

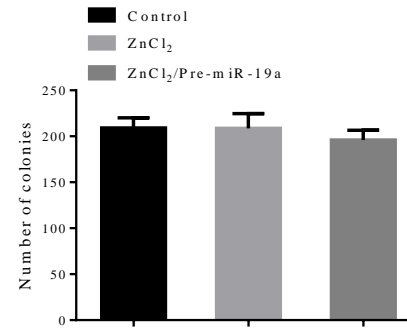
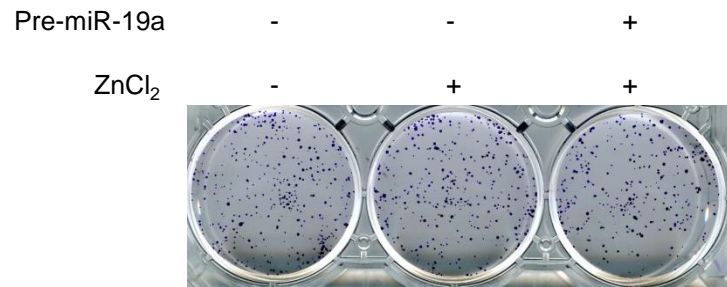
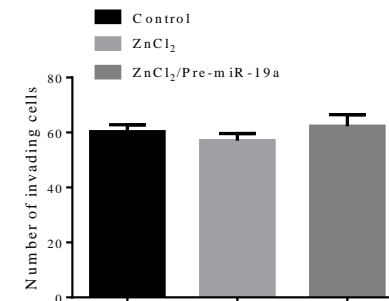
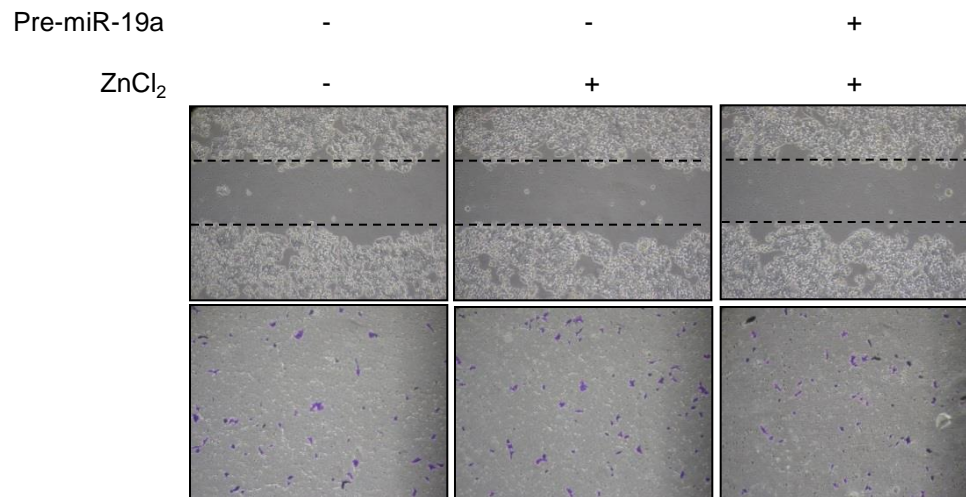
Supplemental Figures

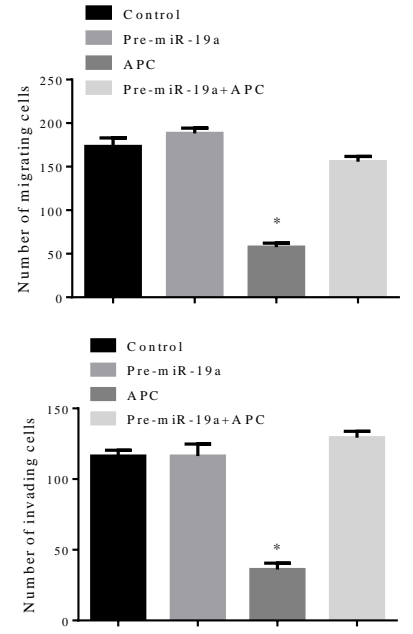
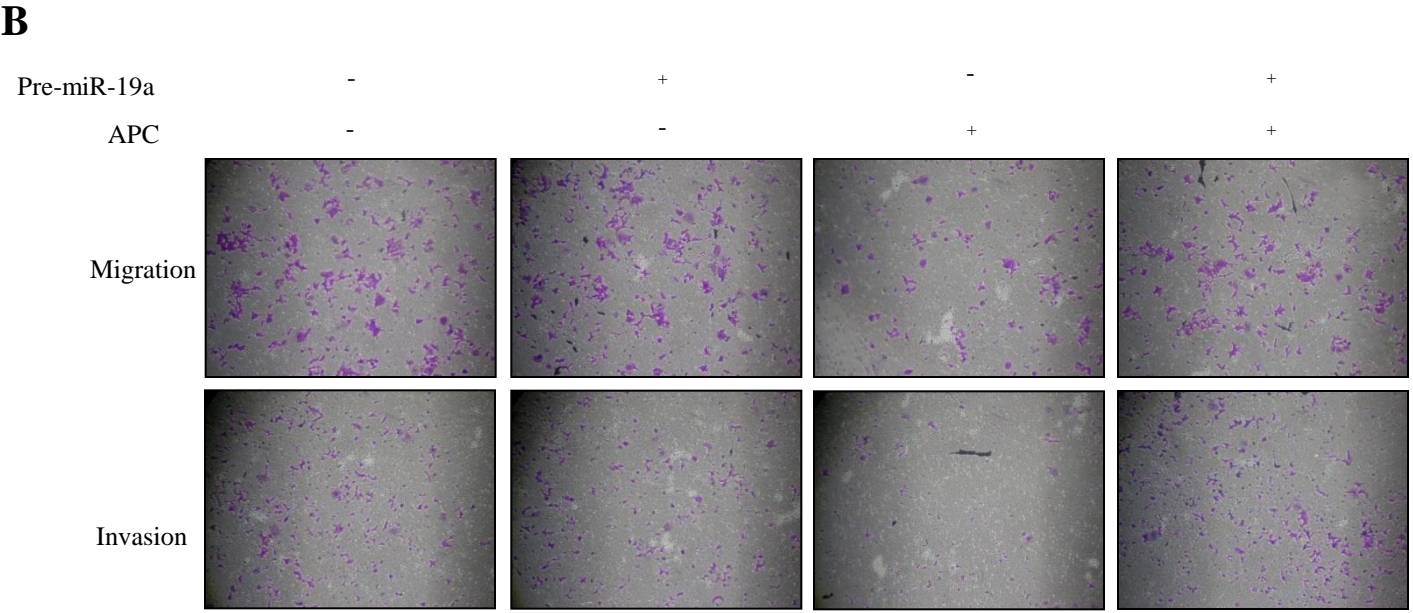
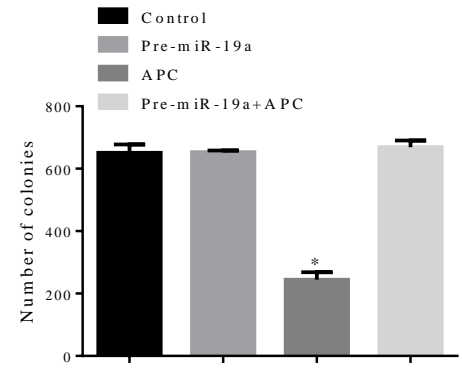
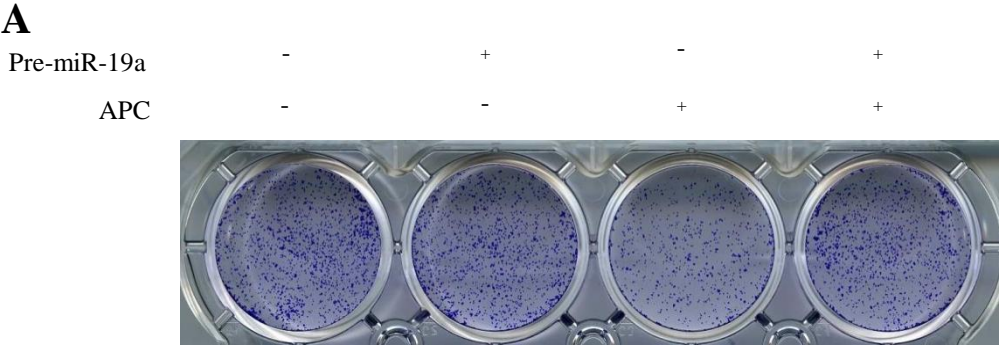
Figure S1. Expression of miR-19a did not affect β -Gal/HT29 cell growth, migration and invasion. β -Gal/HT29 cells were transfected with pre-miR-19a in the culture medium containing ZnCl_2 and then subjected to colony growth (A), wound healing and 2-chamber cell invasion assay (B).

Figure S2. APC repressed cell growth, migration and invasion were abrogated by enforced expression of miR-19a. HCT15 cells, which carry APC mutation, were transfected with APC together with and without pre-miR-19a and then assayed for colony growth (A), cell migration and invasion (B).

Figure S3. Knockdown of miR-19a reduces colony growth, cell migration and invasion in HCT15 cells. Anti-miR-19a and control oligo were introduced into HCT15 cells and then subjected to colony formation (A), wound healing and 2-chamber invasion assays (B).

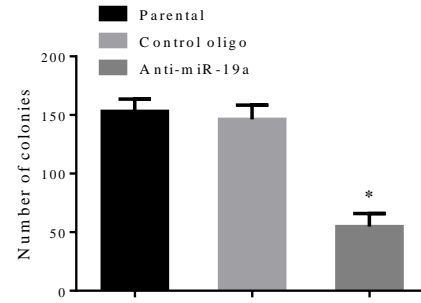
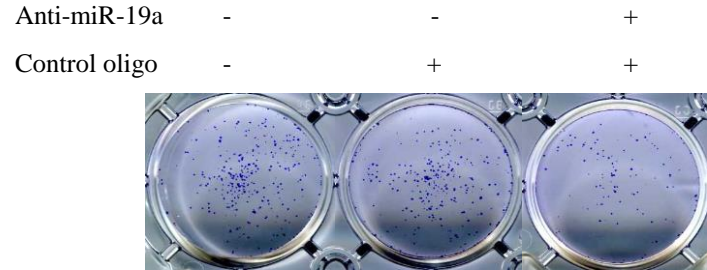
Figure S4. Expression of miR-19a in NCI-H508 cells promotes colony growth, cell migration and invasion. Pre-miR-19a and control oligo were introduced into NCI-H508 cells, in which both APC and β -catenin are wild type. After incubation for 36 hours, cells are assayed for colony formation (A), cell migration and invasion (B).

A**B**

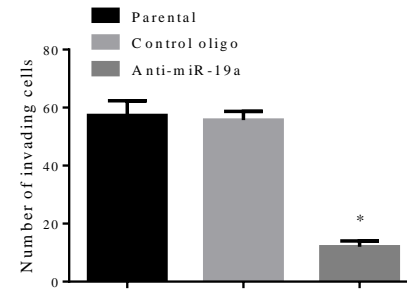
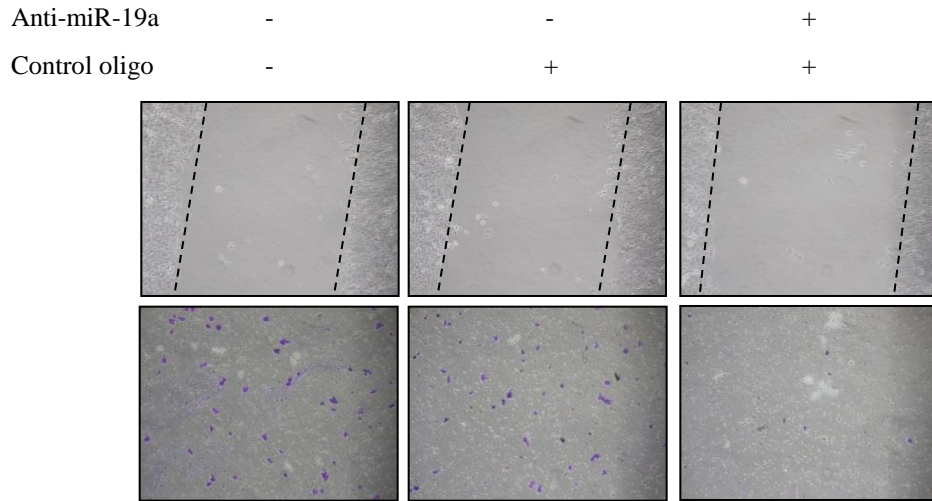


Supplemental Fig. 3

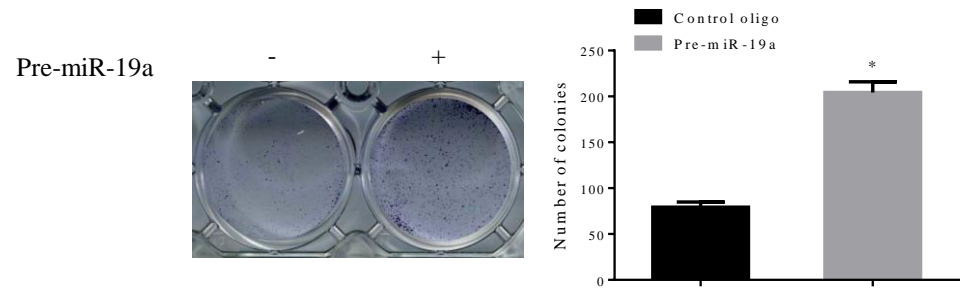
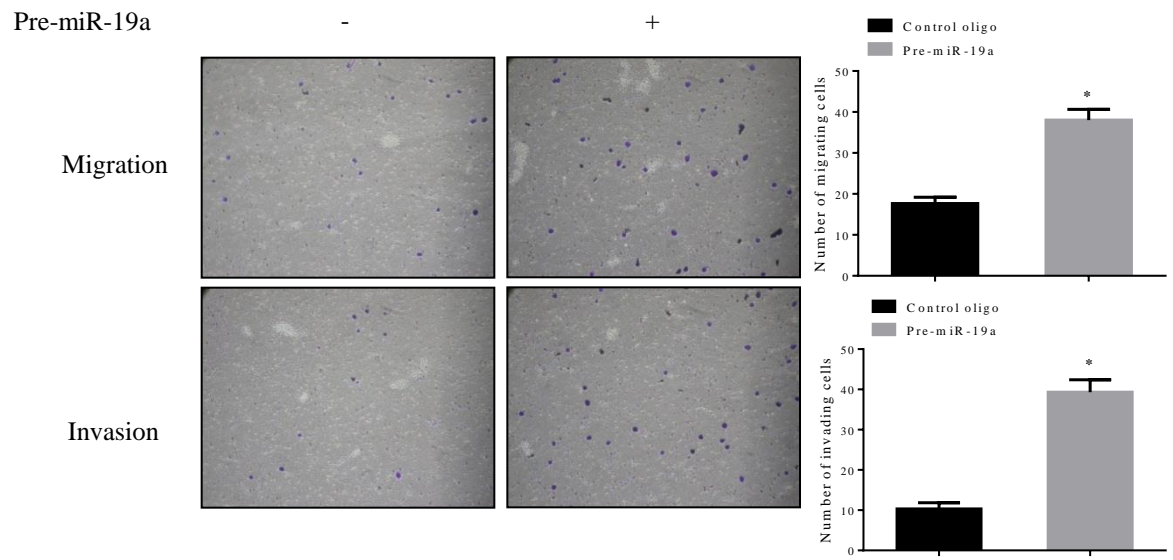
A



B



Supplemental Fig. 3

A**B**

Supplemental Fig. 4