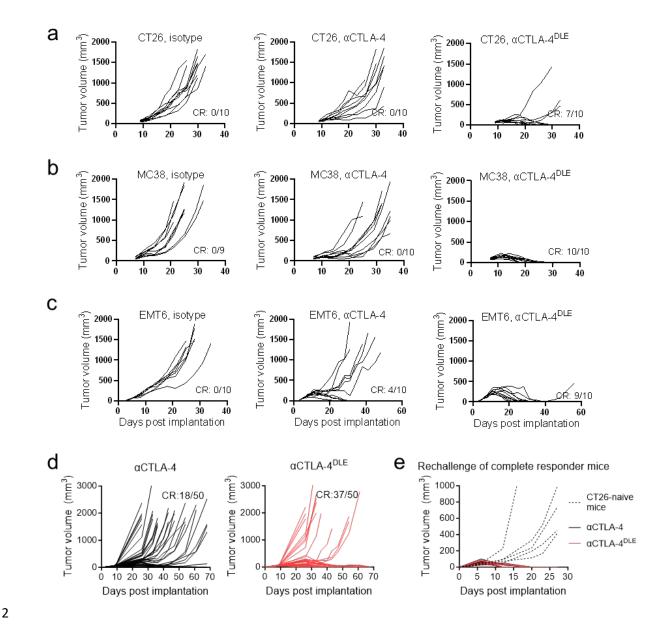
1 Supplementary Figure S2



Supplementary Figure S2. αCTLA-4^{DLE} promotes superior anti-tumor immunity and T cell infiltration in the tumor microenvironment versus parental αCTLA-4 in tumor-bearing mouse models. (a) Individual tumor growth of Balb/C mice bearing CT26 subcutaneous tumors (~50-75 mm³; n=10 mice/group) treated once intraperitoneally with 10 μg of indicated antibodies, (b) C57BL/6 mice bearing MC38 subcutaneous tumors (~120 mm³; n=9-10 mice/group) treated once intraperitoneally with 100 μg of indicated antibodies and (c) EMT6 breast orthotopic tumors

(~100 mm³; n=10/group) treated once weekly for three weeks with 100 μg antibody. (d) Individual tumor growth curves and number of complete responders (no measurable tumor) of Balb/C mice bearing CT26 subcutaneous tumors (~60 mm³; n=50 mice/group) and treated intraperitoneally with a single dose (100 μg) of antibody, as indicated. (e) Tumor growth following rechallenge of complete responder mice with CT26 cells (1x106) in the contralateral flank compared with tumor

growth in CT26-naive control mice.

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