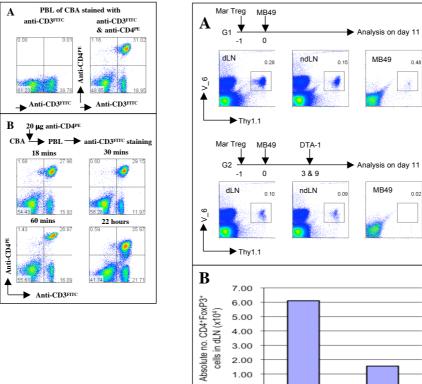
S-Fig.1 MB49-residential Treg cells of host origin DTA-1-treated Untreated 0.933 ±0.210 0.575 ±0.29 шет, тошет, тошет, з 10¹ 10² 10³ Fo×p3-FITC

S-Fig.2 S-Fig.3

0.48



3.00 2.00 1.00 0.00

G1

G2

Legends for supplementary figures

S-Figure 1. DTA-1 treatment also reduces the percentage of tumour-infiltrated Treg cells of host origin.

This experiment was conduced as described in Fig.3C. On day 11, MB49 tumours from individual mice in untreated (G2) and DTA-1-treated group (G4) were stained with anti-V β 6^{PE}, anti-Thy1.1^{PerCP} and anti-CD4^{APC} followed by intracellular staining of anti-Foxp3^{FITC}. Percentage of tumour-infiltrated Treg cells of host origin (Foxp3⁺Thy1.1⁻CD4⁺) of individual mice in each group is shown as mean ±SEM. One of representative of two independent experiments is shown.

S-Figure 2. Binding to CD4 T cells by injected PE-conjugated anti-CD4 mAb.

A. The PBL from a group of CBA mice (n=3) were stained with anti-CD3^{FITC} alone or together with anti-CD4^{PE}. The dot-plots show CD3 versus CD4 expression. FACS profile from one representative of three mice is shown.

B. A group of CBA mice (n=4) were i.v. injected with 20 μg of anti-CD4^{PE}. 18, 30, 60 mintues or 22 hours later, the PBL were taken from individual mice and stained with anti-CD3^{FTTC} only. The dot-plots show CD3 versus CD4 expression. FACS profile from one representative of four mice is shown. This experiment was repeated for three times and similar results were obtained.

S-Figure 3. Accumulation of adoptively transferred HY-specific Treg cells in MB49 tumours is profoundly impaired following DTA-1 treatment.

A. CD4⁺CD25⁺ cells purified from Rag^{+/-} Marilyn Thy1.1 mice were adoptively transferred to two groups of Thy1.2⁺ B6 females by i.v. injection (2.6x10⁶/mouse). On the same day, the mice in G1 and G2 were inoculated s.c. with MB49 cells

 $(5x10^5/mouse)$. On day 3 and 8, the mice in G2 were treated with 50 µg of DTA-1 by i.p. injection, and On day 11, dLN, ndLN and MB49 tumours from individual mice in each group were stained with anti-V $\beta6^{PE}$, anti-Thy1.1^{PerCP} and anti-CD4^{APC} followed by intracellular staining of anti-Foxp3^{FITC}. Percentage of donor cells (Thy1.1⁺V $\beta6^+$) of one representative mouse from each group is shown.

B. The absolute numbers of donor Treg cells in dLN of G1 and G2 were calculated by total cell number x % of gated lymphocytes x % of donor cells. One representative mouse from each group is shown.