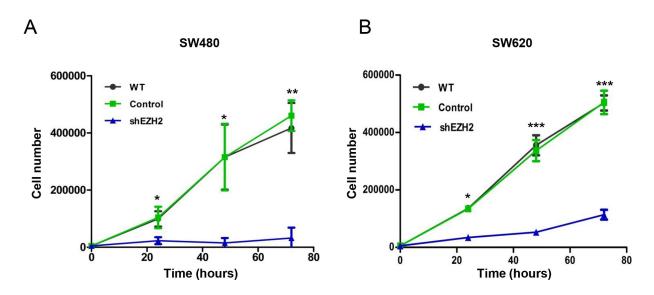
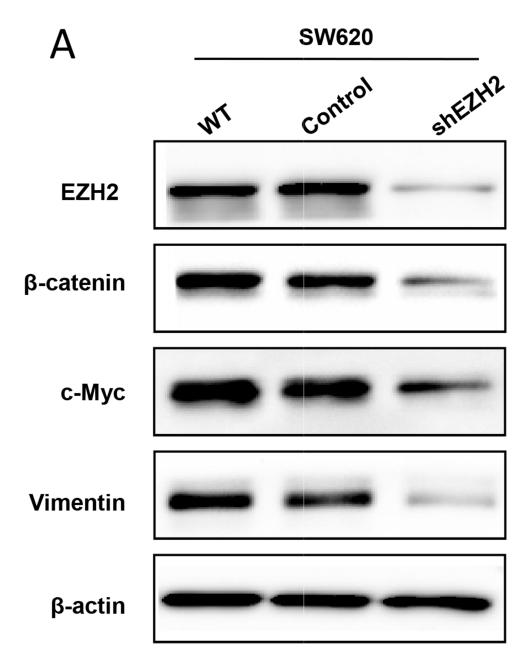
EZH2 promotes colorectal cancer stem-like cell expansion by activating $p21^{cip1}$ -Wnt/ β -catenin signaling

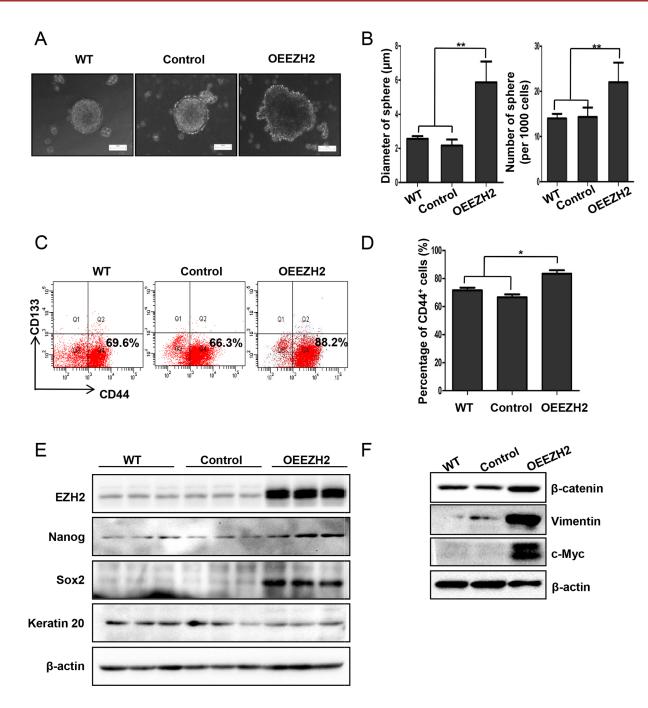
SUPPLEMENTARY FIGURES



Supplementary Figure S1: EZH2 promoted proliferation and clonogenicity of CRC cells. EZH2 was knocked down in **A.** SW480 and **B.** SW620. After 24, 48 and 72 hr of incubation, cell proliferation was measured by cell counting assay.



Supplementary Figure S2: Silencing EZH2 expression inactived Wnt/ β -catenin signaling pathway in SW620 cell line. A. The expression of β -catenin, vimentin and c-myc were detected by western blot after knockdown EZH2 in SW620 cells.



Supplementary Figure S3: Overexpression EZH2 promoted stemness of LoVo cell line. A. Wild type (WT), scramble OERNA (Control) and OEEZH2 of LoVo cells were plated in low-serum non-adherent culture conditions. Images were obtained by microscopy at $10 \times \text{magnification}$ and were representative of all mammospheres formed (Scale bar = $100 \, \mu \text{m}$). B. The size and number of LoVo cell spheres were analyzed. Data were presented as the mean ± S.D. of three independent experiments. C. CD133+/CD44+ population was analyzed by flow cytometry analysis in LoVo cells with or without EZH2 overexpression. D. The statistical results of three independent experiments were presented. E. After EZH2 knockdown, the expression of EZH2, Nanog, Sox2, and Keratin 20 was examined by western blot. F. The expression of β-catenin, vimentin and c-myc were detected by western blot after overexpression EZH2 in LoVo cells.