

Lck Availability during Thymic Selection Determines the Recognition Specificity of the T Cell Repertoire

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SUMMARY

Thymic selection requires signaling by the protein tyrosine kinase Lck to generate T cells expressing $\alpha\beta$ T cell antigen receptors (TCR). For reasons not understood, the thymus selects only $\alpha\beta$ TCR that are restricted by major histocompatibility complex (MHC)-encoded determinants. Here, we report that Lck proteins that were coreceptor associated promoted thymic selection of conventionally MHC-restricted TCR, but Lck proteins that were coreceptor free promoted thymic selection of MHC-independent TCR. Transgenic TCR with MHC-independent specificity for CD155 utilized coreceptor-free Lck to signal thymic selection in the absence of MHC, unlike any transgenic TCR previously described. Thus, the thymus can select either MHC-restricted or MHC-independent $\alpha\beta$ TCR depending on whether Lck is coreceptor associated or coreceptor free. We conclude that the intracellular state of Lck determines the specificity of thymic selection and that Lck association with coreceptor proteins during thymic selection is the mechanism by which MHC restriction is imposed on a randomly generated $\alpha\beta$ TCR repertoire.

INTRODUCTION

Antigen receptors on cells of the adaptive immune system must be capable of recognizing existing pathogens as well as new pathogens that will arise in the future. To do so, T and B lymphocytes use gene recombination to randomly generate antigen receptors with hugely diverse recognition specificities (Davis and Bjorkman, 1988). Although generated by the same recombination machinery, antigen receptors on mature T and B lymphocytes recognize fundamentally different types of antigenic

ligands. Antigen receptors on B lymphocytes recognize conformational epitopes on native antigenic proteins and glycolipids, as do antigen receptors on the minor subset of T cells bearing $\gamma\delta$ T cell antigen receptors (TCR) (Chien and Königshofer, 2007). However, antigen receptors on the major subset of T cells bearing $\alpha\beta$ TCR do not recognize conformational antigenic epitopes but instead recognize linear peptide fragments of antigenic proteins bound to major histocompatibility complex (MHC)-encoded determinants, the feature of antigen recognition referred to as MHC restriction (Davis and Bjorkman, 1988). MHC restriction is unique to $\alpha\beta$ TCR and allows identification of cells containing intracellular pathogens, foreign proteins, or genetic mutations. As a result, MHC restriction is critical for T lymphocyte recognition and function, but how it is imposed on a randomly generated $\alpha\beta$ TCR repertoire remains a major unsolved problem.

Two different explanations have been proposed for the exclusive expression of MHC-restricted $\alpha\beta$ TCR on mature T cells. The germline model of MHC restriction proposes that MHC restriction is intrinsic to germline-encoded $\alpha\beta$ TCR structural elements (Feng et al., 2007; Garcia et al., 2009; Marrack et al., 2008; Scott-Browne et al., 2009). According to the germline model, specific amino acids in the complementary determining regions (CDR) 1 and 2 of TCR α and TCR β have been conserved during evolution because they contact MHC chains and impose MHC specificity on $\alpha\beta$ TCR recognition. Consequently, $\alpha\beta$ TCR are limited by germline imposed structural constraints to be MHC-specific and to bind only to MHC-dependent ligands, with the exception of a few exceedingly rare $\alpha\beta$ TCR that cross-reactively bind an MHC-independent ligand with very low affinity (Bard et al., 1989; Hanada et al., 2011; Rao et al., 1984). In contrast to the germline model, the selection model of MHC restriction proposes that MHC restriction is the result of TCR-signaled thymic selection and is not an intrinsic feature of $\alpha\beta$ TCR structure (Collins and Riddle, 2008; Tikhonova et al., 2012; Van Laethem et al., 2007, 2012). According to the selection model, $\alpha\beta$ TCR are randomly generated in the thymus so that preselection CD4⁺CD8⁺ (double positive, DP) thymocytes express $\alpha\beta$ TCR

with a huge diversity of recognition specificities. However, only MHC-restricted $\alpha\beta$ TCR signal DP thymocytes to undergo thymic selection because CD4 and CD8 coreceptors on DP thymocytes sequester the signaling protein tyrosine kinase (PTK) p56^{lck} (Lck) (Haughn et al., 1992) so that only $\alpha\beta$ TCR with the same MHC specificity as either CD4 or CD8 coreceptor proteins can access Lck (Doyle and Strominger, 1987; Normont et al., 1988).

Lck is a Src family PTK that is expressed in all T lineage cells and inserts in the inner leaf of their plasma membrane as a result of being myristoylated or palmitoylated at its amino terminus (Paige et al., 1993). Membrane Lck initiates TCR signal transduction by first phosphorylating signaling motifs in the TCR complex and then phosphorylating ZAP-70 PTK molecules that are recruited into the TCR complex by the phosphorylated signaling motifs (Abraham et al., 1991; Acuto and Cantrell, 2000; Chan et al., 1992; Gascoigne and Palmer, 2011; Nika et al., 2010). Although p59^{fyn} (Fyn) can also initiate TCR signaling, Lck is the PTK that initiates TCR signaling in the vast majority of developing thymocytes (Palacios and Weiss, 2004). Notably, membrane Lck has two cytosolic cysteines that mediate noncovalent interactions with cysteines in the cytosolic tails of CD4 and CD8 (Rudd et al., 1988; Shaw et al., 1989; Turner et al., 1990; Veillette et al., 1988). As a result, membrane Lck can be either coreceptor associated or coreceptor free if it is not bound to CD4 or CD8. In preselection DP thymocytes, membrane Lck is overwhelmingly coreceptor associated because of the large quantities of both CD4 and CD8 coreceptors expressed, with the result that $\alpha\beta$ TCR signaling in DP thymocytes is only initiated by coengagement of TCR with coreceptor proteins (Wiest et al., 1996).

The selection model of MHC restriction predicts that MHC-independent $\alpha\beta$ TCR would signal thymic selection and generate an MHC-independent peripheral $\alpha\beta$ TCR repertoire if preselection thymocytes contained coreceptor-free Lck instead of coreceptor-associated Lck. This prediction was tested in Quad-KO mice that were simultaneously both MHC-deficient (*b2m*^{-/-}*H2-Ab1*^{-/-}) and coreceptor-deficient (*Cd4*^{-/-}*Cd8*^{-/-}) because Quad-KO thymocytes contained Lck that was necessarily coreceptor free (Van Laethem et al., 2007). Thymocytes in Quad-KO mice were found to be signaled in vivo and many thymocytes were signaled intensely enough to undergo clonal deletion. Bcl-2 transgene (Bcl-2^{Tg}) expression prevented clonal deletion in Quad-KO mice and resulted in large numbers of peripheral T cells expressing $\alpha\beta$ TCR with MHC-independent recognition specificities (Tikhonova et al., 2012; Van Laethem et al., 2007). Two MHC-independent $\alpha\beta$ TCR were cloned from Quad-KO.Bcl-2Tg mice and displayed antibody-like recognition properties that fundamentally differed from conventionally MHC-restricted TCR. The two MHC-independent $\alpha\beta$ TCR did not recognize peptide/MHC (pMHC) complexes but instead bound to conformational epitopes on the membrane protein CD155, and they did so independently of MHC with an affinity that resembled that of primary antibodies (Tikhonova et al., 2012).

The generation in Quad-KO mice of peripheral T cells bearing MHC-independent $\alpha\beta$ TCR fulfilled predictions of the thymic selection model and contradicted requirements of the germline model. Nevertheless, central issues related to MHC-independent thymic selection were not resolved, including the ligand-dependence of thymic selection by coreceptor-free Lck.

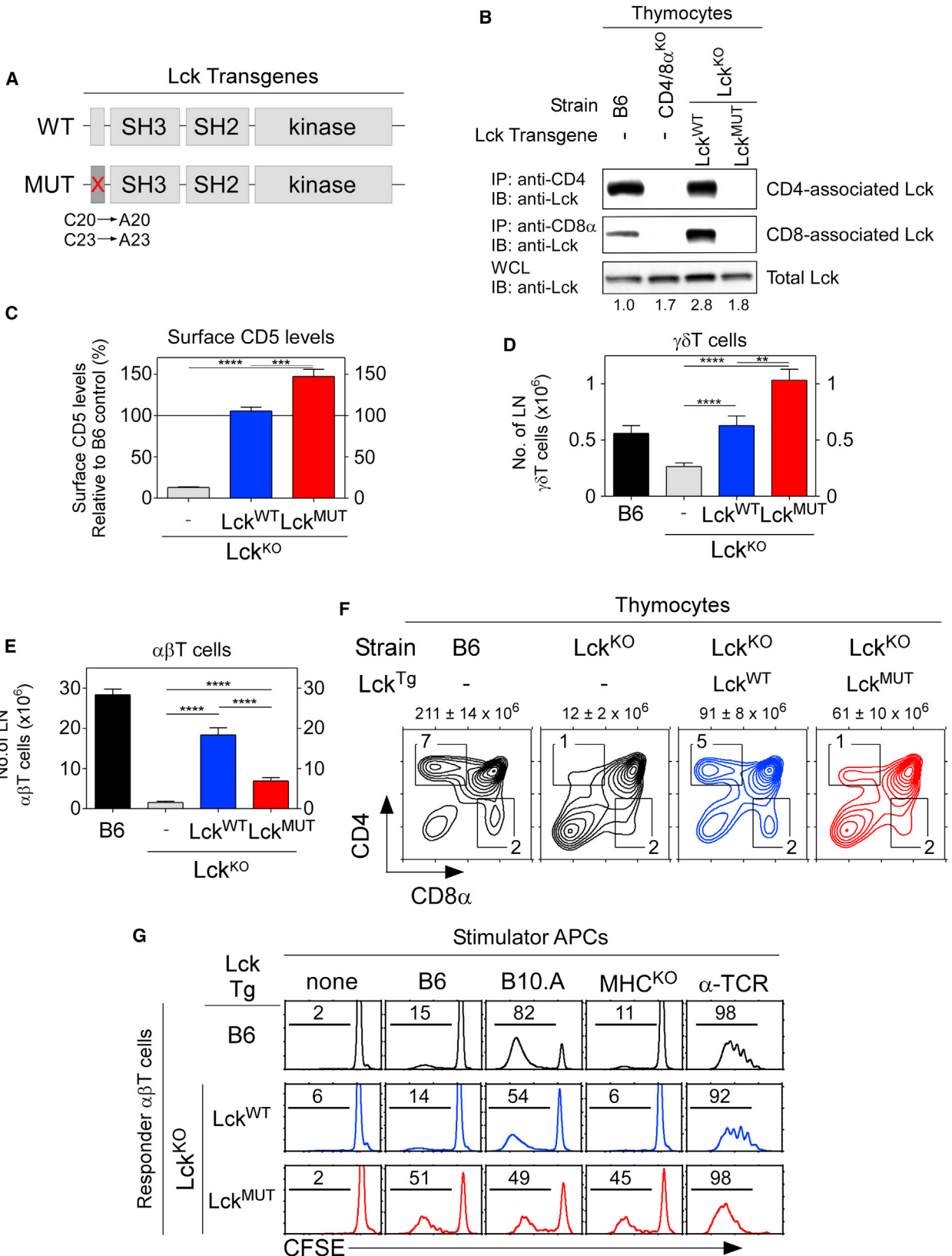
We undertook the present study to evaluate the impact of Lck availability on thymic selection and to assess the differentiation of MHC-independent $\alpha\beta$ TCR in the thymus. We now report that generation of an MHC-independent $\alpha\beta$ TCR repertoire only required mutation of the two cysteines in Lck's N-terminal region that promote association with CD4 and CD8 coreceptors in the thymus. We also demonstrate that, unlike coreceptor-associated Lck that promoted thymic selection of MHC-restricted $\alpha\beta$ TCR, coreceptor-free Lck preferentially promoted thymic selection of MHC-independent $\alpha\beta$ TCR. Moreover we characterize thymic selection of two MHC-independent transgenic $\alpha\beta$ TCR specific for different epitopes on the self-protein CD155. Thymic selection of these two MHC-independent $\alpha\beta$ TCR was signaled by coreceptor-free Lck upon engagement of CD155 ligands, even in MHC-deficient mice. Thus, this study demonstrates that thymic selection of either MHC-independent or MHC-restricted $\alpha\beta$ TCR depends on the presence or absence of coreceptor-free Lck, and it is Lck's association with CD4/CD8 coreceptor proteins during thymic selection that is responsible for thymic selection of an exclusively MHC-restricted $\alpha\beta$ TCR repertoire.

RESULTS

Signaling and Selection in the Thymus by Coreceptor-free Lck

MHC restriction is the cardinal feature of $\alpha\beta$ TCR selected in the thymus but its basis remains uncertain. The Lck sequestration model of thymic selection (Van Laethem et al., 2007) proposes that MHC restriction is not intrinsic to $\alpha\beta$ TCR structure but is the result of thymic selection (Figure S1 available online). According to this hypothesis, $\alpha\beta$ TCR specific for MHC-independent ligands fail to transduce positive selection signals because Lck is bound to the cytosolic tails of CD4/CD8 coreceptor proteins that sequester Lck away from all TCR except those that engage peptide-MHC (pMHC) ligands together with CD4/CD8 coreceptor proteins (Figure S1). A prediction of the Lck sequestration model is that coreceptor-free Lck would promote signaling by MHC-independent TCR in the thymus and selection of MHC-independent $\alpha\beta$ T cells in the periphery.

To determine if coreceptor-free Lck promoted thymic selection of MHC-independent $\alpha\beta$ TCR, we constructed hCD2-driven transgenes encoding either wild-type Lck proteins (Lck^{wt}) or mutant Lck proteins (Lck^{mut}) whose cysteines at positions 20 and 23 were converted to alanines to impair binding to coreceptor proteins (Figure 1A) (Turner et al., 1990). Both Lck transgenes were introduced into Lck^{KO} (*Lck*^{-/-}) mice to generate mice referred to as Lck^{wt}Tg and Lck^{mut}Tg mice, respectively. Immunoblotting of thymocyte lysates revealed that Lck^{mut}Tg thymocytes contained Lck in amounts that were intermediate between B6 and Lck^{wt}Tg mice (Figure 1B). Immunoprecipitation of thymocyte lysates with coreceptor-specific antibodies further revealed that Lck^{mut} proteins bound neither CD4 nor CD8, unlike Lck^{wt} proteins that bound to both CD4 and CD8 (Figure 1B). Nevertheless, CD5 surface expression was restored in Lck^{KO} mice by both Lck^{mut} and Lck^{wt} transgenic proteins, with Lck^{mut}Tg thymocytes expressing higher CD5 levels as an indication of stronger in vivo signaling by Lck^{mut} Tg transgenic proteins (Figure 1C).



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To assess the function of Lck transgenic proteins in thymic selection, we examined $\gamma\delta$ and $\alpha\beta$ T cell generation (Figures 1D and 1E). $\gamma\delta$ TCR arise at the DN stage of thymocyte development when CD4/CD8 coreceptors are not expressed, so Lck^{wt} and Lck^{mut} proteins would both be available to $\gamma\delta$ TCR. Indeed $\gamma\delta$ T cell generation that was reduced in Lck^{KO} mice was restored by both Lck^{mut} and Lck^{wt} transgenic proteins, although significantly more $\gamma\delta$ T cells were present in Lck^{mut} transgenic mice, another indication of stronger in vivo signaling by Lck^{mut} transgenic proteins (Figure 1D).

Unlike $\gamma\delta$ TCR, $\alpha\beta$ TCR are expressed at the DP stage of thymocyte differentiation when CD4/CD8 coreceptor proteins are present that bind Lck. Interestingly, $\alpha\beta$ T cell generation that was abrogated in Lck^{KO} mice was only poorly restored by Lck^{mut} proteins compared to Lck^{wt} proteins, with significantly fewer peripheral $\alpha\beta$ T cells and SP thymocytes in Lck^{mut}Tg mice (Figures 1E and 1F). Impaired $\alpha\beta$ T cell generation by Lck^{mut} proteins contrasted markedly with robust $\gamma\delta$ T cell generation in the same mice and raised the possibility that $\alpha\beta$ T cells might have been signaled so strongly by coreceptor-free Lck that they underwent clonal deletion. However, introduction of the prosurvival Bcl-2^{Tg} to impair in vivo clonal deletion (Pobezinsky et al., 2012) revealed that Lck^{mut}Tg.Bcl-2^{Tg} mice still contained fewer $\alpha\beta$ T cells and SP thymocytes than Lck^{wt}Tg.Bcl-2^{Tg} mice (Figures S2A and S2B). Consequently, coreceptor-free Lck^{mut} proteins signaled the generation of fewer $\alpha\beta$ T cells than coreceptor-associated Lck proteins despite stronger in vivo signaling.

We then assessed the specificity of peripheral $\alpha\beta$ T cells generated in Lck^{mut}Tg mice in in vitro mixed lymphocyte reactions. Unlike B6 and Lck^{wt}Tg $\alpha\beta$ T cells that only reacted against MHC-expressing stimulator cells, Lck^{mut} $\alpha\beta$ T cells reacted vigorously against both MHC-deficient and MHC-expressing stimulator cells (Figure 1G). That is, $\alpha\beta$ T cells generated by coreceptor-free Lck in Lck^{mut}Tg mice expressed $\alpha\beta$ TCR that recognized ligands independently of MHC on MHC-deficient stimulator cells that were also present on MHC-expressing stimulator cells. These results demonstrate that coreceptor-free Lck during thymic selection is all that is required to generate an MHC-independent $\alpha\beta$ TCR repertoire.

Signaling of MHC-Independent $\alpha\beta$ T Cells in the Thymus

To further characterize MHC-independent $\alpha\beta$ TCR selection and signaling in the thymus, we examined thymocytes from so-called Quad^{KO} mice that were MHC-deficient as well as deficient in both CD4 and CD8 coreceptors (*Cd4*^{-/-}*Cd8*^{-/-}*b2m*^{-/-}*H2-Ab1*^{-/-}) (Van Laethem et al., 2007). Endogenous Lck proteins in Quad^{KO} thymocytes were necessarily coreceptor free because coreceptor proteins were not expressed. As indicated by CD5 surface

expression levels, coreceptor-deficient Quad^{KO} thymocytes were strongly signaled in vivo whereas coreceptor-sufficient MHC^{KO} thymocytes were not (Figure 2A). Importantly, MHC-independent in vivo signaling of Quad^{KO} thymocytes required $\alpha\beta$ TCR and Lck proteins, as deficiency in TCR α , RAG2, or Lck reduced or abolished CD5 upregulation (Figure 2A).

Lck Sequestration by Coreceptor Proteins on Developing Thymocytes

Because MHC-independent TCR signaling of Quad^{KO} thymocytes was transduced by coreceptor-free Lck, we wished to determine biochemically if CD4/CD8 coreceptors, by binding Lck, would sequester Lck away from $\alpha\beta$ TCR. To do so, we introduced hCD2-driven CD4 transgenes encoding either full-length (444) or tailless (44T) CD4 proteins (Van Laethem et al., 2007) into Quad^{KO} mice to generate 444.Quad^{KO} and 44T.Quad^{KO} mice whose endogenous Lck proteins bound to 444 but not 44T proteins as revealed by anti-CD4 immunoprecipitation of thymocyte lysates (Figures 2B and 2C). Most importantly, endogenous Lck could also be immunoprecipitated by anti-TCR β but only in thymocytes in which Lck was not bound to CD4 coreceptor proteins (Figure 2C), biochemically documenting that Lck binding to CD4 prevented Lck association with TCR.

Lck Sequestration Impairs Coreceptor-Independent TCR Signaling

To determine the impact of Lck sequestration on thymocyte signaling, we compared TCR signaling in 444.Quad^{KO} and 44T.Quad^{KO} thymocytes. By in vitro assessment, 44T.Quad^{KO} thymocytes with coreceptor-free Lck could be triggered to mobilize intracellular calcium by anti-TCR crosslinking alone (Figure 2D, top), whereas 444.Quad^{KO} thymocytes lacking coreceptor-free Lck required anti-TCR/CD4 cocrosslinking (Figure 2D, bottom). These in vitro results demonstrated that Lck sequestration markedly impaired coreceptor-independent TCR signaling. By in vivo assessment, TCR signaling in MHC-deficient thymi was greater in 44T.Quad^{KO} than 444.Quad^{KO} thymocytes as revealed by dramatically higher CD5 expression, greater TCR ζ phosphorylation, and greater amounts of TCR-associated ZAP-70 proteins (Figures 2E and 2F). These in vivo results reveal that Lck sequestration markedly impairs TCR signaling in MHC-deficient thymi.

Thymic Selection of MHC-Independent TCR

To better understand $\alpha\beta$ TCR signaling of thymic selection in the absence of MHC, we generated TCR transgenes encoding $\alpha\beta$ TCR from two T cell clones that were derived from Quad^{KO}Bcl-2^{Tg} mice with specificity for an MHC-independent ligand

Figure 1. Effect of Coreceptor-free and Coreceptor-Associated Lck on Thymic Selection

(A) hCD2-driven transgenic constructs encoding wild-type (top) and C20A/C23A mutant (bottom) Lck proteins.

(B) Abrogation of Lck binding to coreceptor proteins. Thymocyte lysates were precipitated with anti-CD4 or anti-CD8 α and blotted for Lck. Numbers indicate total Lck band intensities relative to B6 (set equal to 1.0). Data represent four experiments.

(C) CD5 levels on thymocytes relative to control B6 mice (mean \pm SE, n = 7 mice/group).

(D and E) Numbers of $\gamma\delta$ (D) and $\alpha\beta$ (E) LN T cells (mean \pm SE, n = 9 mice/group).

(F) Numbers above profiles indicate thymus cellularity (mean \pm SE, n = 10 mice/group).

(G) Ligand specificity of $\alpha\beta$ T cells in mixed lymphocyte reactions as assessed by proliferation-induced CFSE dye dilution on day 4. Numbers represent percentage of cells that underwent at least one division. Data represent 3 experiments.

****p < 0.0001; ***p < 0.001; **p < 0.01. See also Figures S1 and S2.

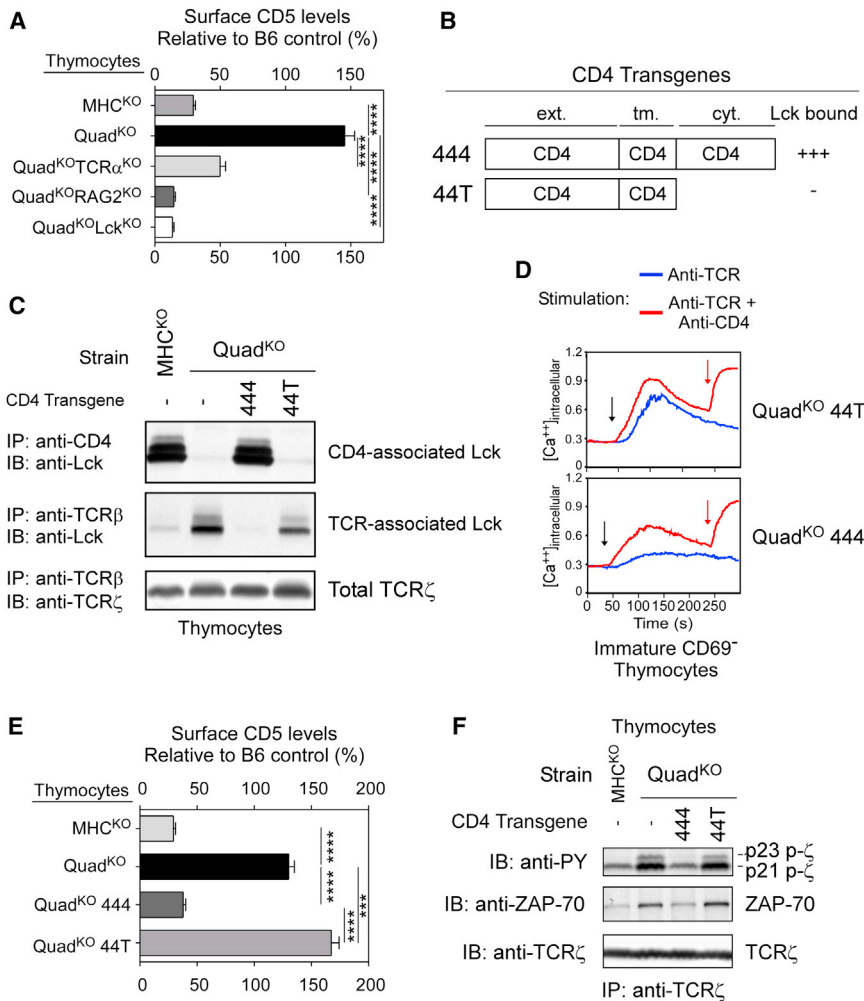


Figure 2. Lck Sequestration and TCR Signaling in the Thymus

(A) CD5 on thymocytes relative to B6 (set at 100%). Mean ± SE, n = 3 mice/group.

(B) hCD2-driven transgenic constructs encoding wild-type (444) and tailless (44T) CD4 proteins named for their extra-cellular, transmembrane, and cytosolic domains. (C) Lck is sequestered away from TCR on thymocytes by the cytosolic tail of coreceptor proteins. Thymocyte lysates were precipitated by anti-CD4 or anti-TCRβ and blotted for Lck or TCRζ. Data represent three experiments.

(D) In vitro TCR induced calcium mobilization. Thymocytes were loaded at 31°C with Indo-1 and coated with biotinylated anti-TCR (5 μg/ml) either alone or together with biotinylated anti-CD4 (1 μg/ml). Crosslinking by avidin (black arrow) and addition of ionomycin (red arrow) are indicated. Data represent two experiments.

(E) In vivo signaling of thymocytes. CD5 on thymocytes is shown relative to control B6 mice (mean ± SE, n = 10 mice/group).

(F) In vivo signaling of thymocytes assessed biochemically. Lysates were precipitated with anti-TCRζ and blotted for phospho-zeta, ZAP70, or TCRζ. Data represent two experiments.

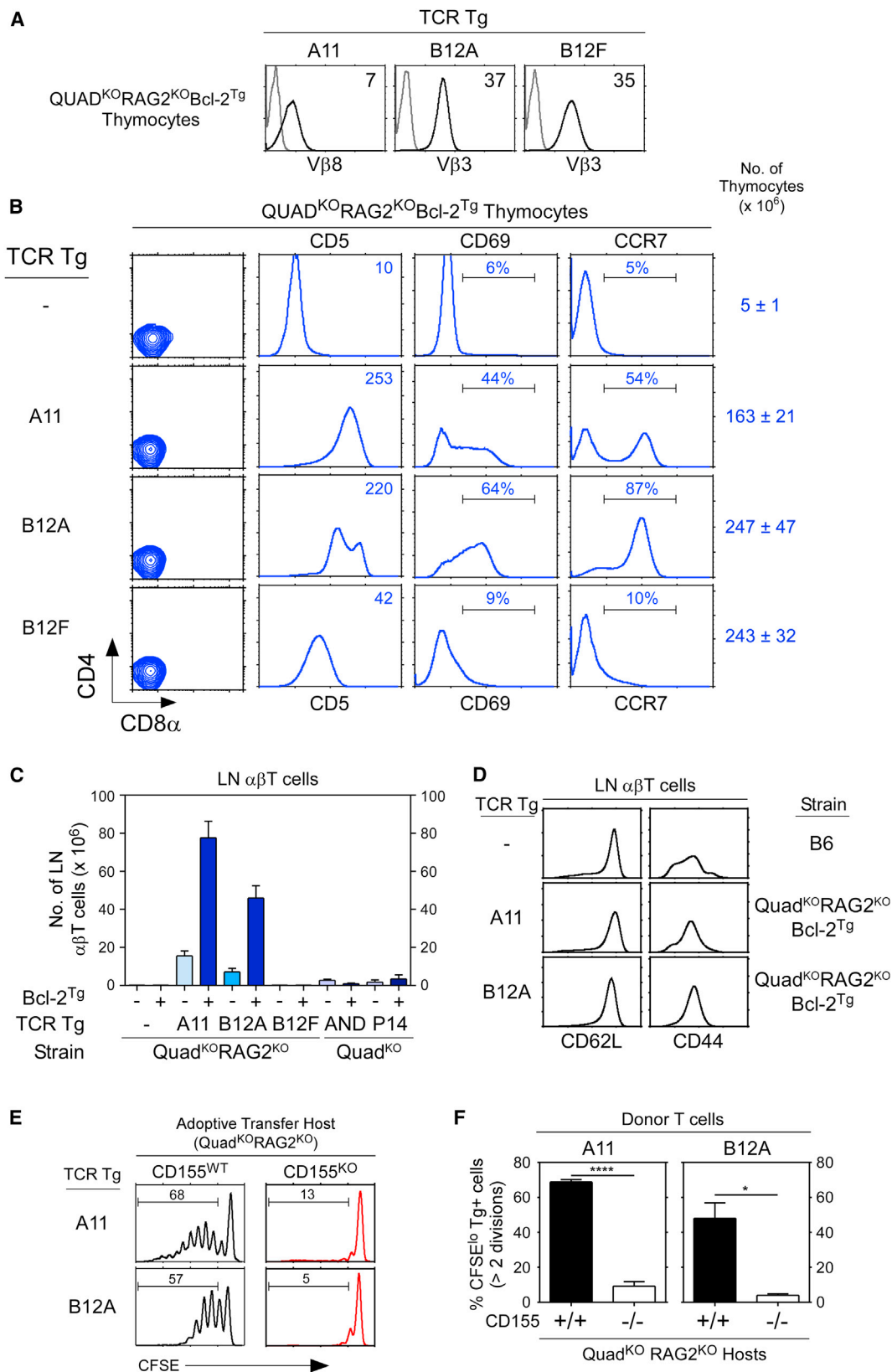
****p < 0.0001; ***p < 0.001.

(Figure S3A) (Tikhonova et al., 2012). The T-hybridoma cells were derived from Quad^{KO} mice expressing the Bcl-2^{Tg} that protected strongly signaled MHC-independent αβTCR from clonal deletion (Van Laethem et al., 2007). Three hCD2-driven TCR transgenes were constructed from the A11 and B12 T-hybridoma cell lines described previously (Tikhonova et al., 2012) and are referred to as A11, B12A, and B12F. The A11 TCR was TCR-Vα8Vβ8, whereas B12A and B12F TCR were identically TCR-Vα4Vβ3 but with different VαJα-encoded CDR3 sequences (Figure S3B). A11 and B12A TCR recognized different conformational epitopes on the self-ligand CD155, the murine analog of the human polio virus receptor, and bound to CD155 proteins independently of MHC with affinities that were unusually high for αβTCR (Tikhonova et al., 2012). However, the B12F TCR displayed no known ligand specificity (Figure S3C). In vivo, all three TCR transgenes (A11, B12A, and B12F) induced TCR expression on developing thymocytes (Figure 3A).

To determine if these TCR could signal positive selection in the absence of MHC, we expressed them in the same host mice from which they were originally derived, i.e., Quad^{KO}Bcl-2^{Tg} mice, except that transgenic host mice were additionally

and CCR7 that indicated positive selection signaling, whereas B12F TCR only minimally upregulated CD5 expression consistent with pre-TCR signaling (Figure 3B). The inability of B12F TCR to markedly upregulate CD5, CD69, and CCR7 expression in Quad^{KO} thymocytes indicated that coreceptor-free Lck only transduced positive selection signals by TCR that engaged an intrathymic ligand. Indeed, coreceptor-free Lck in Quad^{KO} thymocytes did not transduce positive selection signals by either MHC-II restricted AND TCR or MHC-I restricted P14 TCR whose MHC ligands were not expressed in the thymus (Figure S3D).

Because A11 and B12A TCR signaled positive selection in the absence of MHC, A11 and B12A TCR transgenic mice should contain mature αβT cells in the MHC-deficient periphery (Figure 3C). In fact, A11 and B12A TCR generated peripheral LNT cells in huge numbers (50–80 million) in transgenic host mice expressing Bcl-2^{Tg} and in substantial numbers (10–20 million) without Bcl-2^{Tg} (Figure 3C). The LNT cells generated by A11 and B12A TCR displayed a CD44^{lo}CD62L^{hi} naive phenotype, indicating that their huge numbers were due to extensive thymic selection and not to extensive peripheral expansion (Figure 3D). In contrast, MHC-restricted AND and P14 TCR did not generate



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mature T cells in MHC- and coreceptor-deficient mice (Figure 3C).

To document that A11 and B12A transgenic TCR displayed the same ligand specificity as the T-hybridomas from which they were derived (Tikhonova et al., 2012), we examined the ability of A11 and B12A TCR to trigger lymphopenia-induced T cell proliferation upon CD155 recognition. A11 and B12A LNT cells from transgenic host mice that were not lymphopenic were labeled with the intracellular dye CFSE and then transferred into lymphopenic host mice (Quad^{KO}Rag^{KO}) that either expressed or lacked CD155 (Figures 3E and 3F). Assessed 1 week after transfer, both A11 and B12A T cells proliferated vigorously in CD155^{+/+} but not CD155^{-/-} lymphopenic hosts (Figures 3E and 3F), documenting that both transgenic TCR recognized CD155.

We conclude that A11 and B12A TCR transgenes encode MHC-independent and CD155-specific TCR that are uniquely capable of signaling positive selection in thymi that lack MHC expression, unlike any $\alpha\beta$ TCR previously identified.

Identification of the Thymic-Selecting Ligand

Having discovered transgenic $\alpha\beta$ TCR that signaled positive selection in MHC-deficient thymi, we wished to identify the selecting ligand responsible. Because CD155 was recognized by both A11 and B12A TCR and was a self-protein in host transgenic mice, we expressed A11 and B12A TCR in transgenic host mice in which thymocytes would or would not encounter CD155. Interestingly, A11 and B12A TCR signaled positive selection only in CD155^{+/+} but not CD155^{-/-} transgenic host mice as indicated by CD5, CD69, and CCR7 upregulation (Figure 4A) and by thymocyte differentiation into mature CD69⁺CCR7⁺ cells (Figure 4B). Moreover, A11 and B12A TCR generated peripheral T cells only in CD155^{+/+} transgenic host mice, whereas CD155^{-/-} transgenic host mice were devoid of peripheral T cells (Figure 4C). Thus CD155 is the thymic selecting ligand for both A11 and B12A TCR.

It was then important to determine which cells in the thymus expressed CD155. CD155 was expressed by all elements in the thymus (i.e., cortical thymic epithelial cells [cTECs], medullary thymic epithelial cells [mTECs], and thymocytes) but thymocytes expressed the most (Figure 4D). To determine which thymic cellular element(s) induced positive selection signaling, we constructed four different mixed bone marrow (bm) chimeras in which CD155-deficient A11 thymocytes only encountered CD155 on other cells in the thymus. All four groups of chimeras were constructed by injecting equal mixtures of donor bm stem cells from A11 and B12F transgenic mice into irradi-

ated Quad^{KO}Rag^{KO} host mice. LNT cells in chimeric host mice were examined 8–10 weeks later (Figure 4E). It should be appreciated that B12F thymocytes could not contribute to peripheral T cell numbers that resulted entirely from positive selection by A11 TCR. We found the greatest numbers of peripheral T cells resulted from CD155 expressed on both bm-derived elements and TEC (Figure 4E, group 1), followed by CD155 expressed only on bm-derived cells (Figure 4E, group 3), followed by CD155 expressed only on TEC (Figure 4E, group 2). Thus the number of peripheral A11 T cells corresponded to the overall amount of CD155 expressed in the thymus and did not correspond to CD155 expression on any particular thymic element.

Because ligands on lymphoid elements in the thymus have been shown to positively select innate-like $\alpha\beta$ T cells that are characterized by expression of the transcription factor PLZF (Constantinides and Bendelac, 2013; Lee et al., 2011), we assessed CD155-selected $\alpha\beta$ T cells for expression of *Zbtb16*, the gene that encodes PLZF, and for *Sox13*, a gene expressed by many $\gamma\delta$ -lineage T cells (Melichar et al., 2007). However, CD155-selected A11 and B12A T cells neither expressed *Zbtb16* nor *Sox13* (Figure S4).

We conclude that CD155 is the thymic-selecting ligand for A11 and B12A TCR, and A11 and B12A peripheral T cells are neither innate-like nor $\gamma\delta$ -like T cells.

Effect of Coreceptor Expression on MHC-Independent Selection

Our results demonstrated that $\alpha\beta$ TCR can engage MHC-independent ligands in the thymus and can utilize coreceptor-free Lck to signal positive selection. We then asked if thymocyte expression of CD4 and CD8 coreceptor proteins would affect selection signaling by MHC-independent transgenic TCR.

We introduced A11, B12A, and B12F TCR transgenes into coreceptor-sufficient (*Cd4^{+/+}Cd8^{+/+}*) mice that were MHC^{KO}RAG^{KO}Bcl-2^{Tg} (Figure 5A). Contrary to our expectation, A11 and B12A transgenic TCR did signal positive selection in coreceptor-sufficient mice as indicated by marked upregulation of CD5, CD69, and CCR7 (Figure 5A). Indeed, A11 and B12A TCR signaled coreceptor-sufficient thymocytes to differentiate into SP thymocytes that were mostly CD4SP (Figure 5A). MHC-independent positive selection signaling by A11 and B12A TCR also generated peripheral LN T cells in huge numbers (70–100 million) with Bcl-2^{Tg} and in substantial numbers (10–20 million) without Bcl-2^{Tg} (Figure 5B). In contrast, AND and P14 MHC-restricted TCR did not generate T cells in MHC-deficient mice despite CD4/CD8 coreceptor expression (Figure 5B).

Figure 3. Thymic Selection of MHC-Independent $\alpha\beta$ TCR in the Absence of Both MHC and Coreceptors

(A) TCR expression on thymocytes from transgenic host mice (Quad^{KO}RAG2^{KO}Bcl-2^{Tg}). Black lines indicate TCR expression, gray lines indicate control staining. Numbers in panels indicate mean fluorescence intensity (MFI). Data represent five experiments.

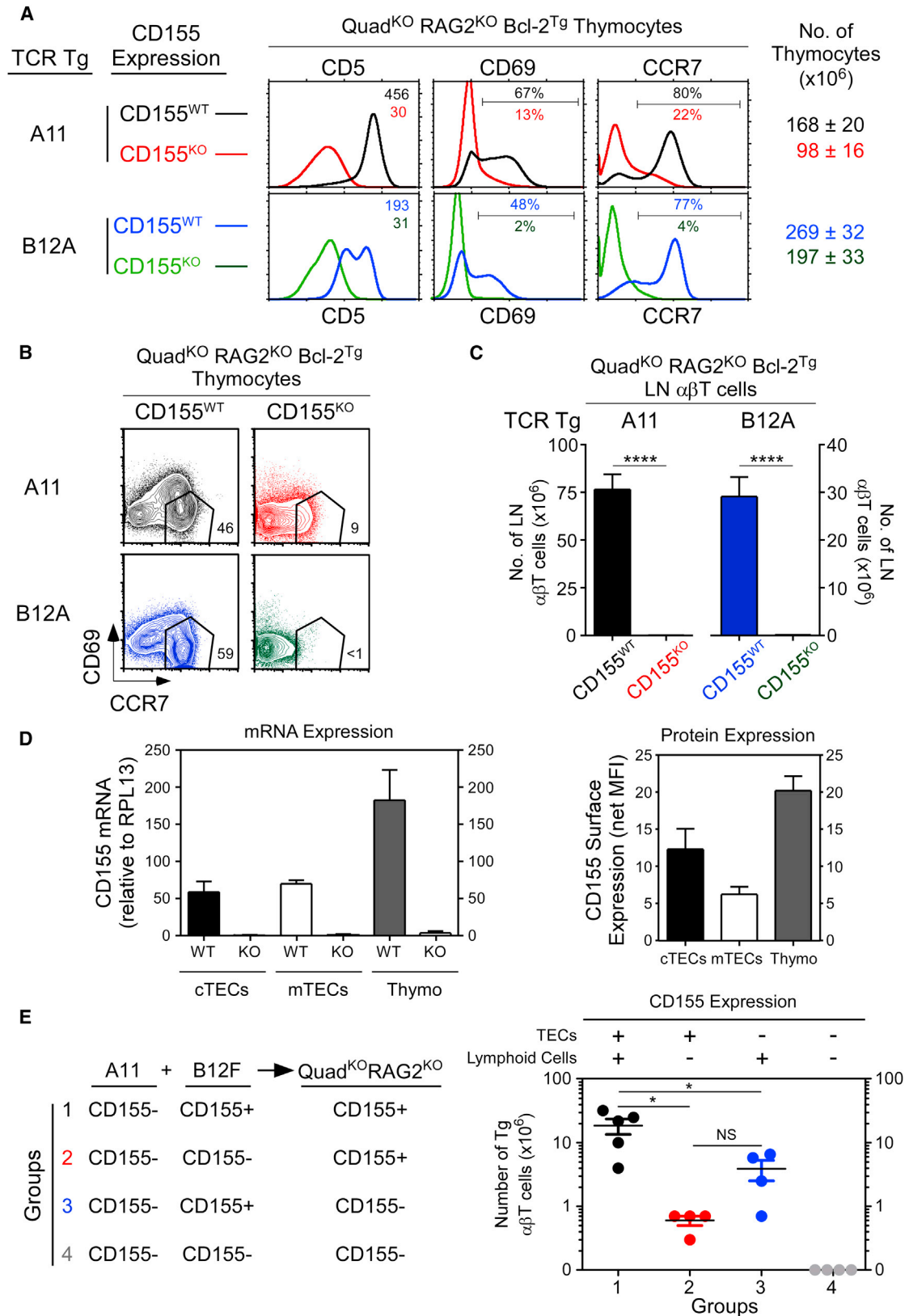
(B) Thymocyte profiles from host transgenic mice expressing transgenic TCR. Numbers in CD5 histograms indicate frequency of positive cells. Thymus cellularity is shown as mean \pm SE (n = 5 mice/group).

(C) Numbers of $\alpha\beta$ LN T cells in TCR transgenic mice (mean \pm SE, n = 5 mice/group).

(D) Expression of CD62L and CD44 on $\alpha\beta$ T cells from TCR transgenic mice. Data represent three experiments.

(E and F) Lymphopenia-induced proliferation triggered by A11 and B12A TCR is dependent on CD155 recognition. A11 and B12A $\alpha\beta$ T cells were CFSE labeled and injected (5×10^6 cells/mouse) into CD155^{+/+} or CD155^{-/-} lymphopenic hosts for 1 week. Numbers in histograms (E) represent the percentage of cells that underwent at least two divisions and summarized in (F). Mean \pm SE of three mice/group.

****p < 0.0001; *p < 0.05. See also Figure S3.



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To understand MHC-independent A11 and B12A TCR signaling of thymocytes in coreceptor-sufficient mice, we considered that transgenic $\alpha\beta$ TCR differed from endogenous $\alpha\beta$ TCR in first being expressed on thymocytes at the DN stage of differentiation when Lck was necessarily coreceptor free. Consequently, we wondered if A11 and B12A transgenic TCR initiated positive selection signaling at the DN stage of thymocyte differentiation. To identify positive selection signaling in TCR transgenic DN thymocytes, we decided on three criteria that would distinguish positive selection from pre-TCR signaling: (1) marked upregulation of both CD5 and CCR7, because pre-TCR signaling minimally upregulates only CD5, (2) ligand-dependence, because pre-TCR signaling is ligand-independent, and (3) TCR specificity, because pre-TCR signaling is induced by transgenic $\alpha\beta$ TCR regardless of recognition specificity.

We observed that A11 and B12A TCR signaling in immature HSA^{hi}DN thymocytes markedly upregulated expression of both CD5 and CCR7 (Figure 5C); such upregulation occurred in *CD155*^{+/+} but not *CD155*^{-/-} mice and was ligand-dependent (Figure 5C). Such upregulation did not occur with B12F TCR and so was TCR-specific (Figure 5D). Thus, A11 and B12A TCR had indeed initiated positive selection signaling in DN thymocytes. Interestingly, thymocytes signaled by A11 and B12A TCR underwent progressive changes in CD4/CD8 and CD69/CCR7 expression expected of thymocytes signaled to undergo positive selection by conventional $\alpha\beta$ TCR (Figure S5A). Moreover, like peripheral CD4 and CD8 T cells selected by conventional $\alpha\beta$ TCR, A11 and B12A CD4 T cells expressed the helper-lineage genes *Zbtb7b* and *Cd40l*, whereas A11 and B12A CD8 T cells expressed the cytotoxic-lineage genes *Runx3d* and *Tbx21* (Figure S5B). Thus, thymocyte differentiation and lineage specification signaled by MHC-independent $\alpha\beta$ TCR resembled that of conventional $\alpha\beta$ TCR.

We think that positive selection signaling by A11 and B12A TCR is initiated in DN thymocytes that had already been pre-TCR signaled to differentiate into DP cells and that positive selection signaling continues until coreceptor proteins are expressed in sufficient amounts to sequester Lck away from the TCR. As confirmation that positive selection signaling by transgenic TCR was initiated in DN thymocytes, DP thymocytes in B12A transgenic mice were not preselection CD5^{lo}CCR7⁻ cells as in unselected B12F transgenic mice, but instead were CD5^{hi}CCR7⁺ cells that had received positive selection signals at their prior (i.e., DN) stage of thymocyte development (Figure 5D). As confirmation that Lck sequestration by coreceptor proteins could inhibit early positive selection signaling by transgenic TCR, prematurely-expressed CD4 transgenes inhibited

generation of mature (CCR7^{hi}CD24^{lo}) thymocytes in B12A transgenic mice (Figure 5E). Interestingly, early Lck sequestration by 444 CD4 transgenic proteins also inhibited $\gamma\delta$ T cell generation in non-TCR transgenic mice (Figure S5C).

We conclude that positive selection signaling by MHC-independent TCR is mediated by coreceptor-free Lck that, in coreceptor-sufficient mice, is present in DN and early (i.e., coreceptor-low) DP thymocytes.

Lck Availability and the Specificity of Positive Selection

Because coreceptor-free Lck transduced positive selection signals by MHC-independent TCR, we wished to determine if coreceptor-free Lck also transduced positive selection signals by MHC-restricted TCR. To do so, we first examined MHC-sufficient mice whose thymocytes were coreceptor-deficient so that endogenous Lck would be coreceptor-free (Figure 6A). Coreceptor-deficiency did not interfere with peripheral T cell generation by A11 and B12A MHC-independent TCR, but it abrogated peripheral T cell generation by MHC-restricted AND and P14 TCR despite *Bcl-2*^{T9} expression (Figure 6A, left). Note that AND and P14 TCR did generate peripheral T cells in coreceptor-sufficient mice (Figure 6A, right). Thus, unlike positive selection signaling by MHC-independent TCR, positive selection signaling by MHC-restricted TCR required coreceptor proteins.

To determine if the requirement by MHC-restricted TCR for coreceptor proteins reflected, at least partly, a requirement for coreceptor-associated Lck, we introduced A11 and AND transgenic TCR into *Lck*^{-/-} mice that expressed coreceptor proteins as well as MHC. Relative to *Lck*^{+/+} mice, peripheral T cell generation by A11 and AND TCR were both significantly reduced in *Lck*^{-/-} mice (Figure 6B). The few peripheral T cells present in *Lck*^{-/-} mice were generated by *Fyn*, as A11 *Lck*^{-/-}*Fyn*^{-/-} mice were essentially devoid of T cells (Figure 6B). We then reconstituted *A11.Lck*^{-/-} and *AND.Lck*^{-/-} mice with coreceptor-free *Lck*^{mut} transgenic proteins (see Figures 1A and 1B) and compared positive selection signaling by *Lck*^{mut} and endogenous Lck proteins as indicated by CD69 upregulation and CD4SP thymocyte generation (Figure 6C). Coreceptor-free *Lck*^{mut} proteins were far more effective than endogenous *Lck*^{+/+} proteins in transducing positive selection signals by MHC-independent A11 TCR, whereas *Lck*^{mut} proteins were far less effective for MHC-restricted AND TCR (Figure 6C). In fact, coreceptor-free *Lck*^{mut} proteins fully restored peripheral T cell generation in *Lck*^{-/-} mice by A11 TCR (>95%), but only minimally restored peripheral T cell generation by AND TCR (<20%) (Figure 6D). Thus coreceptor-free Lck was optimal for positive selection signaling by MHC-independent A11 TCR but

Figure 4. Identification of CD155 as the Thymic Selecting Ligand

(A) In vivo positive selection signaling by A11 and B12A TCR requires CD155. Numbers in CD5 histograms indicate MFI. Numbers in CD69 and CCR7 histograms indicate frequency of positive cells. Thymus cellularity is shown as mean \pm SE (n = 5 mice/group).

(B) CD69 versus CCR7 profiles of A11 and B12A thymocytes. Numbers indicate frequency of mature CD69⁻CCR7^{hi} thymocytes (n = 5 mice/group).

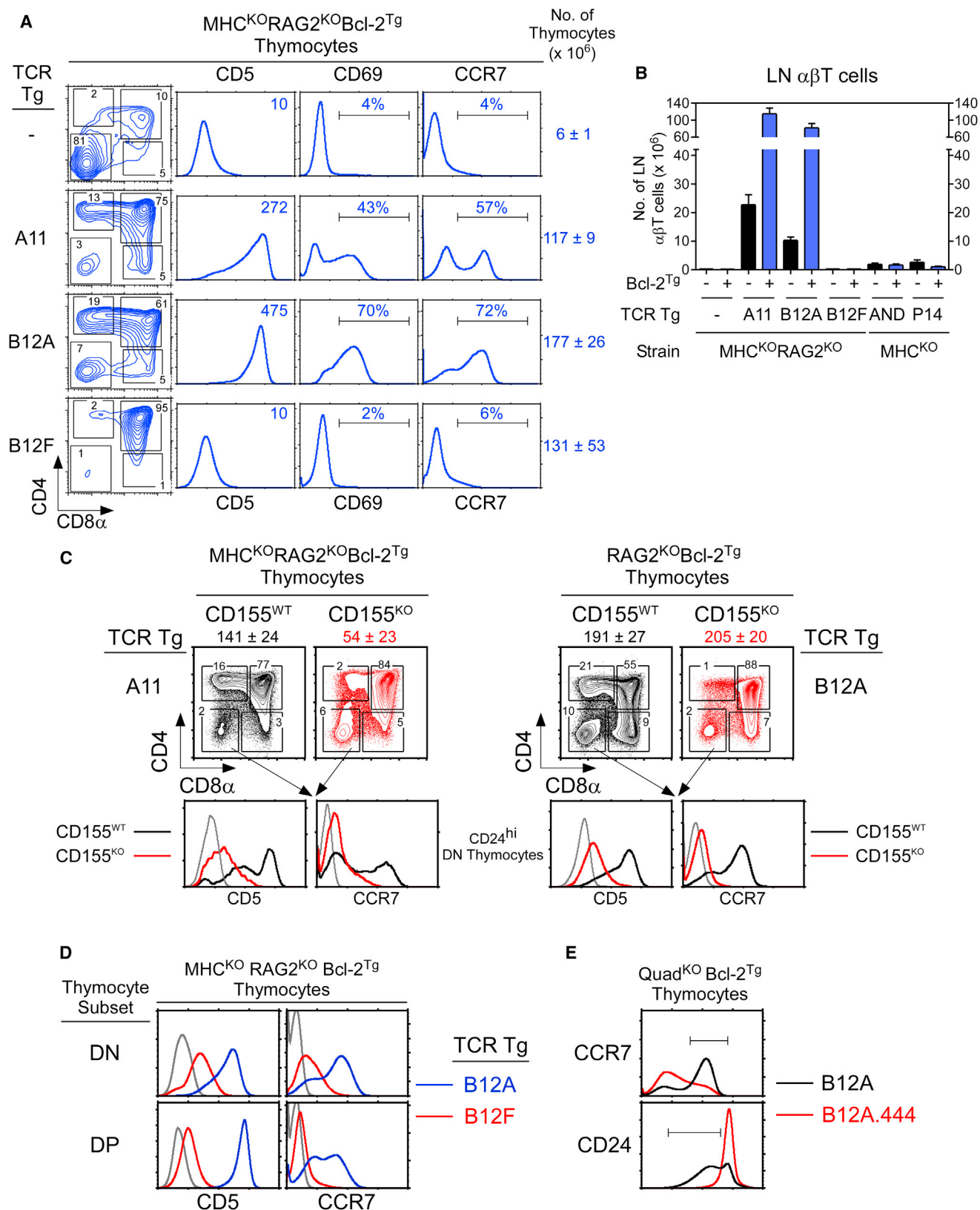
(C) Numbers of $\alpha\beta$ LN T cells (mean \pm SE of 5 mice/group).

(D) CD155 expression by electronically sorted thymic elements by quantitative PCR (left panel) and surface protein expression (right panel). Mean \pm SE (n = 3).

(E) Thymic elements that induce CD155-specific thymic selection. Four groups of mixed donor bone marrow chimeras were constructed with 1:1 mixtures of A11 and B12F donor cells and injected into irradiated host (*Quad*^{KO}*RAG2*^{KO}) mice that expressed or lacked CD155. Numbers of A11 $\alpha\beta$ T cells in LN were determined after 8–10 weeks. Each circle represents one mouse (n = 4).

*p < 0.05; ****p < 0.0001; NS, not significant.

See also Figure S4.



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was only minimally effective for positive selection signaling by MHC-restricted AND TCR.

We conclude that coreceptor-free Lck preferentially promotes positive selection signaling by MHC-independent TCR, whereas coreceptor-associated Lck is required for positive selection signaling by MHC-restricted TCR.

Structural TCR Constraints on MHC-Independent Thymic Selection

The germline concept of MHC restriction proposed that germline-encoded amino acid (aa) residues in CDR1 and CDR2 of TCR α and TCR β have been evolutionarily conserved to engage MHC molecules and to impose MHC specificity on thymic selection (Garcia et al., 2009; Marrack et al., 2008; Scott-Browne et al., 2009). A prediction of the germline concept is that evolutionarily conserved CDR2 aa would not promote TCR selection by MHC-independent ligands. To test this prediction, we constructed hCD2-driven transgenes in which the CDR2 region of the A11 TCR-V β 8 chain contained a point mutation in an evolutionarily conserved aa, either Y48 or E54. A11 TCR variants (referred to as A11^{WT}, A11^{Y48A}, and A11^{E54A}) were introduced into MHC-deficient (MHC^{KO}RAG^{KO}Bcl-2^{Tg}) host mice that were CD155^{+/+} and all three TCR variants were expressed at comparable surface levels (Figure 7A). All three TCR variants reconstituted the cellularity of host thymi and induced the generation of DP thymocytes, indicative of pre-TCR signaling (Figure 7A). However, positive selection signaling that was robust in A11^{WT} thymocytes was absent in A11^{Y48A} and reduced in A11^{E54A} thymocytes (Figure 7A). A11^{Y48A} thymocytes did not upregulate CD5, CD69, or CCR7 (Figure 7A, middle panels), did not differentiate into CD4SP cells (Figure 7A, left panel), and did not differentiate into mature CD69⁻CCR7⁺ cells (Figure 7A, right panel). Biochemical analysis revealed that in vivo TCR signaling was absent in A11^{Y48A} thymocytes as TCR ζ was neither phosphorylated nor associated with ZAP-70 (Figure 7B). In the lymphoid periphery, peripheral T cells were absent in A11^{Y48A} and reduced in A11^{E54A} transgenic mice (Figure 7C). These results reveal that the evolutionarily conserved Y48 aa residue, and to a lesser degree the E54 aa residue, contributed to CD155-specific positive selection signaling by A11 TCR, even though MHC recognition was not involved.

DISCUSSION

The present study demonstrates that thymic selection imposes MHC-restriction on a randomly generated $\alpha\beta$ TCR repertoire that is not intrinsically MHC-specific. It was specifically Lck's availability during thymic selection that determined if the thymus

selected a mature $\alpha\beta$ TCR repertoire that was MHC-restricted or selected one that was MHC-independent. Coreceptor-associated Lck promoted thymic selection of conventionally MHC-restricted $\alpha\beta$ TCR, but coreceptor-free Lck preferentially promoted thymic selection of MHC-independent $\alpha\beta$ TCR. This study also characterized two unique transgenic TCR, A11 and B12A, that utilized coreceptor-free Lck to signal positive selection and to generate mature $\alpha\beta$ T cells in the absence of MHC, unlike any $\alpha\beta$ TCR previously described. The selecting ligand in the thymus for both TCR was the self-ligand CD155 for which these MHC-independent TCR had high affinity (Tikhonova et al., 2012). Finally, the evolutionarily conserved and germline-encoded TCR-V β 8 residues Y48 and E54 in CDR2 β were shown to be necessary for MHC-independent thymic selection of A11, suggesting that these TCR residues might have been conserved during evolution for reasons other than MHC recognition. Thus, the thymus can select either MHC-restricted or MHC-independent TCR, and it is Lck's association with CD4/CD8 coreceptor proteins on preselection thymocytes that limits the thymus to selecting an exclusively MHC-restricted $\alpha\beta$ TCR repertoire.

Coreceptor-associated and coreceptor-free Lck are alternative states of membrane Lck that were mutually exclusive in preselection DP thymocytes. Association with CD4 and CD8 coreceptors physically sequestered membrane Lck away from $\alpha\beta$ TCR, documenting that membrane Lck molecules in preselection DP thymocytes were present in insufficient amounts to bind to coreceptor proteins as well as TCR complexes. Because membrane Lck bound to the cytosolic tails of CD4 and CD8 coreceptors whose external domains interacted with MHC-II and MHC-I determinants, respectively, DP thymocytes could only be signaled by $\alpha\beta$ TCR that engaged intrathymic pMHC ligands together with one or the other coreceptor, whereas coreceptor-free Lck was available to all $\alpha\beta$ TCR. In fact, when preselection thymocytes contained coreceptor-free Lck, either because of mutation of Lck's N-terminal domain cysteines that eliminated coreceptor binding or because of absent coreceptor protein expression, MHC-independent $\alpha\beta$ TCR were able to signal thymic selection and generate an MHC-independent $\alpha\beta$ TCR repertoire. Thus, this study establishes that MHC-independent $\alpha\beta$ TCR do exist in the thymus but are normally prevented from signaling thymic selection by Lck sequestration.

This analysis leads to the insight that TCR gene rearrangements may be specifically timed during normal development to determine the recognition specificities of the TCR expressed by mature $\gamma\delta$ and $\alpha\beta$ T cells. TCR- γ , TCR- δ , and TCR- β gene loci rearrange at the DN stage of thymocyte differentiation but TCR- α gene loci do not rearrange until the DP stage. As a result, $\gamma\delta$ TCR and preTCR complexes (consisting of TCR β and invariant

Figure 5. Positive Selection Signaling by MHC-Independent TCR in Coreceptor-Sufficient Mice Lacking MHC

(A) Thymocyte profiles. Numbers in CD4 versus CD8 α plots indicate cell frequencies. Numbers in CD5 histograms indicate MFI. Numbers in CD69 and CCR7 histograms indicate frequency of positive cells. Thymus cellularity is shown as mean \pm SE (n = 6 mice/group).

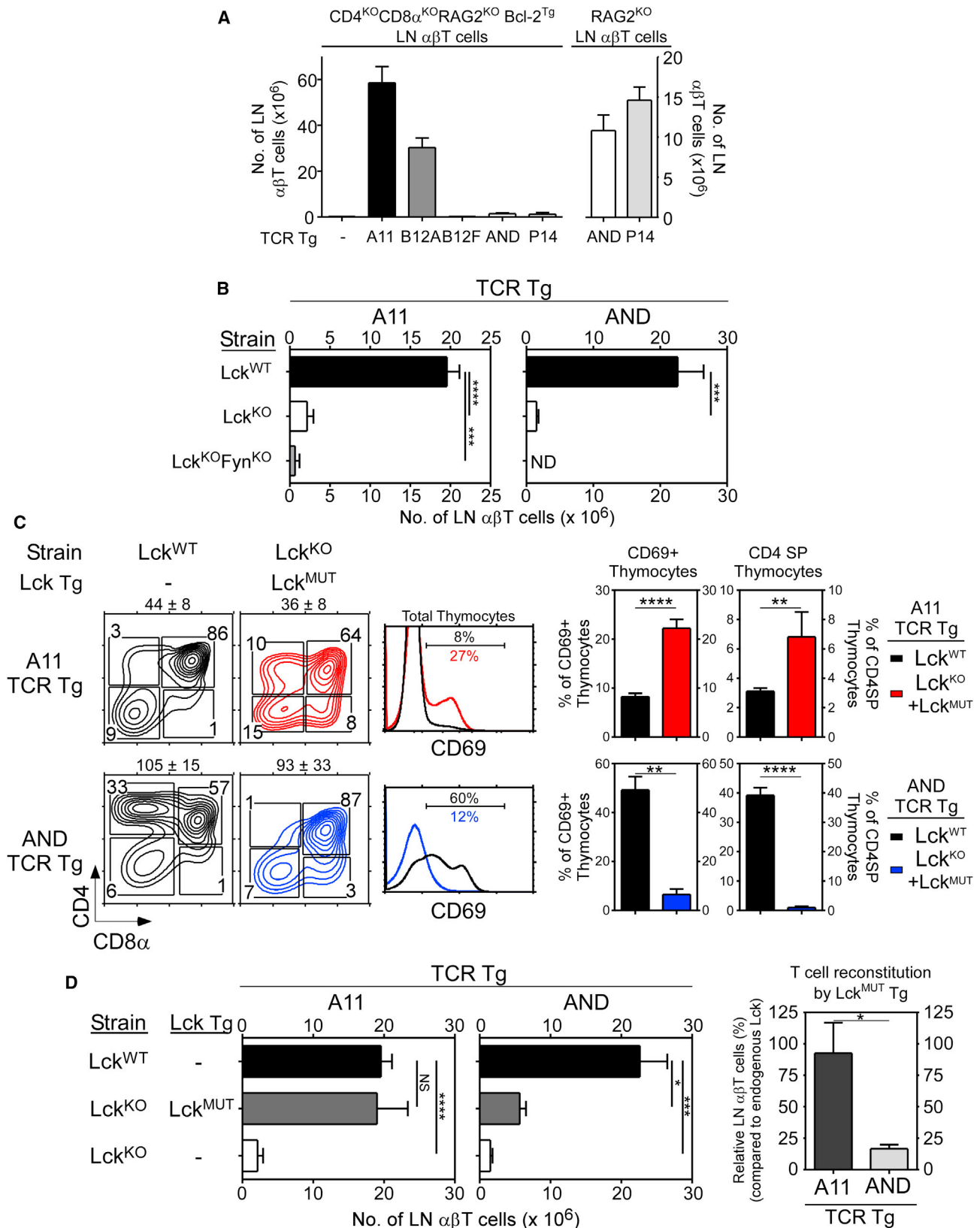
(B) Numbers of $\alpha\beta$ LN T cells (mean \pm SE, n = 6 mice/group).

(C) Positive selection signaling occurs in DN thymocytes and is ligand specific. CD4 versus CD8 α profiles (top) and CD5 and CCR7 expression on CD24^{hi}DN thymocytes (bottom). Data represent four experiments.

(D) Early positive selection signaling is TCR specific. CD5 and CCR7 expression on gated DN and DP thymocytes. Data represent four experiments.

(E) Early positive selection signaling is inhibited by early CD4 expression. CCR7 and CD24 expression on thymocytes from B12A and B12A.444 transgenic mice. Data represent four individual mice/group.

See also Figure S5.



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preT α chains) are formed prior to CD4/CD8 coreceptor expression on DN thymocytes that contain coreceptor-free Lck, whereas $\alpha\beta$ TCR complexes are formed after CD4/CD8 coreceptor expression on DP thymocytes that contain coreceptor-associated Lck. In fact, we suggest that the timing of endogenous $\gamma\delta$ and $\alpha\beta$ TCR expression has specifically evolved to allow different TCR complexes selective access to either coreceptor-free or coreceptor-associated Lck so that ligand recognition by $\gamma\delta$ TCR would be mainly MHC-independent and ligand recognition by $\alpha\beta$ TCR would be MHC-restricted.

This study also characterized two different transgenic $\alpha\beta$ TCR that signaled thymic selection in the absence of MHC, demonstrating the existence of $\alpha\beta$ TCR that do not require MHC for thymic selection. Even though these TCR were MHC-independent, thymic selection signaling by A11 and B12A TCR strictly required engagement of a unique selecting ligand in the thymus that we identified as the self-protein CD155. Identification of CD155 as the unique thymic selecting ligand for A11 and B12A TCR contrasts with the inability to identify unique thymic selecting ligands for all but one conventionally MHC restricted TCR (Lo et al., 2009). We suspect this is because A11 and B12A bind to CD155 with relatively high affinity—and therefore with a high degree of specificity—as a result of their binding to ligand independently of coreceptors. In contrast, we think that MHC-restricted TCR bind to their thymic pMHC selecting ligands with extremely low affinity—and therefore with a low degree of specificity—because these TCR must co-engage those ligands together with coreceptor proteins in order to access coreceptor-associated Lck molecules for transduction of thymic selection signals.

Affinity considerations also explain why coreceptor-free Lck proteins preferentially promoted thymic selection of MHC-independent rather than MHC-restricted TCR. Although coreceptor-free Lck was available to all $\alpha\beta$ TCR, coreceptor-free Lck promoted thymic selection signaling by MHC-independent A11 TCR much more efficiently than MHC-restricted AND TCR. In our view, coreceptor-free Lck is inefficient in promoting TCR signal transduction because coreceptor-free Lck must be passively captured within microclusters of ligand-engaged $\alpha\beta$ TCR, whereas CD4/CD8 coreceptors bring coreceptor-associated Lck directly into microclusters of pMHC-engaged $\alpha\beta$ TCR (Holdorf et al., 2002). As a result, signaling by coreceptor-free Lck requires $\alpha\beta$ TCR with significantly higher ligand affinities than signaling by coreceptor-associated Lck. In fact, the two MHC-independent $\alpha\beta$ TCR characterized in this study bind their MHC-independent selecting ligand CD155 with high affinity that is in the 150–200 nM range. Consequently, we think that coreceptor-free Lck preferentially promotes thymic selection of MHC-independent $\alpha\beta$ TCR because MHC-independent $\alpha\beta$ TCR bind their selecting ligand in the thymus with relatively high affinity

whereas MHC-restricted $\alpha\beta$ TCR bind their selecting ligands in the thymus with very low affinity.

Affinity considerations can also explain why coreceptor-free Lck proteins generated fewer polyclonal $\alpha\beta$ T cells than coreceptor-associated Lck proteins. This experimental observation may appear discordant with the a priori expectation that coreceptor-free Lck would promote signaling by $\alpha\beta$ TCR with a broader range of ligand specificities than coreceptor-associated Lck. However, $\alpha\beta$ TCR that bind thymic selecting ligands with sufficiently high affinity to utilize coreceptor-free Lck would be less frequent than $\alpha\beta$ TCR that bind thymic selecting ligands with the much lower affinity needed to utilize coreceptor-associated Lck. Consequently, we do not think that the fewer polyclonal $\alpha\beta$ T cells generated by coreceptor-free than coreceptor-associated Lck is indicative of the relative numbers of MHC-independent and MHC-restricted $\alpha\beta$ TCR expressed in the preselection repertoire.

Positive selection signaling by MHC-independent $\alpha\beta$ TCR in the thymus resembled that of MHC-restricted $\alpha\beta$ TCR in their requirements for the evolutionarily conserved Y48 and E54 residues in the CDR2 region of V β 8 TCR (Feng et al., 2007; Garcia et al., 2009; Marrack et al., 2008). Positive selection signaling by a transgenic MHC-restricted V β 8 TCR was previously shown to require Y48 and E54, arguably because of their roles as MHC contact residues that promoted TCR engagement of pMHC selecting ligands in the thymus and fulfilling a prediction of the germline model of MHC restriction (Scott-Browne et al., 2009). However, the current study showed that the transgenic MHC-independent A11 V β 8 TCR also required Y48 and E54 for optimal positive selection signaling, even though MHC was not involved. Reciprocally, it was recently shown that an MHC-restricted TCR repertoire could still be generated without evolutionarily conserved germline-encoded CDR1 and CDR2 sequences (Holland et al., 2012). We suggest that germline-encoded CDR1 and CDR2 amino acid residues may have been conserved in evolution for reasons unrelated to their role as MHC contact residues but possibly related to their role in maintaining the integrity of the TCR combining site.

As thymic differentiation of MHC-independent $\alpha\beta$ TCR has not previously been characterized, it is worth noting that, because transgenic MHC-independent $\alpha\beta$ TCR are expressed in DN thymocytes and signal in the absence of coreceptors, transgenic MHC-independent $\alpha\beta$ TCR initiated positive selection signaling in thymocytes at the DN stage of differentiation. However, we think that DN thymocytes that were signaled to undergo positive selection had already been signaled by their pre-TCR to differentiate into DP thymocytes, in which case positive selection signaling continued until coreceptor protein expression was high enough to sequester Lck and extinguish further

Figure 6. Effect on Thymic Selection of Coreceptor-Associated and Coreceptor-free Lck

- (A) Numbers of $\alpha\beta$ LN T cells in coreceptor-deficient (left side) or coreceptor-sufficient (right side) mice (mean \pm SE, n = 3 mice/group).
 (B) Numbers of $\alpha\beta$ LN T cells in kinase-deficient mice (mean \pm SE, n = 3 mice/group). ND, not done.
 (C) Profiles of thymocytes with endogenous Lck (black lines) or Lck^{mut} (colored lines) proteins. Numbers in CD4 versus CD8 α plots indicate cell frequencies. Numbers in CD69 histograms indicate frequency of positive cells and is summarized in right panels (mean \pm SE, n = 3 mice/group).
 (D) Numbers of $\alpha\beta$ LN T cells in TCR transgenic mice with endogenous Lck (black bar) or Lck^{mut} (gray bar) proteins. Reconstitution by Lck^{mut} proteins relative to endogenous Lck proteins (set at 100%) is displayed in the right panel (mean \pm SE, n = 3 mice/group).

*p < 0.05; **p < 0.01; ***p < 0.001; ****p < 0.0001; NS, not significant.

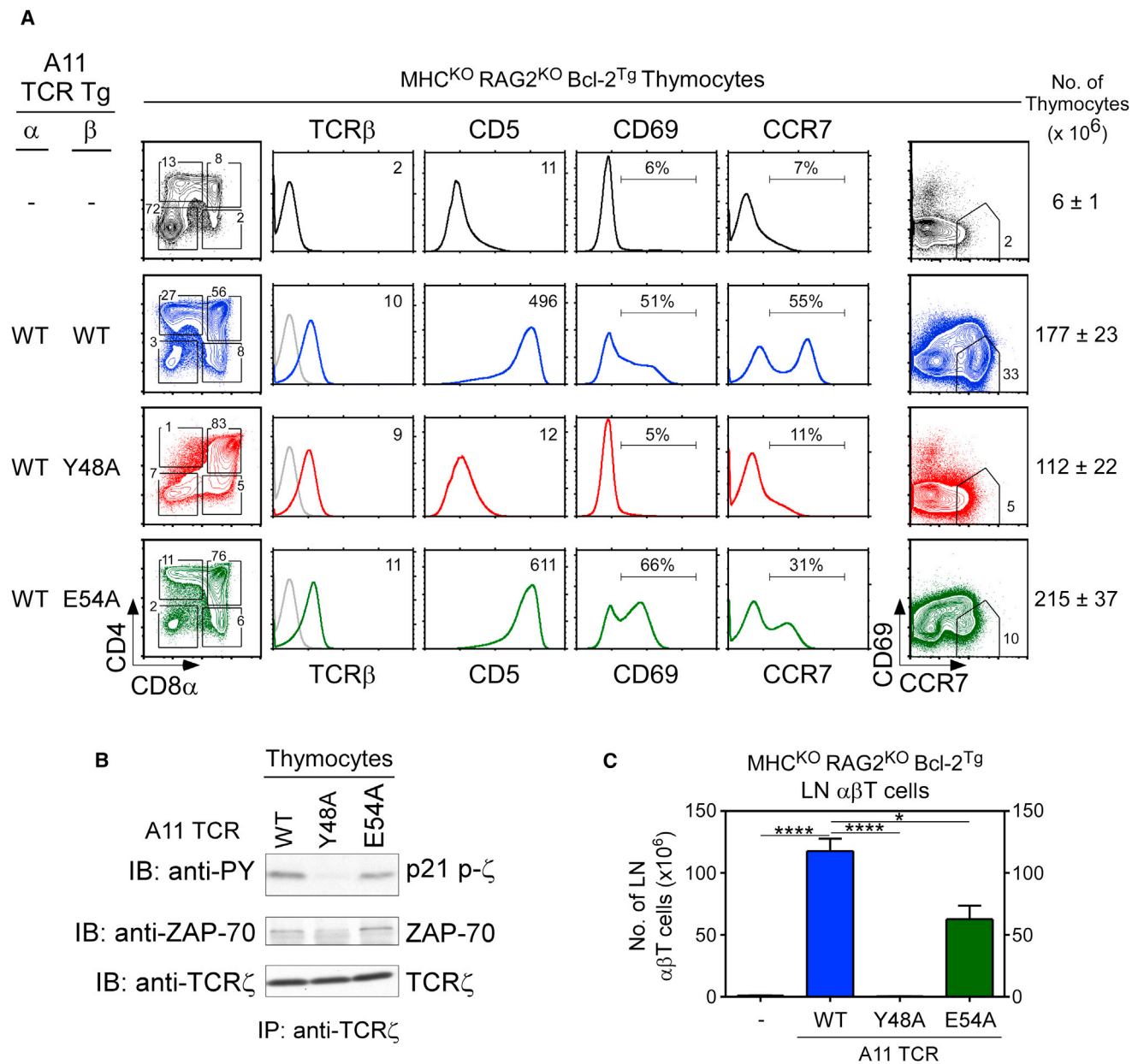


Figure 7. Role of Evolutionarily Conserved CDR2 β Residues in Thymic Selection by an MHC-Independent TCR

(A) Thymocyte profiles from MHC-deficient mice expressing A11 TCR whose TCR β chain was wild-type or contained a point mutation in CDR2, either Y48A or E54A. Numbers in TCR β and CD5 histograms indicate MFI. Numbers in CD69 and CCR7 histograms indicate frequency of positive cells. Numbers in CD69 versus CCR7 plots indicate frequency of mature cells. Thymus cellularity is shown as mean \pm SE (n = 4 mice/group).

(B) Assessment of in vivo signaling of thymocytes. Unstimulated thymocyte lysates were precipitated with anti-TCR ζ and blotted for phospho-zeta, ZAP70, or TCR ζ . Data represent two experiments.

(C) LN $\alpha\beta$ T cell numbers (mean \pm SE, n = 4 mice/group).

*p < 0.05; ****p < 0.0001.

MHC-independent TCR signaling. Nevertheless, thymocytes signaled by transgenic MHC-independent $\alpha\beta$ TCR underwent similar changes in CD4/CD8 and CD69/CCR7 expression in the thymus as thymocytes signaled by conventional MHC-restricted $\alpha\beta$ TCR and gave rise to similar CD4/helper- and CD8/cytotoxic-lineage T cells in the periphery. Thus, thymocyte

development and lineage-specification appear to progress identically regardless of the phenotypic stage in thymocyte differentiation that $\alpha\beta$ TCR-mediated positive selection signaling occurs.

In conclusion, this study has documented the existence of MHC-independent $\alpha\beta$ TCR in the thymus and identified the basis for thymic selection of an MHC-restricted $\alpha\beta$ TCR repertoire.

This study has also demonstrated that generation of an unconventional MHC-independent $\alpha\beta$ TCR repertoire expressed by peripheral T cells requires the presence of coreceptor-free Lck during thymic selection. As a result, TCR gene rearrangements appear to be carefully timed during development in the thymus so that $\gamma\delta$ TCR specifically encounter coreceptor-free Lck and $\alpha\beta$ TCR specifically encounter coreceptor-associated Lck so that T cells expressing these different TCR complexes will primarily recognize MHC-independent and MHC-restricted ligands, respectively.

EXPERIMENTAL PROCEDURES

Animals

C57BL/6 (B6), *B2m*^{-/-}, *Rag2*^{-/-}, *Lck*^{-/-}, *Fyn*^{-/-}, *Tcr α* ^{-/-}, and *Cd4*^{-/-} mice were purchased from Jackson Laboratories. B10.A mice were obtained from NCI. *CD8 α* ^{-/-} and *H2-Ab1*^{-/-} mice were bred in our own animal colony, as were AND TCR transgenic mice (Kaye et al., 1989), P14 TCR transgenic mice (Pircher et al., 1989), and hBcl-2 transgenic mice (Linette et al., 1996). CD4 transgenic mice expressing full-length 444 or tailless 44T transgenic CD4 proteins were generated as previously described (Van Laethem et al., 2007), as were *CD155*^{-/-} mice (Maier et al., 2007). Animal care was in accordance with National Institutes of Health (NIH) guidelines.

New transgenic mouse strains constructed for this study were generated by cloning gene-specific cDNAs into the human CD2 transgenic vector to achieve T cell specific expression. For wild-type and mutant Lck transgenes, full-length Lck cDNA was cloned from purified B6 $\alpha\beta$ T cells in which residues C20 and C23 were both mutated to alanine using the QuikChange XL site-directed mutagenesis kit (Stratagene). cDNAs encoding Lck^{wt} and Lck^{mut} proteins were each cloned into the transgenic vector and injected into B6 oocytes. For TCR transgenes, full-length cDNAs encoding the rearranged TCR α and TCR β chain of each TCR (A11, B12A, B12F) were separately cloned into TCR α and TCR β vectors that were then injected together into B6 oocytes. For A11 TCR transgenes with CDR2 mutations, residues Y48 and E54 in the cDNA encoding the A11 TCR β chain were individually mutated to alanine, cloned into transgenic vectors, and injected together with the A11 TCR α transgenic vector into B6 oocytes. At least two independent founder lines for each transgene were generated and analyzed.

Antibodies and Reagents

For a detailed list see the [Extended Experimental Procedures](#).

Flow Cytometry and Cell Sorting

See the [Extended Experimental Procedures](#) for details.

Calcium Mobilization and Protein Immunoblotting

Assays were performed as described (Van Laethem et al., 2007). For details see the [Extended Experimental Procedures](#).

T Cell Proliferation and Activation

To test T cell reactivity, purified $\alpha\beta$ LNT responder cells (1×10^5 /well) were labeled with CFSE and cocultured for 4 days with LPS-activated and irradiated splenic B stimulator cells (2×10^5 /well). For in vivo proliferation, $5\text{--}10 \times 10^6$ CFSE-labeled T cells were injected into unmanipulated lymphopenic host mice. IL-2 production by T-hybridomas were measured by standard ELISA.

Construction of Mixed Bone Marrow Chimeras

Bone marrow stem cells (10^7 T cell-depleted) from each donor were injected into sublethally irradiated (6 Gy) *Rag2*^{-/-} host mice and analyzed 8–10 weeks later.

Thymic Epithelial Cell Isolation

Thymi were teased apart to release all thymocytes and then treated with collagenase/DNase/dispase for four sequential digestions at 37°C for 10 min each (Gray et al., 2008). Cells were then stained with mAb specific for EpCAM, CD45, MHCII, Ly51, and UEA-1, to distinguish cortical and medullary thymic epithelial cells.

Statistical Analyses

Student's t test with two-tailed distributions was used for statistical analyses. P values of 0.05 or less were considered significant.

SUPPLEMENTAL INFORMATION

Supplemental Information includes Extended Experimental Procedures and five figures and can be found with this article online at <http://dx.doi.org/10.1016/j.cell.2013.08.009>.

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