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CORRIGENDUM

ACAT2 Promotes Cell Proliferation and Associates with Malignant Progression in Colorectal Cancer [Corrigendum]

Weng M, Zhang H, Hou W, Sun Z, Zhong J, Miao C. *Onco Targets Ther.* 2020;13:3477–3488.

The authors have advised Figures 4A and 5K are incorrect. The correct Figures 4 and 5 are shown below.

The authors apologize for these errors and advise that this does not affect the results of the paper.

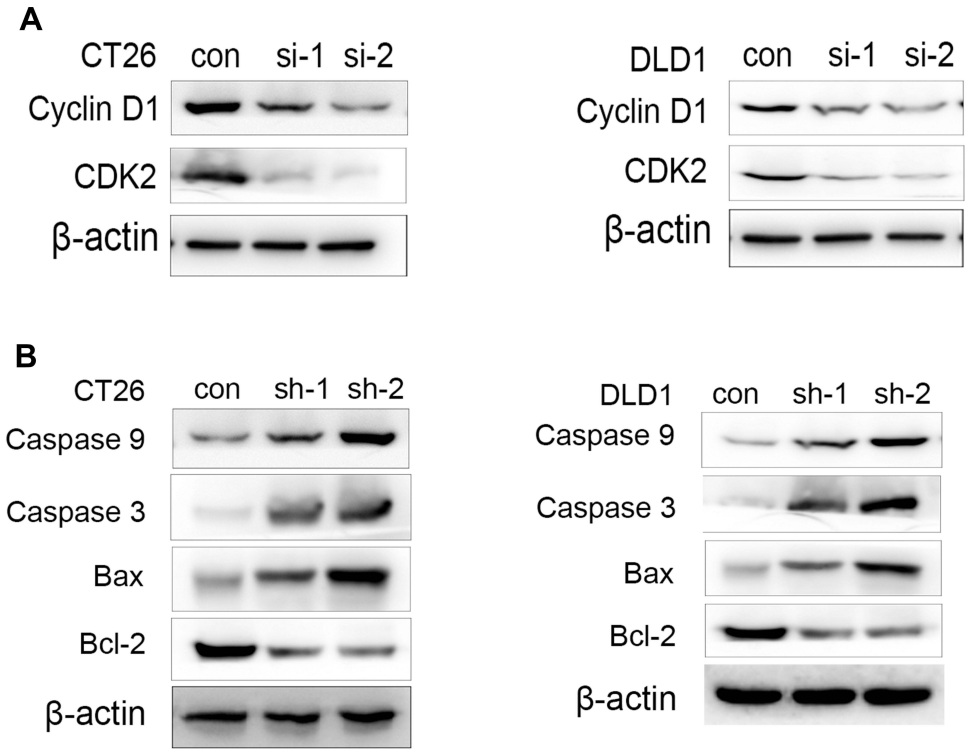

Figure 4 The regulation of some apoptotic marker in CT26 and DLD1 cells by ACAT2 knockdown. (**A**) The expression of cyclin D1 and CDK2 in CT26 or CT26 cells transfected with siRNA-ACAT2 (DLD1 or DLD1 cells transfected with siRNA-ACAT2). (**B**) The expression of caspase 3, caspase 9, Bcl-2 and Bax in CT26 or CT26 cells transfected with siRNA-ACAT2 (DLD1 or DLD1 cells transfected with siRNA-ACAT2).

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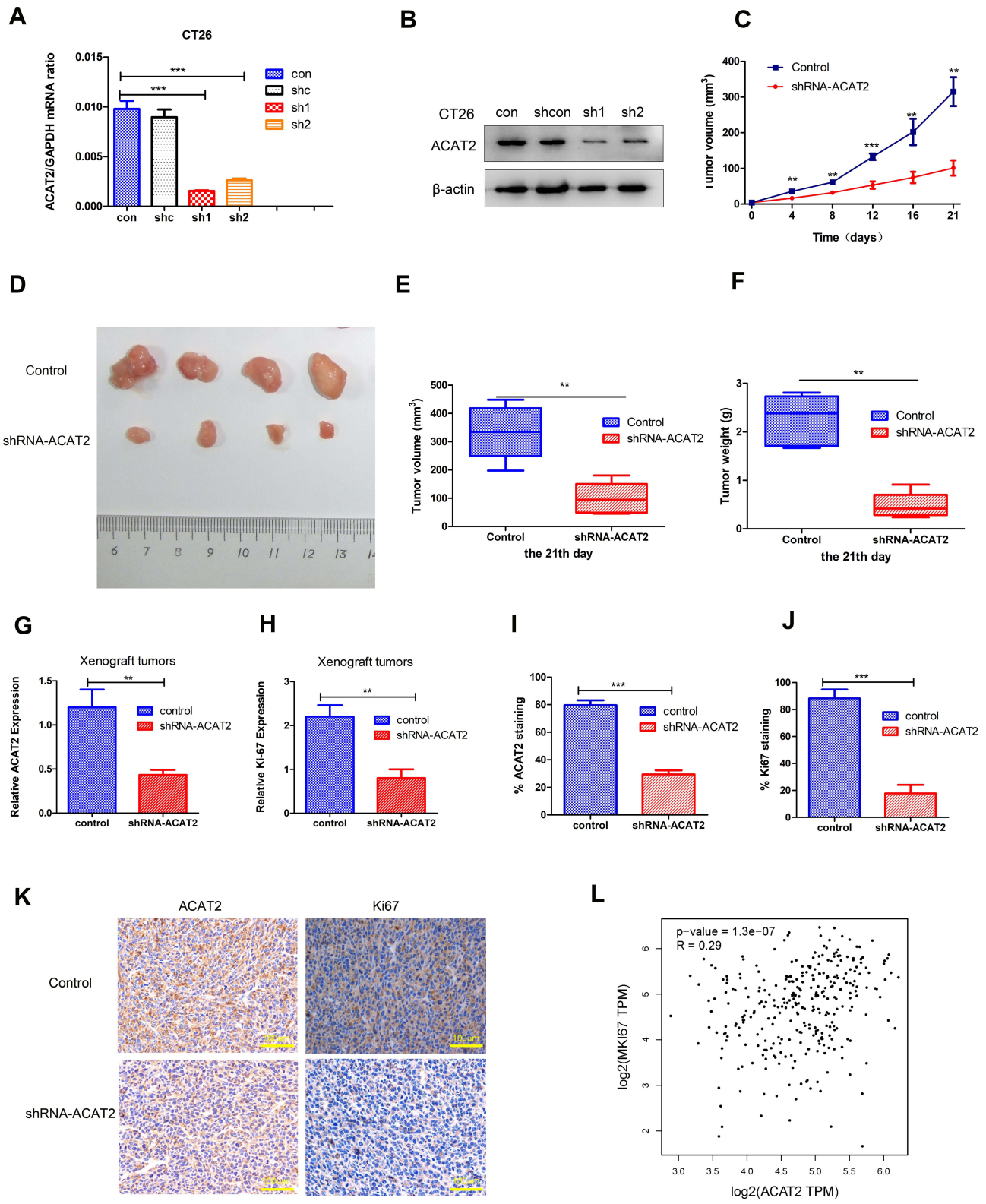

Figure 5 Knockdown of ACAT2 expression suppresses CRC growth and inhibits Ki-67 expression in vivo. (**A, B**) The efficiency of ACAT2 knockdown in CT26 cells was measured by qPCR and Western blotting. (**C, D**) CT26 cells or shRNA-ACAT2 CT26 cells were injected into BALB/c mice. Photograph of dissected tumors (upper: control group; lower: shRNA-ACAT2 group; n=4; P<0.01). The tumor volumes were measured every 3 days. The shRNA-ACAT2 in CT26 cells attenuated tumor growth in mice. (**E, F**) Tumor volumes and tumor weights on the 21st day (P<0.01). (**G, H**) The expression of ACAT2 and Ki-67 in tissues was detected by qPCR (P<0.01). (**I, J**) The graph shows the quantitative analysis of ACAT2 and Ki67 staining. (**K**) The protein expression ACAT2 and Ki67 in dissected tumor samples were evaluated by IHC. Scale bar: 100um. (**L**) Based on TCGA dataset analysis, ACAT2 expression was positively correlated with Ki-67 expression in CRC ($P=1.3\times10^{-7}$). **P<0.01, ***P<0.001; compared with the control group.