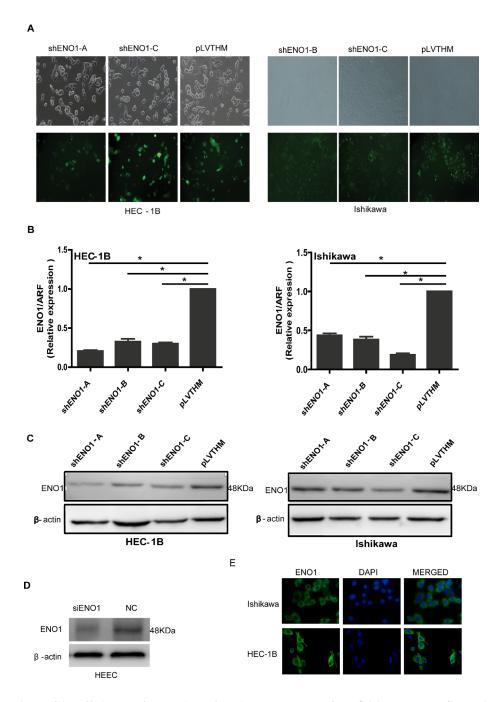
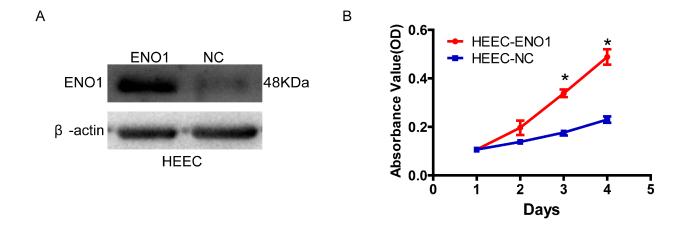
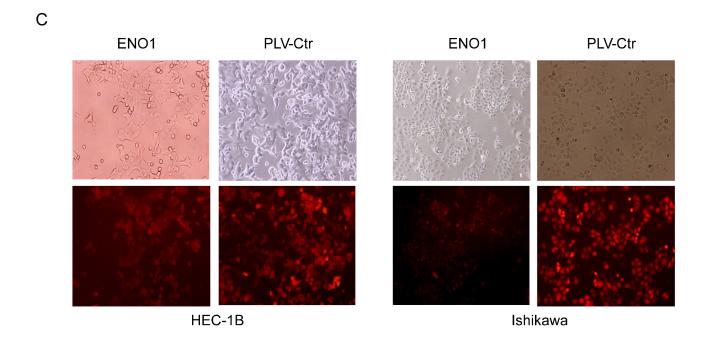
SUPPLEMENTARY FIGURES

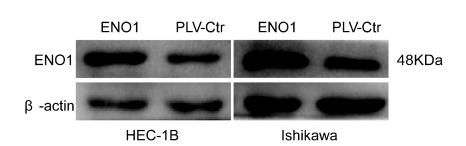


Supplementary Figure S1: Efficiency of shRNA or siRNA knock down of ENO1 in human EC cell lines HEC-1B and Ishikwawa and HEEC. A. HEC-1B and Ishikwawa were transfected by different lentivirus which contain sh-ENO1 and PLVTHM. B. RT-PCR showing transcriptional levels of the ENO1 gene with ARF used as a loading control. Bar graph showed the relative expression of mRNA among the groups. Data were presented as mean \pm SEM for three independent experiments (*P < 0.05). C. Western blotting showing protein expression levels in shENO1 or PLVTHM cells. A representative image of three different experiments was shown. β -actin served as a loading control. D. Western blotting showing protein expression levels in siENO1-HEEC and HEEC-NC cells. Each experiment was repeated three times. E. Immunofluorescence staining of ENO1 in Ishikawa and HEC-1B cells (magnification, \times 1000).

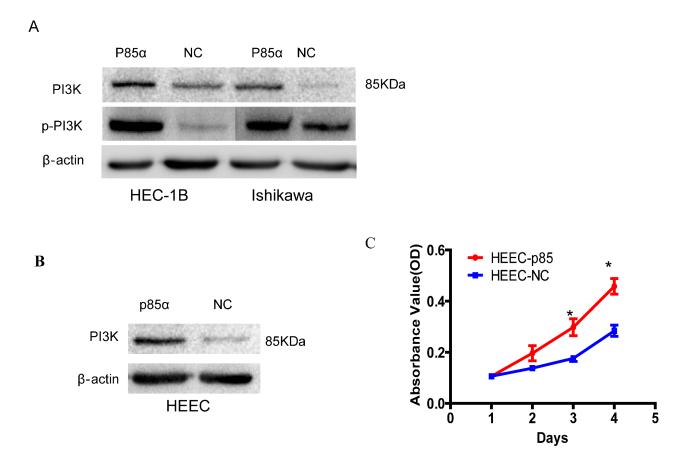
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Supplementary Figure S2: Overexpressing ENO1 in HEEC and EC cells. A. Overexpressing ENO1 increased the protein level of ENO1 in HECC detected by western blotting. β-actin served as the internal control. Each experiment was repeated three times. B. MTT assay showed that elevated ENO1 induced cell proliferation in HEEC. Data were presented as mean \pm SEM for three independent experiments (*P < 0.05). C. HEC-1B and Ishikawa were transfected by lentivirus which contain full-length ENO1 and RFP empty vector (PLV-Ctr). D. Western blotting showing protein expression levels in ENO1 or PLV-Ctr cells. A representative image of three different experiments was shown. β-actin served as a loading control.



Supplementary Figure S3: Overexpression of P85 expression in EC cells and HEEC. A. Overexpressing p85 increased the protein level of PI3K and p-PI3K in EC cells detected by western blotting. β-actin served as the internal control. Each experiment was repeated three times. B. Western blotting showing PI3K protein expression levels in P85 overexpressed HEEC and its control cells. A representative image of three different experiments was shown. β-actin served as a loading control. D. Effect of p85 overexpressed on HEEC cell proliferation as measured by MTT assay. Data were presented as mean \pm SEM for three independent experiments (*P < 0.05).