Additional file 8

Figure S4. RIP assay confirmed that miR-143 was a target of PVT1. (a-b) The correlation among PVT1, miR-143 and Ago2 was determined by analyzing GBC-SD and NOZ cell lysates utilizing RIP with an Ago2 antibody. (c-d) qPCR was performed to detect the PVT1 level in the substrate of RIP assay in miR-143-overexpressing GBC cells. (e) Detection of PVT1 using qPCR in the sample pulled down by biotinylated PVT1 and negative control (NC) probe. (f) Detection of miR-143 using qPCR in the same sample pulled down by biotinylated PVT1 and NC probe. Input was used for normalization. *P < 0.05, **P < 0.01, ***P < 0.001. Error bars indicate mean ± SD.

