



T Cell Receptor Cross-Reactivity between Similar Foreign and Self Peptides Influences **Naive Cell Population Size and Autoimmunity**

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Table 1 of this paper published in the January 2015 issue contains several errors. The Derp1 peptide comes from the Derp1 protein of Dermatophagoides pteronyssinus not the Der f1 protein of Dermatophagoides farinae. The sequence of the MOG peptide in the I-A^b tetramer used in the study was GWYRSPFSRVV not GWYRSPFSRVVVHLY. The most likely 9 amino acid core register for the OVA2C peptide in the I-A^b tetramer used in the study is VHAAHAEIN not AVHAAHAEI, which is the second most likely. The most likely register has a different number of predicted self peptide homologs than the second most likely one, which changed the x axis value for the OVA2C peptide in Figure 4G. However, neither the level of statistical significance of the correlation between the number of self peptide homologs and the number of naive cells specific for foreign peptides shown in Figure 4G nor the conclusion that "TCR cross-reactivity on self-peptide homologs plays some role in determining the size of MHCII-bound foreign peptide-specific CD4⁺ T cell populations" were changed. Thus, none of these errors affect any of the conclusions of the paper. The authors deeply regret these errors and apologize for them. Corrected versions of the table and figure appear here.

Table 1. Sequence of Peptides in I-A ^b Tetramers	
Peptide	Sequence
ESAT6	QQWN <u>FAGIEAAAS</u> A
FliC	VQNR <u>FNSAITNLG</u> NT
STM1540-3#	VYYTTYAPQAT
STM1540-1#	YTTYAPQATSA
eGFP	HDF <u>FKSAMPEGY</u> VQE
OVA3C#	GH <u>AAHAEINEA</u>
RpIF	<u>FVSPAAHII</u>
OVA2C#	QA <u>VHAAHAEIN</u>
3K	E <u>AQKAKANKA</u> VDKA
Calnexin	LV <u>VKNPAAHHA</u> IS
AasF	VSSPAVQES
1G1W	E <u>AGGALANWA</u> VDSA
LLO#	NEK <u>YAQAYPNVS</u>
Derp1	CQIYPPNVNKI
Cda2	HQ <u>YMTALSNEV</u> VF
PmpG-1	YVDPAAAGG
СТВ	N <u>NKTPHAIAA</u> IS
GP66	DI <u>YKGVYQFKS</u> V
2W	E <u>AWGALANWA</u> VDSA
CD4Ag28m	VE <u>IHRPVPGTA</u>
2W109#	E <u>YWGPLPNWV</u>
MOG#	GW <u>YRSPFSRVV</u>



Immunity **Errata**

