Title: Crucial role of stimulator of interferon genes-dependent signaling in house dust mite extract-induced IgE production

Authors: Hiroki Nunokawa^{1,2}, Yusuke Murakami^{1,3}, Takashi Ishii¹, Tomoya Narita^{1,3}, Haruyuki Ishii², Hajime Takizawa², Naomi Yamashita^{1,3}

Affiliations:

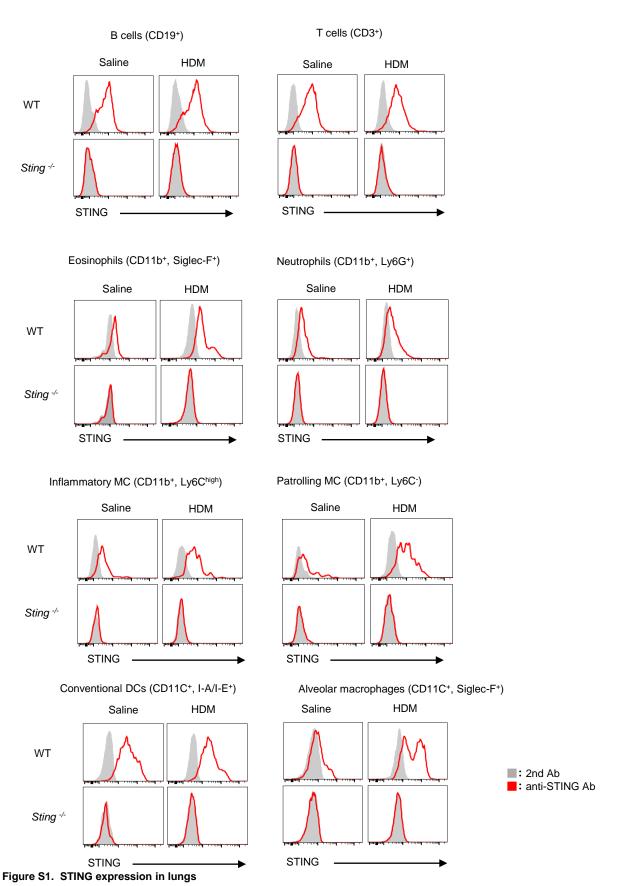
- 1, Research Institute of Pharmaceutical Sciences, Musashino University, Nishitokyo-shi, Tokyo, Japan
- 2, Department of Respiratory Medicine, Kyorin University School of Medicine, Mitaka-shi, Tokyo, Japan
- 3. Faculty of Pharmacy, Department of Pharmaceutical Sciences, Musashino University, Nishitokyo-shi, Tokyo, Japan

Corresponding author: Naomi Yamashita, MD, PhD,

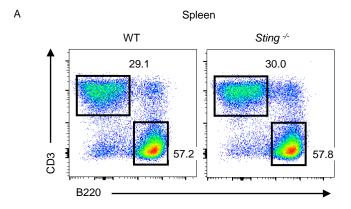
Research Institute of Pharmaceutical Sciences, Musashino University, Nishitokyo-shi, Tokyo, 202-8585, Japan.

Tel: +81-42-468-8647

Email: naoyama@musashino-u.ac.jp



Flow cytometry analysis of STING expression in lungs from WT and $Sting^{-/-}$ mice. B cells, T cells, eosinophils, neutrophils, monocyte subsets, conventional DCs and alveolar macrophages were stained for the markers indicated and analyzed. Gray histograms show 2nd reagent only. Red lines show anti-STING antibody staining. Data are presented from twice independent experiments (n = 2 each). DCs: Dendritic cells, MCs: Monocytes



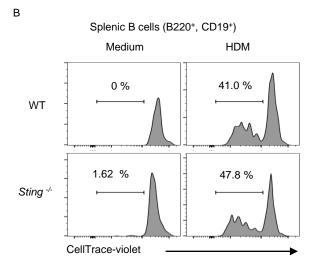


Figure S2. Flow cytometry analysis of B cell proliferation in response to HDM

A, Splenic B cells and T cells were stained for the markers indicated and analyzed. B, Splenic B cells were sorted by indicated markers and treated with the dye. After that, B cells were stimulated with HDM for 3 days and analyzed. The percentages show proliferation level. Data are presented from twice independent experiments (n = 3 each).