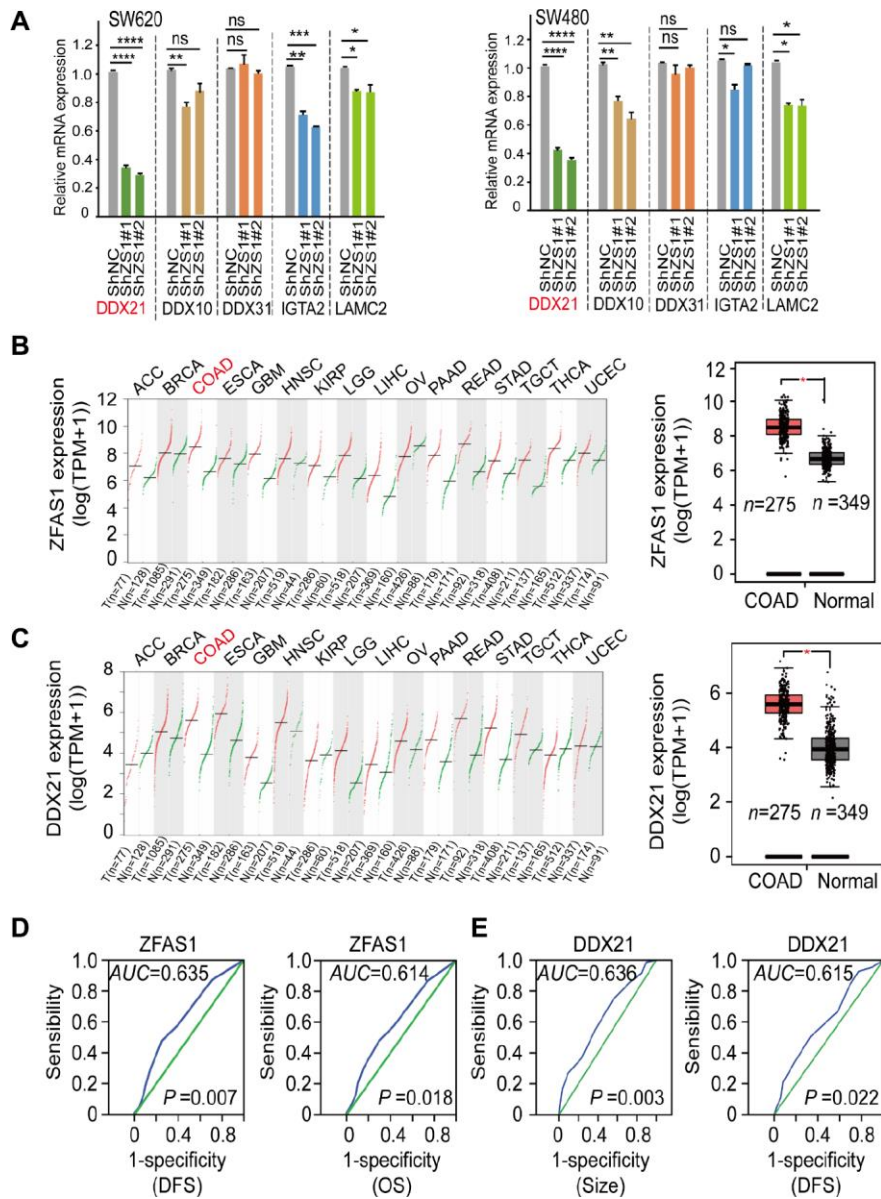
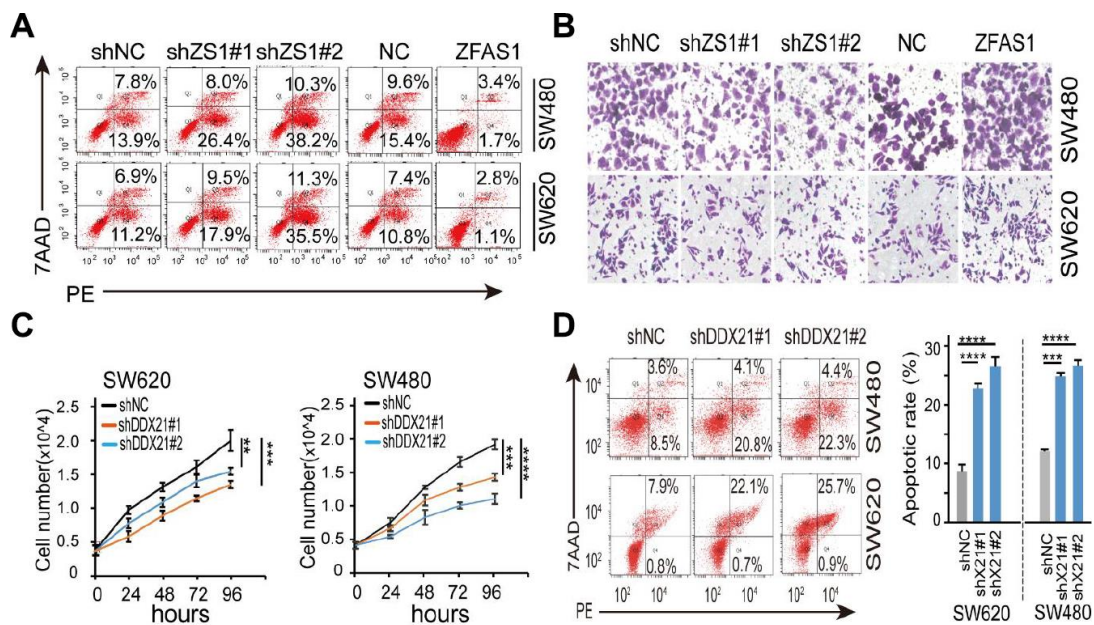


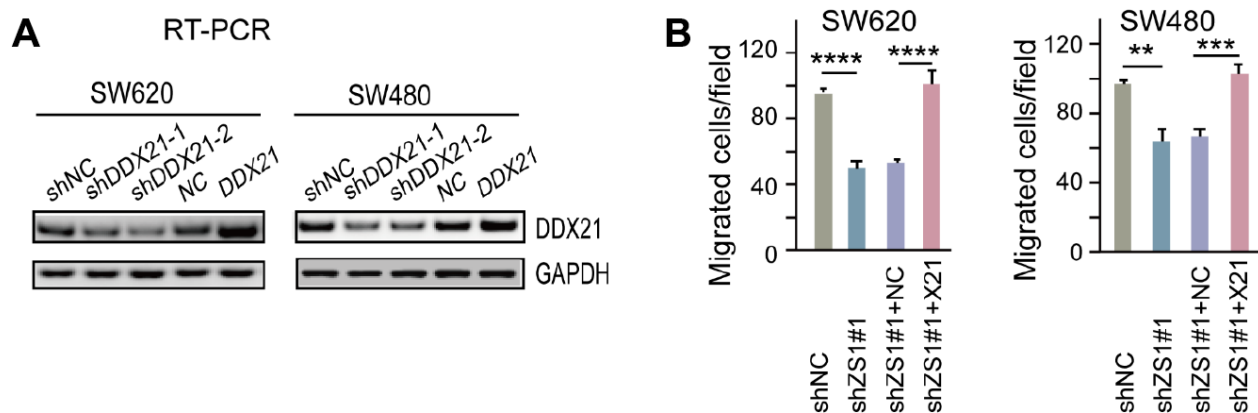
SUPPLEMENTARY FIGURES



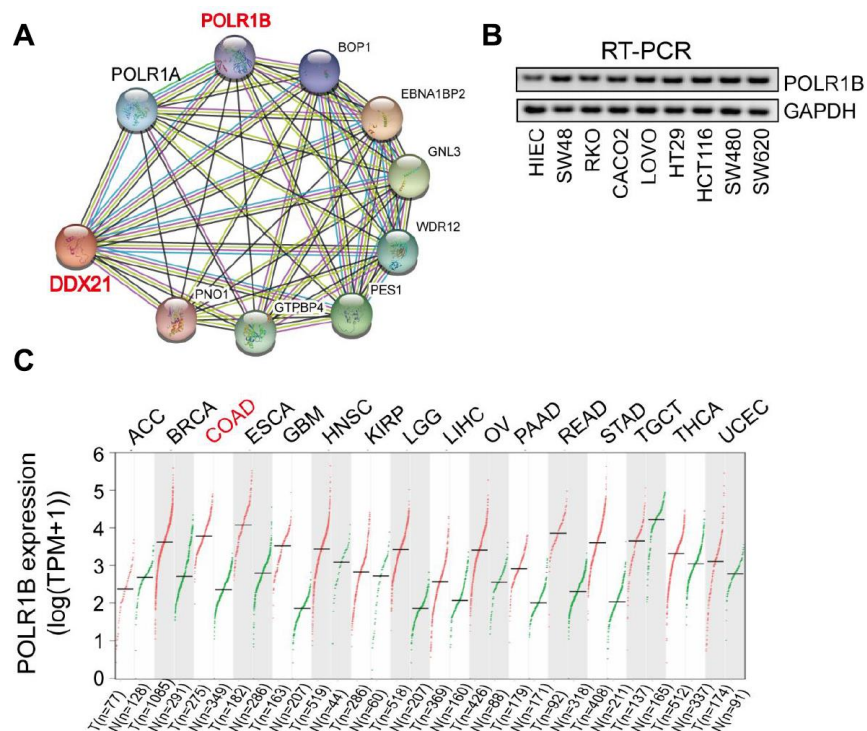
Supplementary Figure 1. The expression of lncRNA ZFAS1 and its correlation with DDX21 expression based on the TCGA datasets and our included paired CRC patient tissues. (A) The mRNA levels of potential target indicators were detected after silencing lncRNA ZFAS1 in SW620 and SW480 cells by qPCR assay. **(B and C)** RNA-seq analysis illustrating the log2 gene expression of lncRNA ZFAS1 and DDX21 in a variety of tumor tissues (left) and comparison the relative expression in CRC patients tissues (n = 275) and the normal controls (n = 375) (right) based on the TCGA data (<http://gepia.cancer-pku.cn/>). **(D and E)** The ROC curves illustrating the cutoff values of lncRNA ZFAS1 and DDX21 expression in paired CRC patient tissues vs. adjacent-tumor control tissues (n=157).



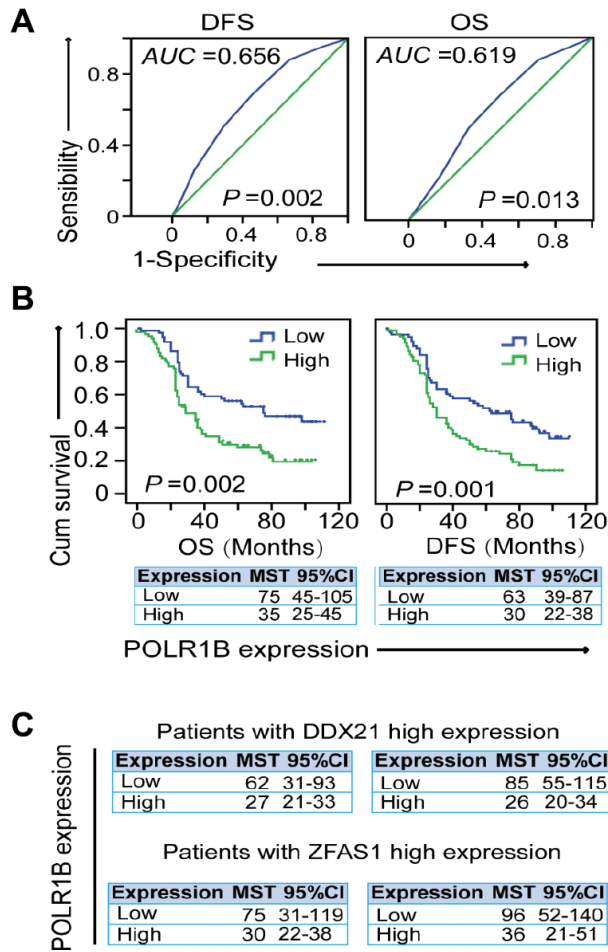
Supplementary Figure 2. The effect of lncRNA ZFAS1 and DDX21 on biological characteristics of CRC cells. (A) The cell apoptotic rates were determined after treated with lncRNA ZFAS1 overexpressing or silencing vector in both SW620 and SW480 cells by Flow cytometry. (B) The invasion ability was determined after ectopic or knockdown lncRNA ZFAS1 in both SW620 and SW480 cells. $n = 3$ independent experiments. (C) Cell number monitoring assays showed the cell proliferation after silencing DDX21. (D) Flow cytometry assays determined the cell apoptotic rates after DDX21 knockdown.



Supplementary Figure 3. DDX21 rescued the CRC cells proliferation inhibition caused by ZFAS1 knockdown. (A) The expression of DDX21 after silencing or overexpression DDX21 in SW620 and SW480 cells by RT-PCR assay. (B) The migrated cells per field were detected after co-transfected with shRNA ZFAS1 and pcDH-DDX21 vector in SW620 and SW480 cells using Trans-well assays.



Supplementary Figure 4. The expression of POLR1B in the CRC cells and the TCGA datasets. (A) STRING (<https://string-db.org/>) predicting the functional protein association networks of DDX21 including POLR1B. **(B)** RT-PCR assay detecting the POLR1B expression in CRC cells including SW480, SW620, HCT116, SW48, CACO2, LOVO, HT29, and RKO cells vs. normal intestinal epithelial HIEC cells. **(C)** RNA-seq analysis illustrating the log₂ gene expression of POLR1B in a variety of tumor tissues based on the TCGA data (<http://gepia.cancer-pku.cn/>).



Supplementary Figure 5. Stratification analysis of the correlation of POLR1B with prognosis of CRC patients. (A) The ROC curves illustrating the cutoff values of POLR1B expression in paired CRC patient tissues vs. adjacent-tumor control tissues (n=157). (B) Kaplan-Meier plot representing the impact of POLR1B high/low expression on the DFS and OS in this cohort (n=157). (C) Stratified Log-rank test showing the MST and 95%CI of POLR1B high/low expression on the DFS and OS based on the patients with DDX21 high expression (upper) or ZFAS1 high expression (lower).