

Supplementary Materials for

Protein kinase D2 is a digital amplifier of T cell receptor-stimulated diacylglycerol signaling in naïve CD8⁺ T cells

María N. Navarro, Carmen Feijoo-Carnero, Alba Gonzalez Arandilla, Matthias Trost, Doreen A. Cantrell*

*Corresponding author. E-mail: d.a.cantrell@dundee.ac.uk

Published 21 October 2014, *Sci. Signal.* **7**, ra99 (2014) DOI: 10.1126/scisignal.2005477

The PDF file includes:

Fig. S1. Quantification of the number of pMHC molecules per RMA-S cell.

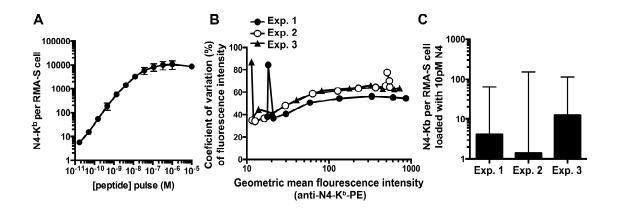


Fig. S1. Quantification of the number of pMHC molecules per RMA-S cell. RMA-S cells were loaded with different amounts of N4 peptide, stained with PE-coupled anti-N4- K^b antibody, and analyzed by flow cytometry. The number of pMHC molecules was determined by measuring the MFI of the PE-coupled anti-N4- K^b antibody with PE-Quantibrite beads. (**A**) The numbers of pMHC molecules presented per RMA cell loaded with the indicated concentrations of N4 peptide. Data are means \pm SD from three independent experiments. (**B**) The coefficients of variation of the MFI data for the different concentrations of N4 peptide used in the each of the three experiments described in (A). (**C**) The distribution (MFI \pm SD) of the number of pMHC molecules presented per RMA cell loaded with 1 pM N4 peptide, which was the lowest concentration of loading peptide for which pPKD2^{Ser873} was detected by flow cytometry.