Supplementary Material

Gastrointestinal microbiome dysbiosis in infant mice alters peripheral CD8$^+$ T cell receptor signaling

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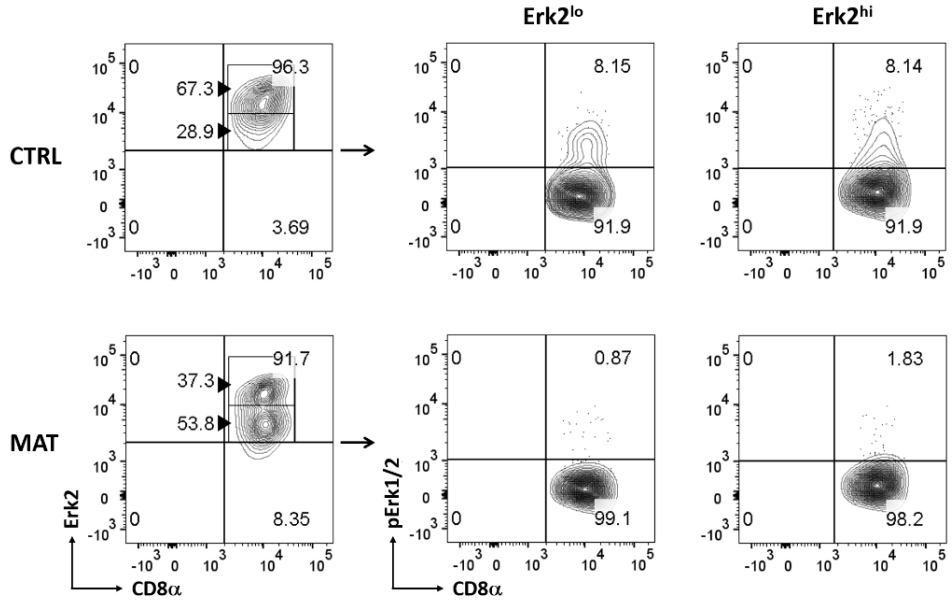
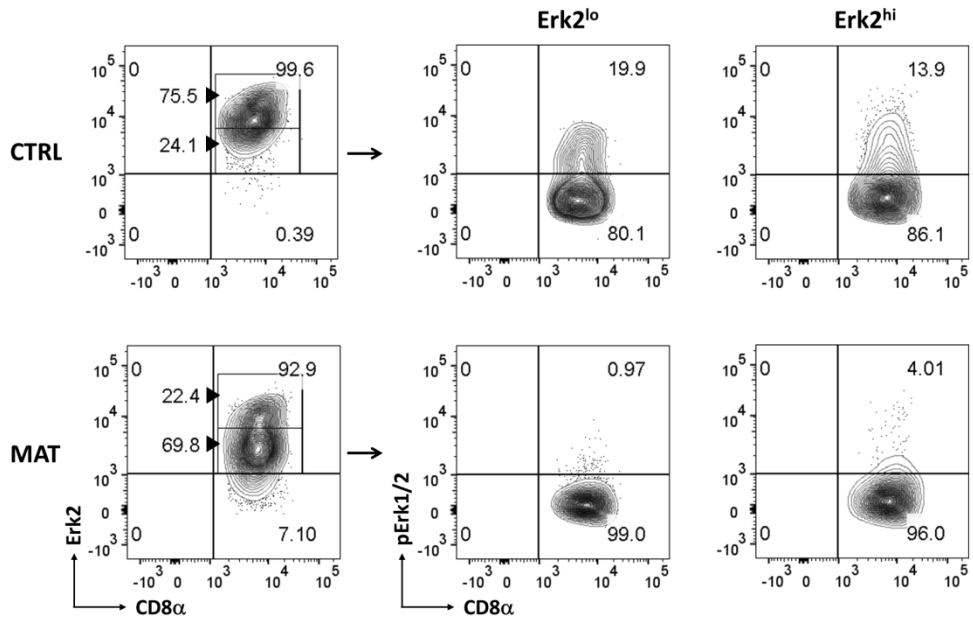
Supplementary Figure 1



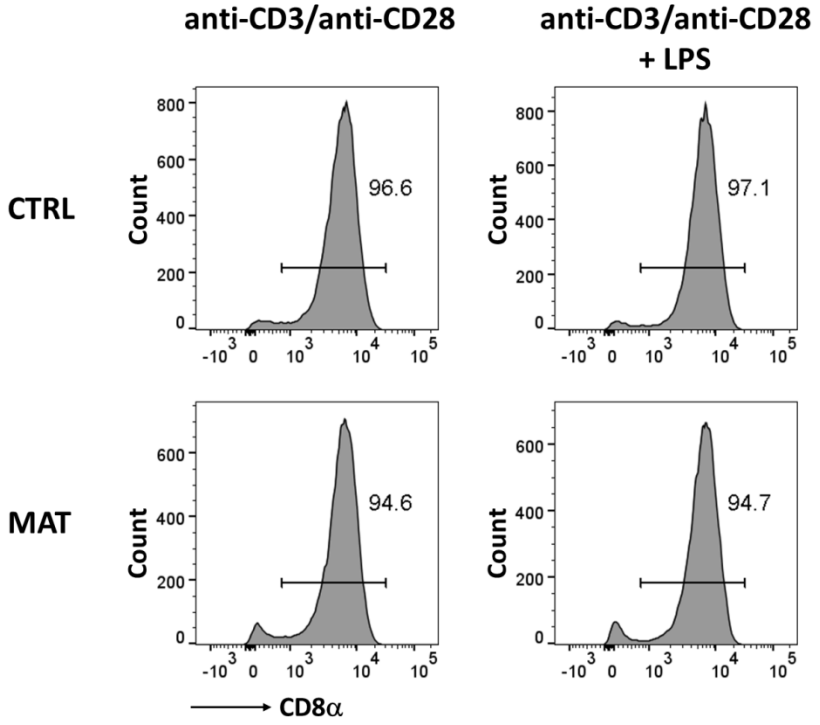
Supplementary Figure 1. MAT effector CD8$^+$ T cells do not sustain phosphorylation of Erk-1/2. Total T cells enriched from the spleens of uninfected dol 15 CTRL and MAT infant mice were stimulated with anti-CD3/anti-CD28 for 72 hours and effector (CD44$^+$) CD8$^+$ T cells were analyzed for the expression of Erk2 and phospho-Erk-1/2 (pErk-1/2) by flow cytometry. The percentage of effector CD8$^+$ T cells expressing low or high levels of Erk2 (Erk2^{lo} and Erk2^{hi}, respectively) and the gating analysis of their corresponding pErk-1/2 expressing populations is shown. Data is representative of two independent experiments.

Supplementary Figure 2



Supplementary Figure 2. Purified MAT effector CD8$^+$ T cells do not sustain phosphorylation of Erk-1/2. Total CD8$^+$ T cells purified from gender-matched pooled spleens of uninfected dol 15 CTRL and MAT infant mice were stimulated with anti-CD3/anti-CD28 for 72 hours and effector (CD44$^+$) CD8$^+$ T cells were analyzed for the expression of Erk2 and phospho-Erk-1/2 (pErk-1/2) by flow cytometry. The percentage of effector CD8$^+$ T cells expressing low or high levels of Erk2 (Erk2^{lo} and Erk2^{hi}, respectively) and the gating analysis of their corresponding pErk-1/2 expressing populations is shown.

Supplementary Figure 3



Supplementary Figure 3. CD8 T cell purity from MAT and CTRL effector T cells generated *in vitro* with or without LPS. Total $CD8^+$ T cells purified from gender-matched pooled spleens of uninfected dol 15 CTRL and MAT infant mice were stimulated with anti-CD3/anti-CD28 with or without *Escherichia coli*-derived LPS (1 μg/ml) for 72 hours. The percentage of $CD8^+$ T cells in these cultures is shown. Data is representative of three independent experiments.