

**Supplementary Table 1. Data Collection and Refinement Statistics**

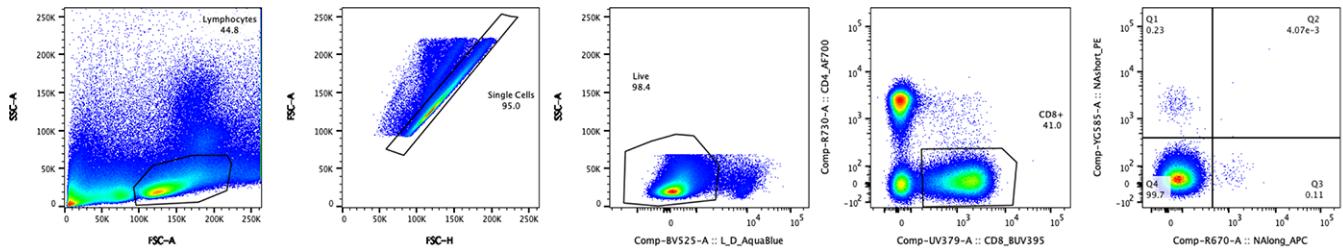
Data Collection Statistics	H-2Db-NA(10)	H-2Db-NA(11)
Temperature	100K	100K
Space group	P2 <sub>1</sub>	P2 <sub>1</sub>
Cell Dimensions (a,b,c)(Å)	68.72, 44.96, 71.77 β = 98.23°	69.01, 44.75, 71.66 β = 98.45°
Resolution (Å)	45.95 – 1.50 (1.53 – 1.50)	45.91 – 1.55 (1.58 – 1.55)
Total number of observations	247455 (12308)	228178 (11040)
Number of unique observations	69128 (3413)	62822 (3067)
Multiplicity	3.6 (3.6)	3.6 (3.6)
Data completeness (%)	99.1 (99.4)	99.5 (99.6)
I/σ <sub>I</sub>	17.6 (2.0)	12.7 (2.0)
R <sub>pim</sub> <sup>a</sup> (%)	2.8 (39.0)	5.6 (43.9)
Refinement Statistics		
Non-hydrogen atoms	3774	3769
Protein	3202	3175
Water	572	594
R <sub>factor</sub> <sup>b</sup> (%)	17.81 (26.52)	18.30 (28.80)
R <sub>free</sub> <sup>b</sup> (%)	20.88 (30.02)	21.25 (31.93)
Rms deviations from ideality		
Bond lengths (Å)	0.013	0.013
Bond angles (°)	1.57	1.58
Ramachandran plot (%)		
Favored	99	99
Allowed	0.79	1.3
Outliers	0	0

<sup>a</sup>R<sub>p.i.m</sub> =  $\sum_{hkl} [1/(N-1)]^{1/2} \sum_i |I_{hkl,i} - \langle I_{hkl} \rangle| / \sum_{hkl} \langle I_{hkl} \rangle$

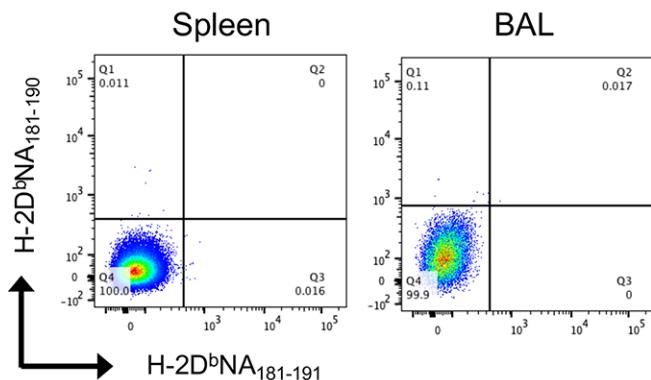
<sup>b</sup> R<sub>factor</sub> =  $\sum_{hkl} ||F_o|| - ||F_c|| / \sum_{hkl} ||F_o||$  for all data except ≈ 5% which were used for R<sub>free</sub> calculation

## Supplemental Fig. 1

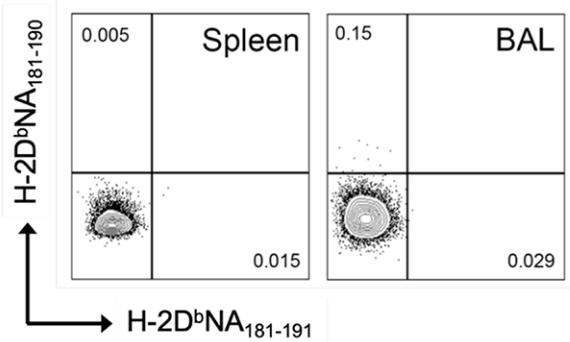
**A**



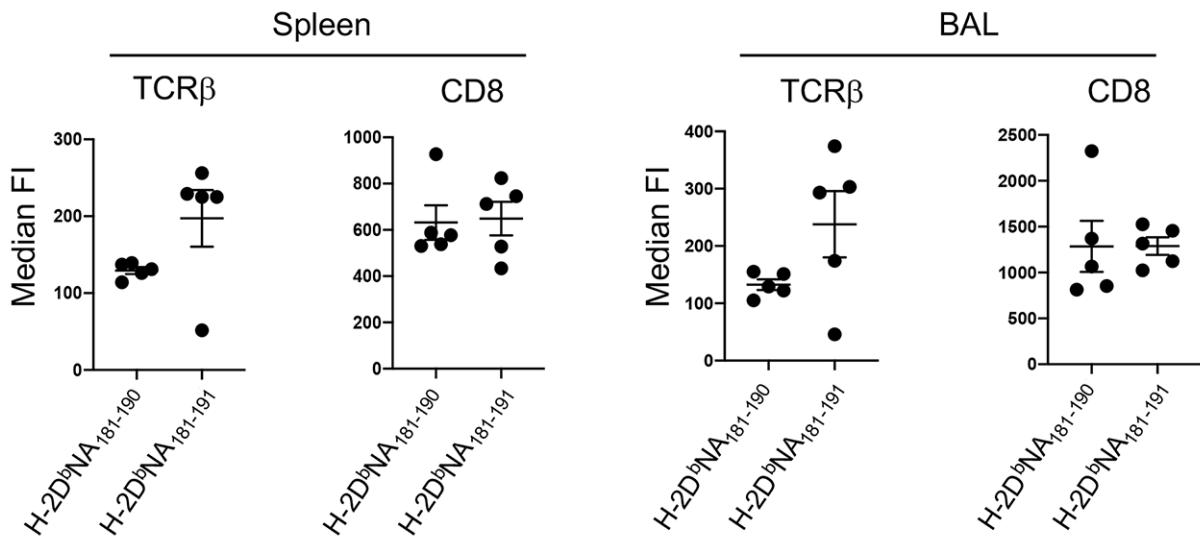
**B**



**C**



**D**



**Supplemental Fig. 1.** (A) Dot plots showing consecutive gating strategy for analysis of tetramer staining on splenic CD8+ T cells. (B) Dot plots showing detection of tetramer staining in spleen and BAL samples stained with all antibodies but no tetramers (FMO control). (C) Tetramer staining on CD4+ T cells from samples shown in Figure 1A. (D) Median fluorescence intensity of TCR $\beta$  and CD8 staining on spleen and BAL samples from tetramer dissociation experiment shown in Figure 2E.