Figure S1. The UCSC Xena tool was used to explore the multi-omics data of the TCGA-COAD and READ datasets. Gene expression (RNAseq) was evaluated using the normalized log₂(FPKM+1), and somatic mutations (deleterious, splice, missense/inframe, intron/RNA, splice, silent and complex or unannotated mutations) were estimated using MuTect2 variant aggregation and masking. Mutations in the checkpoint molecules CTLA-4, CD274 (PD-L1), PDCD1 (PD-1), PDCD1LG2 (PD-L2), ADORA2A, HAVCR2, IDO1, IDO2, LAG3, TIGIT and VTCN1 do not affect their corresponding expression levels.

