Supplementary Figure 1



Supplementary Figure 1. DARA increases functional CD107a expression on NK cells in the killing activity. (A) DARA increased the percentage of CD56⁺CD107a⁺ cells in the mixture of a killing assay against BCBL-1 cells. Antibody- (10 μ g/ml of IgG1 or DARA) coated BCBL-1 cell lines were co-cultured with expanded NK cells from a representative healthy donor at an E:T ratio of 5:1 for 4 hours. The CD107a-expressing NK cells were analyzed by flow cytometry. Anti-CD107a antibody was added at the end of assay. (B) Percentages of CD56⁺CD107a⁺ cells are shown, and were analyzed using the student *t*-test. Data are presented as mean values \pm standard error (SE). * P<0.05, ** P<0.01, *** P<0.001, **** P<0.001 significant levels.

Supplementary Figure 2



Supplementary Figure 2. NK cell lines show killing activity towards K562. (A) The phenotypes of KHYG-1 and N6NK cell lines. (B and C) CFSE-labeling K562 cells were co-cultured with KHYG-1 (CD16⁻) or N6NK (CD16⁺) cells at different E:T ratios for 4 hours. The dead cells were determined using Ghost dye 780 staining. (B) Data of flow cytometric analysis of double Ghost dye 780 and CFSE staining. (C) Percentages of CFSE⁺Ghost Dye⁺ are shown, and were analyzed using the paired *t*-test (n=3). Data are presented as mean values \pm standard error (SE). * P<0.05, ** P<0.01, *** P<0.001.

Supplementary Figure 3



Supplementary Figure 3. DARA demonstrates ADCP in other PELs. MDMs from a representative healthy donor were plated into a well. CFSE-labeled PEL cells were incubated with either DARA or IgG1 antibody for 15 minutes at room temperature. The antibody coating CFSE-labeled PEL cells were added to the well of MDMs (E: T ratios = 0:1, 1:1, 1:5) and further incubated for 2 hours. Phagocytic cells were defined as CD11b⁺CFSE⁺ cells. (A) Data are presented as the % phagocytosis (CD11b⁺CFSE⁺ cells). (B) The data are shown at E: T = 1:5. The data are presented as mean values \pm the standard error (SE). * P<0.05, ** P<0.01, **** P<0.001.