

BRIEF DEFINITIVE REPORT

miR-155 promotes T reg cell development by safeguarding medullary thymic epithelial cell maturation

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During thymocyte development, medullary thymic epithelial cells (mTECs) provide appropriate instructive cues in the thymic microenvironment for not only negative selection but also the generation of regulatory T (T reg) cells. Here, we identify that miR-155, a microRNA whose expression in T reg cells has previously been shown to be crucial for their development and homeostasis, also contributes to thymic T reg (tT reg) cell differentiation by promoting mTEC maturation. Mechanistically, we show that RANKL stimulation induces expression of miR-155 to safeguard the thymic medulla through targeting multiple known and previously uncharacterized molecules within the TGF β signaling pathway, which is recognized for its role in restricting the maturation and expansion of mTECs. Our work uncovers a miR-155-TGF β axis in the thymic medulla to determine mTEC maturity and, consequently, the quantity of tT reg cells and suggests that miR-155 ensures proper tT reg cell development in both cell-intrinsic and -extrinsic manners.

Introduction

Regulatory T (T reg) cells constitute a specialized tolerogenic subset of cells recognized for maintaining immune homeostasis and preventing inappropriate reactivity to self-antigens and innocuous foreign antigens (Josefowicz et al., 2012a). While T reg cells can be generated in the periphery and play a nonredundant role in restraining allergic-type inflammation at mucosal interfaces, those generated in the thymus (tT reg cells) are absolutely critical for controlling systemic and tissue-specific autoimmunity (Josefowicz et al., 2012b). To this end, the thymic medulla represents a specific site for establishing self-tolerance via the generation of tT reg cells in addition to its known role in mediating negative selection (Hinterberger et al., 2010). In the thymic medulla, medullary thymic epithelial cells (mTECs) express high levels of MHCII molecules, tissue-restricted antigens, and costimulatory ligands CD80/CD86 in order to foster an instructive cross-talk between these specialized thymic stromal cells and developing thymocytes (Lucas et al., 2016). Disruption of the aforementioned interactions results in failed de novo generation of tT reg cells (Aschenbrenner et al., 2007; Malchow et al., 2016; Salomon et al., 2000; Tai et al., 2005). The fact that loss of mTECs leads to an explicit defect in the tT reg cell compartment while leaving conventional CD4

single positive thymocytes unaffected further substantiates an indispensable role for the thymic medulla in tT reg cell differentiation (Cowan et al., 2013).

To support the generation of tT reg cells, mTECs themselves need to differentiate properly. Activation of the RelB-dependent noncanonical NF- κ B pathway driven by tumor necrosis factor superfamily cytokines such as receptor activator of NF- κ B ligand (RANKL), CD40 ligand, and lymphotoxin β (LT β) have been shown to be essential for mTEC progenitors to undergo a stepwise differentiation process to generate immature MHCII^{lo}CD80^{lo} mTECs before mature MHCII^{hi}CD80^{hi} mTECs (Akiyama et al., 2008; Hikosaka et al., 2008; Irla et al., 2008). Among them, RANKL stimulation is particularly important for the induction of autoimmune regulator (AIRE), a transcription factor that plays a major role in driving the expression of tissue-restricted antigens in mature mTECs (Anderson et al., 2002; Rossi et al., 2007; Zuklys et al., 2000). On the other hand, TGF β has been shown to play a negative role in restraining mTEC maturation by interfering with the noncanonical NF- κ B pathway (Hauri-Hohl et al., 2014). However, because TGF β can be detected in the thymus shortly after birth, and predominantly in the thymic medulla (Konkel et al., 2014), exactly how mTECs

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shield themselves from persistent TGF β exposure remains unclear (Hauri-Hohl et al., 2014).

MicroRNAs (miRNAs) comprise a class of small noncoding RNAs that regulate gene expression at the posttranscriptional level and whose roles in controlling the development and function of T cells, including T reg cells, are well established (Chong et al., 2008; Cobb et al., 2006; Cobb et al., 2005; Liston et al., 2008; Zhou et al., 2008). It is also now appreciated that miRNAs can regulate thymic T cell differentiation by maintaining a proper thymic microenvironment, where deletion of the miRNA network within TECs severely compromises thymic infrastructure and largely impacts mTECs (Khan et al., 2014; Papadopoulos et al., 2012; Zuklys et al., 2012). Nevertheless, the current understanding of individual miRNAs crucial for controlling different aspects of TEC biology and, more importantly, their impact on thymic T cell development, remains limited (Khan et al., 2015). Here, we show that miR-155, a prominent miRNA known for its diverse functions in various immune cell populations (Vigorito et al., 2013), plays an equally important role in mTECs. Previously, we and others have shown that elevated expression of miR-155 driven by Foxp3 ensures proper T reg cell homeostasis by maintaining their competitive fitness (Lu et al., 2009). Our current work further demonstrates that miR-155 promotes T reg cell development in the thymus by safeguarding mTEC maturation. Mechanistically, RANK signaling induces miR-155 expression in the thymic medulla to alleviate the negative effects that ensue from the continuous presence of intrathymic TGF β via targeting multiple known and previously uncharacterized molecules within this cytokine-signaling pathway. As such, the miR-155-TGF β axis maintains the mature mTEC population and, thus, establishes a thymic microenvironment favorable for tT reg cell development.

Results and discussion

miR-155 promotes tT reg cell development in both T cell-intrinsic and -extrinsic manners

miRNAs play a pivotal role in controlling multiple aspects of T reg cell biology. Among the many miRNAs predominantly expressed in T reg cells, miR-155 is crucial for promoting optimal T reg cell development and homeostasis partly through ensuring responsiveness to IL-2, a cytokine required for thymic and peripheral T reg cell maintenance (Kohlhaas et al., 2009; Lu et al., 2009). Mice harboring a germline deficiency of miR-155 present with diminished T reg cell numbers and frequencies, plus additional studies using mixed bone marrow chimeras have identified a cell autonomous role of miR-155 in controlling T reg cell biology (Kohlhaas et al., 2009; Lu et al., 2009). Consistent with this notion, upon T cell-specific miR-155 ablation (Fig. S1 A), reduced frequencies as well as total numbers of T reg cells in the thymus are also detected (Fig. 1, A–C). Interestingly though, the degree of reduction in T cell-specific miR-155 conditional KO (T-cKO) mice does not fully recapitulate that observed in mice completely devoid of miR-155 (~30% in T-cKO versus ~60% in miR-155 null; Fig. 1 D; Kohlhaas et al., 2009; Lu et al., 2009; Sánchez-Díaz et al., 2017). These results suggest that loss of miR-155 expression in other non-T cell populations may also

contribute to the impaired tT reg cell phenotype observed in mice containing miR-155 germline deficiency.

miR-155 is preferentially expressed in mature mTECs and is induced by RANKL stimulation

mTECs function as a key stromal cell population crucial for the generation of T reg cells in the thymus, and so it is possible that miR-155 promotes tT reg cell development by regulating mTEC biology. In support of this notion, a previous miRNA profiling study has demonstrated that miR-155 is expressed at elevated levels in mTECs compared with cortical thymic epithelial cells (cTECs), another population of thymic stromal cells critical for early thymocyte development (Khan et al., 2015). In line with this work, our analyses of different thymic resident cell subsets also revealed higher expression levels of miR-155 in mTECs relative to cTECs, albeit lower than that in CD45⁺ immune cells (Fig. 2, A and B). Moreover, in scrutiny of two major subsets within the mTEC population, CD80^{lo}MHCII^{lo} and CD80^{hi}MHCII^{hi} (Fig. 2 C), with the latter comprising the more mature subset essential for tT reg cell generation, we found that miR-155 expression is restricted to the CD80^{hi}MHCII^{hi} mTEC compartment. Together, these data point to a potential role of miR-155-mediated regulation of mTEC maturation.

Among different tumor necrosis factor superfamily members that have been characterized in mTEC development, RANKL-RANK is the foremost determinant of mTEC development (Akiyama et al., 2008). To examine whether elevated miR-155 expression in mature mTECs is induced by RANK signaling, primary CD80^{lo}MHCII^{lo} immature mTECs were isolated and stimulated with RANKL. Along with prior work (Rossi et al., 2007), we noted an induction of AIRE mRNA in these cells as early as 6 h following RANKL stimulation (Fig. 2 E). Moreover, along with up-regulation of AIRE, an increase in the primary transcript of miR-155 (*pri-miR-155*) was also readily detectable (Fig. 2 E). Contrarily, a reduction in *pri-miR-155* was observed concomitantly with diminished expression of AIRE in mTECs isolated from mice treated with RANKL-blocking antibodies (Fig. 2 F). Collectively, these results locate RANK signaling in driving miR-155 expression in mTECs and imply functional relevance for miR-155 during mTEC development.

Expression of miR-155 in TECs is required to maintain optimum mTEC maturation and tT reg cell development

To investigate the potential role of miR-155 in mTEC maturation and its subsequent impact on tT reg cell development, we generated mice with TEC-specific ablation of miR-155 by crossing miR-155 floxed mice (*miR-155^{fl}*) to FOXN1-Cre mice (Fig. S1, B–D). Considering that deletion of miR-155 is not restricted to mTECs but rather encompasses the entire thymic epithelium, we first sought to determine whether the cellularity and phenotype of the thymic epithelia would be impacted by the loss of miR-155. As depicted in Fig. 3 A (and data not shown), we did not detect any alterations in total thymic cellularity, including proportions of cTECs and mTECs, upon deletion of miR-155 in TECs relative to control WT or T-cKO mice. In contrast, we found that the frequency of CD80^{hi}MHCII^{hi} mature mTECs was significantly reduced in TEC-specific miR-155 conditional knockout mice

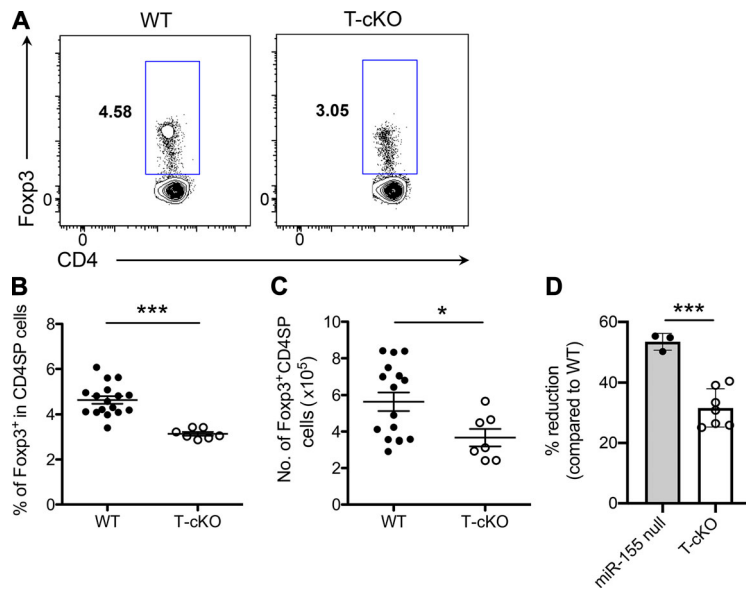


Figure 1. miR-155 promotes tT reg cell development in both T cell-intrinsic and -extrinsic manners. (A–C) FACS analysis (A), frequencies (B), and absolute numbers (C) of total Fop3+ T cells in the thymus of 5–6-wk-old WT and T-cKO mice. (D) Percentages of the reduction of tT reg cell frequencies (on the basis of corresponding WT controls) in miR-155 null and T-cKO mice. Each symbol represents an individual mouse, and the bar represents the mean. Data are pooled from at least three independent experiments. Results of Student's *t* test: *, *P* < 0.05; ***, *P* < 0.001.

(TEC-cKO) mice in comparison to both WT and T-cKO controls, supporting our proposed function of miR-155 during mTEC maturation (Fig. 3 B). Accompanied by a decrease in mature mTECs that also express AIRE, we detected an increase in the frequency of a subset of AIRE⁺CCL21⁺ mTECs in TEC-cKO mice (Fig. S2 A). Unlike AIRE-expressing mTECs, the development of this specific mTEC subset relies on LT β receptor signaling and has been implicated in recruiting (and/or retaining) positively

selected CCR7⁺ thymocytes to the medulla (Lkhagvasuren et al., 2013; Zhang and Bhandoola, 2014). On the other hand, no significant change was observed in DCLK1-expressing thymic tuft cells (Fig. S2 B), another mTEC subset that is transcriptionally distinct from the AIRE⁺ mature mTECs (Bornstein et al., 2018). Interestingly, despite the reduction of mature mTECs in TEC-cKO mice, on a per-cell basis, levels of AIRE in mature mTECs were comparable between TEC-cKO and control mice (Fig. S2, C

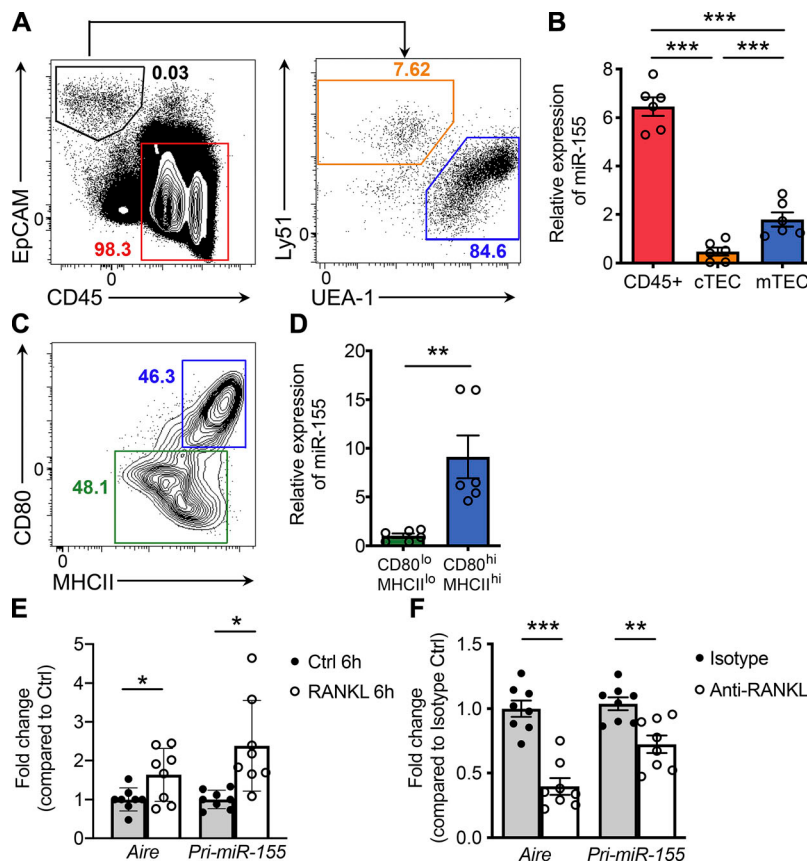


Figure 2. RANKL stimulation results in elevated miR-155 expression in mature mTECs. (A) Representative FACS profiles with gating strategy for different thymic subsets are shown. Hematopoietic cells were defined as CD45⁺EpCAM⁺. TECs were defined as CD45⁺EpCAM⁺ and further divided into cTEC (Ly51⁺UEA-1⁻) and mTEC (Ly51⁺UEA-1⁺) populations. (B) qPCR analyses of miR-155 expression in hematopoietic cells, cTECs, and mTECs. (C) Immature and mature mTECs within the mTEC population were further discriminated based on the levels of expression of CD80 and MHCII molecules. (D) qPCR analyses of miR-155 expression in CD80^{lo}MHCII^{lo} immature mTECs and CD80^{hi}MHCII^{hi} mature mTECs. (E and F) qPCR analyses for Aire and pri-miR-155 expressions in sorted primary immature mTECs stimulated with 500 ng/ml RANKL for 6 h in vitro (E) or in total mTECs isolated from mice administrated with anti-RANKL or isotype control antibodies (F). The n-fold changes on the basis of corresponding untreated control groups were shown. Each symbol represents an individual mouse, and the bar represents the mean. Data are pooled from at least three independent experiments. Results of Student's *t* test: *, *P* < 0.05; **, *P* < 0.01; ***, *P* < 0.001.

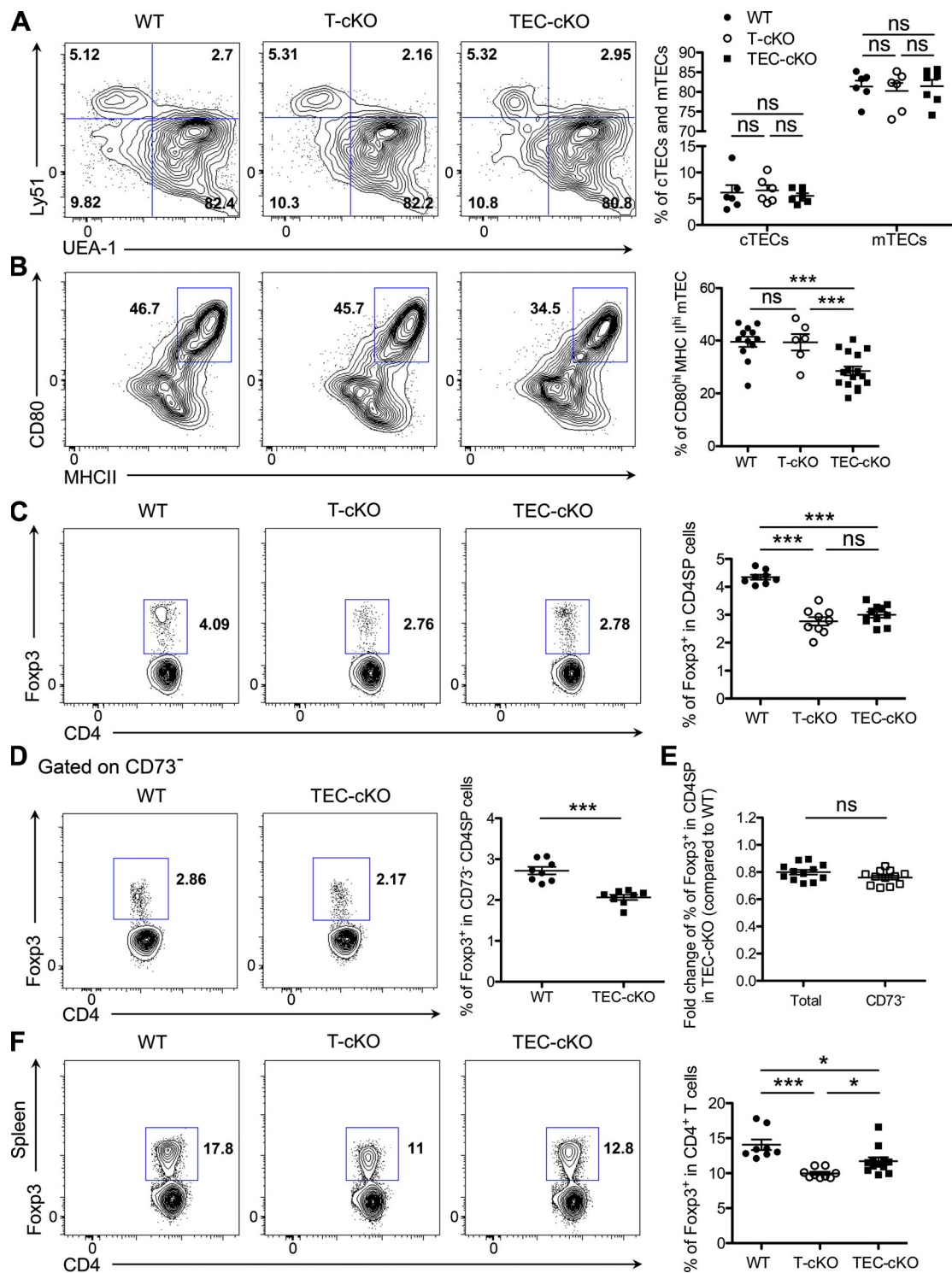


Figure 3. miR-155 deficiency in TECs leads to a diminished mature mTEC population and impaired tT reg cell development. (A) FACS analysis and frequencies of Ly51⁺ UEA-1⁻ cTEC and Ly51⁺ UEA-1⁺ mTEC populations from 5–6-wk-old WT, T-cKO, and TEC-cKO mice. (B) FACS analysis and frequencies of CD80^{hi} MHCII^{hi} mTECs from 5–6-wk-old WT, T-cKO, and TEC-cKO mice. (C and D) FACS analysis and frequencies of total Foxp3⁺ T reg cells (C) or CD73⁻ Foxp3⁺ T reg cells (D) in the thymus of 5–6-wk-old WT, T-cKO, and TEC-cKO mice. (E) Fold change of percentage of total or CD73⁻ T reg cells in TEC-cKO mice compared with WT littermates. (F) FACS analysis and frequencies of Foxp3⁺ T reg cells in the spleen of 5–6-wk-old WT and TEC-cKO mice. Each symbol represents an individual mouse, and the bar represents the mean. Data are pooled from at least three independent experiments. Results of Student's *t* test: *, *P* < 0.05; ***, *P* < 0.001; ns, not significant.

and D), implying that miR-155 does not directly regulate AIRE expression despite its role in mTEC maturation.

Due to the aforementioned role of mature mTECs in tT reg cell development and the impairment in mature mTECs from mice bearing TEC-specific ablation of miR-155, we next asked whether the generation of tT reg cells is similarly affected in TEC-cKO mice. Analogous to what has been discerned in T-cKO mice, a comparable reduction in tT reg cell frequency was detected in TEC-cKO mice (Fig. 3 C), highlighting that miR-155 as expressed in T cells and TECs contributes equally to T reg cell development in the thymus. Further supporting this notion, a similar reduction in the frequency of nascent tT reg cells in TEC-cKO mice was also seen when mature recirculating CD73⁺ T reg cells were excluded from the total tT reg cell population (Fig. 3, D and E). It should be noted that while miR-155 deletion in mTECs results in a reduction in tT reg cell frequency, the proliferative capacity as well as expression levels of Foxp3 and other T reg cell-associated molecules were unaffected in these T reg cells, relative to WT controls (Fig. S3). In the spleen, even though the frequency of T reg cells in TEC-cKO mice remained lower than WT controls, significantly more T reg cells in the spleen of TEC-cKO mice were seen when compared with T-cKO mice (Fig. 3 F). While the difference between the results obtained from the thymus and the spleen is intriguing, it is not surprising, seeing as miR-155 is a known crucial regulator conferring competitive fitness to T reg cells both in the thymus and in the periphery in a cell autonomous manner (Lu et al., 2009). Taken together, these data clearly demonstrate an indispensable role of TEC-derived miR-155 in controlling mTEC maturation and, subsequently, the generation of T reg cells in the thymus.

miR-155 limits TGF β signaling via targeting multiple components of the TGF β signaling pathway

The TGF β signaling cascade has previously been reported to play a regulatory role in limiting the establishment and function of the thymic medulla, specifically by influencing the differentiation of mTECs (Konkel et al., 2014). In addition, miR-155 has also been implicated in regulating TGF β signaling by directly targeting *Tgfb2* and *Smad2* in human lung fibroblasts and in THP-1 monocyte cell lines, respectively (Chu et al., 2017; Louafi et al., 2010). Therefore, it is possible that miR-155 also promotes mTEC maturation by restraining TGF β signaling. Indeed, we observed increased levels of both *Tgfb2* and *Smad2* in mature mTECs isolated from TEC-cKO mice, indicating that these two genes are also subject to regulation by miR-155 in mature mTECs (Fig. 4, A and B). Consistently, we detected higher levels of *P21*, an established TGF β -induced gene (Datto et al., 1995), in mature mTECs devoid of miR-155. The levels of *Myc*, a gene known to be repressed by TGF β activity (Frederick et al., 2004), were, alternatively, reduced (Fig. 4, C and D). Overall, these results are suggestive of enhanced TGF β signaling in the absence of miR-155-mediated gene regulation.

Since miRNAs usually exert their regulatory effects by targeting multiple genes in a shared pathway or protein complex to ensure biological impact (Ebert and Sharp, 2012), we sought to define additional targets in the TGF β signaling pathway that may be controlled by miR-155 in mature mTECs. After analyzing

previous results obtained from high-throughput sequencing of RNAs isolated by cross-linking immunoprecipitation (HITS-CLIP), a biochemical approach that affords the identification of functional miRNA-mRNA interaction in a given tissue/cell sample (Loeb et al., 2012), we identified SMAD3, a molecule that acts cooperatively with SMAD2 to form major TGF β signaling transducers, as another potential miR-155 target (Fig. 4, E and F). Supporting this notion, our luciferase reporter studies confirmed that miR-155 directly represses SMAD3 (Fig. 4 G), and mature mTECs isolated from TEC-cKO mice express significantly higher amounts of *Smad3* transcript (Fig. 4 H). Moreover, by taking similar approaches, we also identified and confirmed RNF111 as a direct target of miR-155 (Fig. 4, E-H). RNF111 (or “Arkadia”) is an E3 ubiquitin ligase recognized for its role in enhancing TGF β responses by promoting the degradation of c-SKI, a known negative regulator of TGF β signaling that blocks TGF β -driven transcriptional activation and repression by forming an inhibitory complex with SMAD proteins (Sharma et al., 2011; Suzuki et al., 2004). Thus, our data show that through targeting multiple requisite components within the TGF β signaling pathway, ranging from the receptor to major signal transducers and to the E3 ligase that degrades the TGF β signaling inhibitor, miR-155 acts as a key molecule involved in attenuating TGF β signaling in mTECs amid their maturation process.

Diminished mTECs and tT reg cells in TEC-cKO mice are largely rescued by partial TGF β receptor deletion

While our results demonstrate that TEC-specific miR-155 ablation leads to enhanced TGF β signaling in mTECs, it remains obscure as to whether loss of miR-155-dependent regulation of TGF β signaling is responsible for the mTEC and tT reg cell phenotypes observed in TEC-cKO mice. Previously, it has been shown that short-term systemic pharmacological inhibition of TGF β signaling with the TGF β receptor I (TGF β RI) kinase inhibitor selectively increases the number of mature mTECs (Hauri-Hohl et al., 2014). Interestingly, however, such treatment did not seem to rescue the defective tT reg cell phenotype observed in mice with TEC-specific miR-155 ablation; only a modest increase of tT reg cell frequencies was detected in TEC-cKO mice following treatment of the TGF β RI kinase inhibitor compared with the untreated group (Fig. 5, A and B). It should be noted that TGF β signaling in thymocytes has previously been reported to be needed for both induction of Foxp3 and the differentiation of tT reg cells (Konkel et al., 2014). It is therefore possible that any positive effect resulting from blockade of TGF β signaling in mTECs is masked by the loss of TGF β -driven tT reg cell induction in mice with systemic TGF β RI kinase inhibitor administration.

To directly examine the role of the miR-155-TGF β signaling axis in mTECs and its subsequent impact on tT reg cell development, we opted for a genetic approach by removing one allele of *Tgfb2* specifically in the thymic epithelia of TEC-cKO mice (TEC-cKO/*Tgfb2*^{fl/+}). As shown in Fig. 5, C and D, a significantly enlarged mature CD80^{hi}MHCII^{hi} mTEC population was seen in TEC-cKO/*Tgfb2*^{fl/+} mice compared with TEC-cKO mice. Accor-

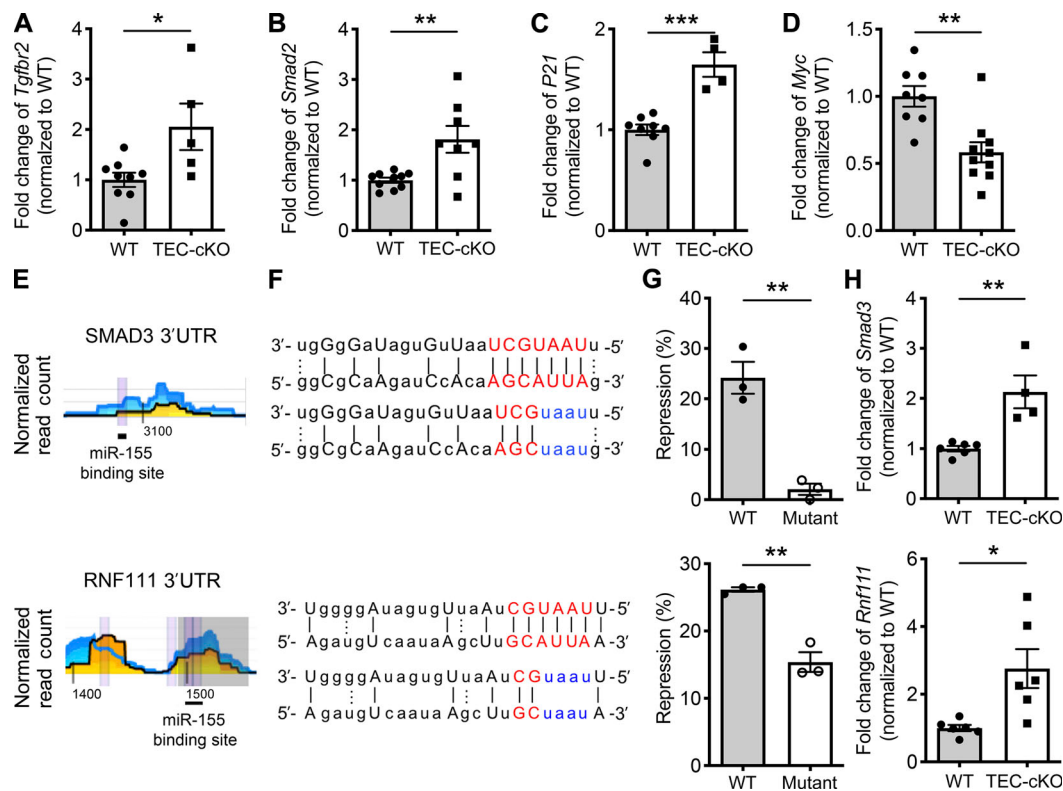


Figure 4. miR-155 regulates TGFβ signaling in mature mTECs via targeting multiple key components. (A–D) qPCR analyses for the expressions of *Tgfb2* (A), *Smad2* (B), *P21* (C), and *Myc* (D) in sorted CD80^{hi}MHCII^{hi} mature mTECs from 5–6-wk-old WT and TEC-cKO mice. The n-fold changes on the basis of each corresponding WT controls were shown. **(E and F)** High-throughput sequencing of RNAs isolated by cross-linking immunoprecipitation analyses (the underlying numbers represent the nucleotide position related to the start of the 3'UTR (E) and sequence alignments (F) of the putative miR-155 binding sites in 3'UTR of SMAD3 (upper) and RNF111 (lower). Mutations of the corresponding miR-155 target sites are shown in blue. **(G)** Ratios of repressed luciferase activity of cells with SMAD3 3'UTR (upper) and RNF111 3'UTR (lower) with or without mutations in the seed sequences in the presence of miR-155 compared with cells transfected with empty vector. **(H)** qPCR analyses of the expressions of *Smad3* (upper) and *Rnf111* (lower) in sorted CD80^{hi}MHCII^{hi} mature mTECs from 5–6-wk-old WT and TEC-cKO mice. Each symbol represents an individual mouse, and the bar represents the mean. Data are pooled from at least three independent experiments. Results of Student's *t* test: *, *P* < 0.05; **, *P* < 0.01; ***, *P* < 0.001.

driving tT reg cell development, substantially augmented tT reg cell frequencies were also detected in accompany with an increase in the mature mTEC population in TEC-cKO mice containing *Tgfb2* heterozygosity (Fig. 5, E and F). Notably, the frequencies of mature mTECs and tT reg cells in TEC-cKO/*Tgfb2*^{f/f} mice remained lower than those in WT controls (Fig. 5, C–F), indicating that additional miR-155 targets independent of TGFβ signaling might also contribute to the mTEC and tT reg cell phenotypes observed in TEC-cKO mice. Nevertheless, our data effectively demonstrate that miR-155 supports mTEC maturation and the resultant tT reg cell development by restricting TGFβ signaling.

In the past decade, the role of miRNAs in controlling diverse immune responses has become a focal point of intense investigation. Previously, we have shown that the same miRNA can impose its gene regulatory effect in different immune cell subsets, enabling them to play their specialized roles in producing a concerted response to a particular environmental stimulus (Cho et al., 2018). Therefore, while numerically speaking, it seems that combining the loss of miR-155 in T cells and TECs largely accounts for the tT reg cell phenotype in mice with the germline miR-155 deletion, it remains possible that impaired function of other immune cells such as dendritic cells (DCs) could also

contribute to the reduction in T reg cell numbers seen in miR-155 null mice. After all, miR-155 deficiency is already known to diminish the antigen-presenting and costimulatory capacities of DCs (Rodriguez et al., 2007), and like mTECs, DCs are also crucial for optimal tT reg cell generation (Perry et al., 2014; Proietto et al., 2008). Still, despite our accumulating knowledge of miRNA-mediated gene regulation in different immune cell populations, relatively limited efforts have been spent on understanding how miRNAs impact the immune system by shaping the environmental cues supplied by nonimmune cells. Here, our results demonstrate that miR-155 can facilitate T reg cell development in the thymus by ensuring the proper maturation of the thymic medulla and, thus, suggest that additional attention be paid to stromal cells residing in the same micro-environments in order to better understand the biological impact of a given miRNA on a selective immunological process.

Materials and methods

Mice

miR-155^{f/f} mice (Hu et al., 2014) were bred with CD4-Cre mice (Lee et al., 2001) and FOXN1-Cre mice (Gordon et al., 2007),

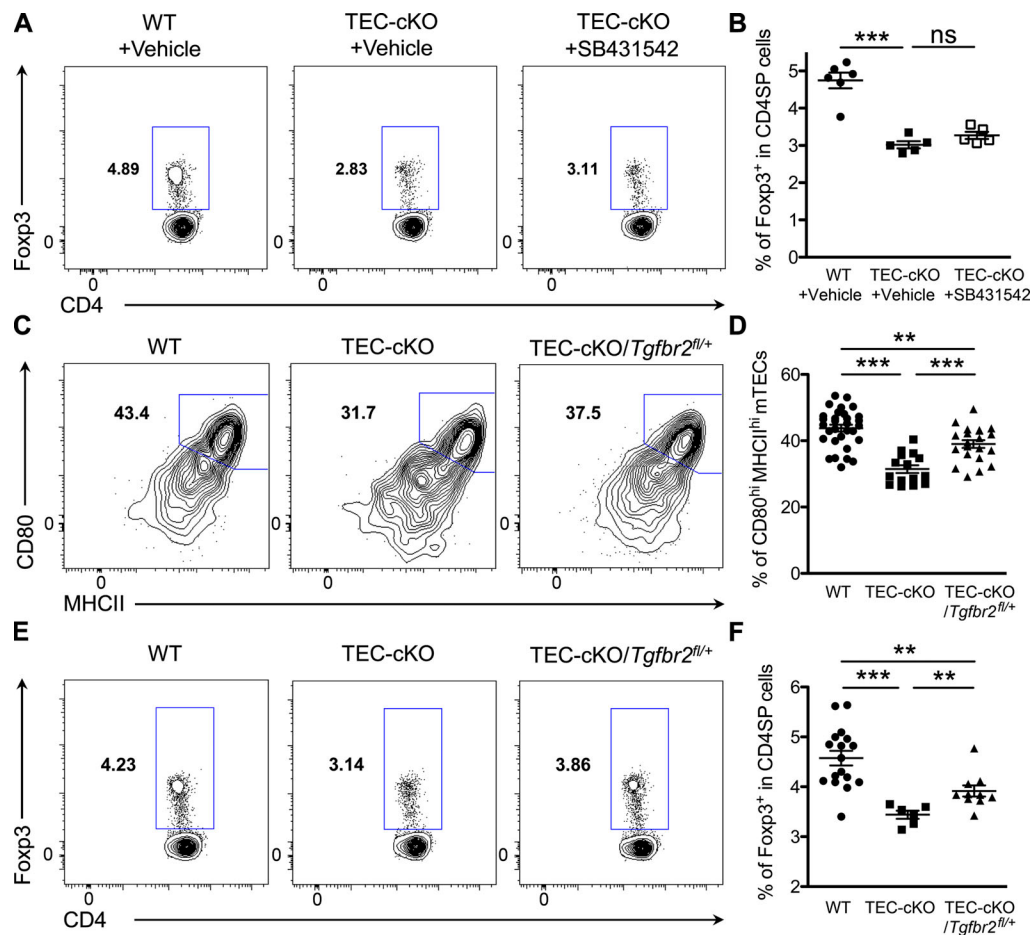


Figure 5. *Tgfr2* heterozygosity in TECs restores mature mTEC and tT reg cell phenotypes in TEC-cKO mice. (A and B) FACS analyses (A) and frequencies (B) of Foxp3⁺ T reg cells in the thymus of 6–8-wk-old WT and TEC-cKO mice treated daily with the TGFβRI kinase inhibitor SB431542 or vehicle only via i.p. injection from day 1 to day 6 and analyzed 12 d after the first injection. (C and D) FACS analysis (C) and frequencies (D) of CD80^{hi}MHCII^{hi} mature mTECs in the thymus of 5–6-wk-old WT, TEC-cKO, and TEC-cKO/*Tgfr2*^{fl/+} mice. (E and F) FACS analysis (E) and frequencies (F) of Foxp3⁺ T reg cells in the thymus of 5–6-wk-old WT, TEC-cKO, and TEC-cKO/*Tgfr2*^{fl/+} mice. Each symbol represents an individual mouse, and the bar represents the mean. Data are pooled from at least three independent experiments. Results of Student's *t* test: **, *P* < 0.01; ***, *P* < 0.001; ns, not significant.

respectively, to obtain mice with T cell-specific deletion (T-cKO) and TEC-specific deletion (TEC-cKO) of miR-155. *Tgfr2*^{fl} mice (Levéen et al., 2002) were bred with TEC-cKO mice to obtain TEC-cKO/*Tgfr2*^{fl/+} mice. Unless otherwise indicated, 5–6-wk-old mice were used. All mice were maintained and handled in accordance with the Institutional Animal Care and Use Guidelines of University of California, San Diego and National Institutes of Health Guidelines for the Care and Use of Laboratory Animals and the Animal Research: Reporting of In Vivo Experiments Guidelines.

Tissue preparation

For T cell analysis, single-cell suspensions of thymus and spleen were prepared by slide mechanical disruption. TECs were isolated as previously described (Seach et al., 2013). In brief, thymus was mechanically disrupted and digested with DNase I (Sigma-Aldrich) and Liberase (Roche). The resulting single-cell suspension was filtered and washed once in magnetic-activated cell sorting buffer. In some experiments, cells were used immediately for FACS analysis. Otherwise, density-gradient centrifugation was performed for TEC enrichment before FACS

analysis or sorting. Briefly, cells were resuspended in 2 ml of 1.115 g/ml isotonic Percoll and overlaid with 1.5 ml of 1.065 g/ml isotonic Percoll, followed by a layer of 1 ml of PBS. Samples were then centrifuged at 600 *g* at 4°C for 30 min with the brakes off. The thymic stroma accumulated between the top and middle layers and was collected and washed for subsequent FACS analysis or cell sorting.

Flow cytometry and antibodies

For FACS analysis, cells were first stained with Ghost Dye Red 780 (Tonbo Biosciences) followed by surface antibody staining for CD4 (RM4-5), CD8a (53-6.7), CD45 (30-F11), EpCAM (G8.8), Ly51 (6C3), Biotinylated UEA-1 (Vector Labs), CD80 (16-10A1), MHCII (M5/114.15.2), TCRβ (H57-597), Streptavidin (eBioscience), CD44 (IM7), CD62L (MEL-14), CD25 (PC61), GITR (DTA-1). Intracellular staining for Foxp3 (FJK-16s), Ki67 (SolA15), AIRE (5H12), CTLA-4 (UC10-4B9), CCL21 (59106), DCLK1 (Abcam polyclonal ab31704) and AlexaFluor-488-conjugated donkey anti-rabbit IgG Ab (Poly4064) was completed after fixation and permeabilization with Foxp3/Transcription Factor Staining Kit according to

manufacturer protocol (Tonbo). Cells were fixed in 2% paraformaldehyde before analysis by flow cytometry. For T reg cell analysis, total tT reg cells were gated on Foxp3⁺ cells from the CD4⁺CD8⁻ population. For nascent tT reg cells, CD73⁻ cells were first gated from the CD4⁺CD8⁻ population, followed by Foxp3⁺ gating. As for TEC analysis, cells were first gated/sorted from the CD45⁺EpCAM⁺ population followed by using Ly51 and UEA-1 to separate cTECs (Ly51⁺UEA-1⁻) and mTECs (Ly51⁻UEA-1⁺). Within the mTECs, immature (or mTEC^{lo}) and mature (or mTEC^{hi}) mTECs were separated by the expression levels of CD80 and MHCII. Moreover, AIRE and CCL21 were used to identify the CCL21⁺AIRE⁻ mTEC subset while DCLK1 was used to label the thymic tuft cells. FACS data were collected by BD LSRFortessa or BD LSRFortessa X-20 (BD Biosciences) and analyzed by FlowJo (Tree Star). Finally, BD BD FACSAria Fusion (BD Biosciences) was used for cell sorting.

Quantitative PCR analysis

Different thymic cell subsets, CD45⁺(CD45⁺EpCAM⁻), cTEC (CD45⁺EpCAM⁺Ly51⁺UEA-1⁻), mTEC (CD45⁺EpCAM⁺Ly51⁻UEA-1⁺) cells, immature mTEC (CD45⁺EpCAM⁺Ly51⁻UEA-1⁺CD80^{lo}MHCII^{lo}), and mature mTEC (CD45⁺EpCAM⁺Ly51⁻UEA-1⁺CD80^{hi}MHCII^{hi}) cells were sorted on a FACSAria Fusion (BD Biosciences), and then total mRNA was isolated using a miRNeasy kit (Qiagen) according to the manufacturer's instructions. To determine expression of miR-155, TaqMan MicroRNA Assay (Thermo Fisher Scientific) was performed. For other gene detection, cDNA was generated using iScript cDNA Synthesis Kit (Bio-Rad), followed by quantitative real-time PCR (qPCR) reactions using SYBR Green PCR Mix (Thermo Fisher Scientific). The primer sequences used were as follows: *Tgfb2*: 5'-GAGAAGCCGCATGAAGTCTG-3' (F), 5'-CATGAAGAAAGTCTCGCCCG-3' (R); *Smad2*: 5'-ATATAGGAAGGGGAGTGCGC-3' (F), 5'-AAACGGCTTCAAACCCCTGG-3' (R); *Smad3*: 5'-ACTTGGACC TACAGCCAGTC-3' (F), 5'-TGCATTCCGGTTGACATTGG-3' (R); *Rnf111*: 5'-CAGCCTTCCACAGTGTGAGA-3' (F), 5'-GGTGTGCTAATG CATGATGG-3' (R); *Myc*: 5'-CACTCACCAGCACAACTACG-3' (F), 5'-GTTCTCTCTGACGTTCCA-3' (R); *P21*: 5'-GCAGATCCACAGCGA TATCC-3' (F), 5'-CAACTGCTCACTGTCCACGG-3' (R); *Aire*: 5'-AGG TCAGCTTCAGAGAAAACCA-3' (F), 5'-TCATTCCAGCACTCAGT AGA-3' (R); *pri-miR-155*: 5'-ACCCTGCTGGATGAACGTAG-3' (F), 5'-CATGTGGGCTTGAAGTTGAG-3' (R); *Gapdh*: 5'-CGTCCCGTAGAC AAAATGGT-3' (F), 5'-TCAATGAAGGGGTGCTTGAT-3' (R); *pri-miR-29a*: 5'-AACCGATTTCAGATGGTGCT-3' (F), 5'-AAGCCTTCTCTG GAAGTGGAC-3' (R).

In vitro RANKL stimulation

CD80^{lo}MHCII^{lo} immature mTECs were sorted on a FACSAria Fusion (BD Biosciences) into complete RPMI, followed by stimulation with 500 ng/ml recombinant mouse RANKL (BioLegend) for 6 h at 37°C.

In vivo mouse treatment

For in vivo RANKL blockade, anti-RANKL antibody (IK22/5) or isotype control antibody (RTK2758) was administrated to 4-wk-old mice at a dose of 250 µg in PBS every other day via i.p. injection for a total of three injections. Mouse thymi were collected for further study 1 d after the last injection. For systemic administration of TGFβRI kinase inhibitor SB431542 (Selleckchem;

10 mg per kg body weight, 0.2 mg in 100 µl of 2% DMSO + 30% PEG300 + PBS) or vehicle control was administrated via i.p. injection once a day from day 1 to day 6 and analyzed 12 d after the first injection.

Luciferase reporter assay

The 3'UTR sequences of SMAD3 and RNF111 were amplified from mouse genomic DNA and cloned into pSiCheck2 vector (Promega). Site-direct mutagenesis (Agilent) was performed to obtain mutants of SMAD3 3'UTR and RNF111 3'UTR, respectively. Indicated 3'UTR WT or mutant plasmids were transfected into HEK293T (ATCC CRL-3216) cells along with either a miR-155-expressing plasmid or a control empty vector. Luciferase activity was determined by the Dual luciferase reporter assay system (Promega) according to the manufacturer's instructions at 24 h after transfection.

Statistical analyses

An unpaired, two-tailed Student's *t* test (or one-way ANOVA for studies with more than two groups) was done on all reported data using Prism software (GraphPad; *, *P* < 0.05; **, *P* < 0.01; ***, *P* < 0.001; ns, not significant). All experiments were performed independently at least three times to ensure the reproducibility of the data.

Online supplemental material

Fig. S1 shows the generation of T-cKO and TEC-cKO mice. Fig. S2 shows the effects of miR-155 deficiency on different mTEC subsets. Fig. S3 shows that tT reg cells developed in TEC-cKO mice exhibited normal phenotypes.

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Supplemental material

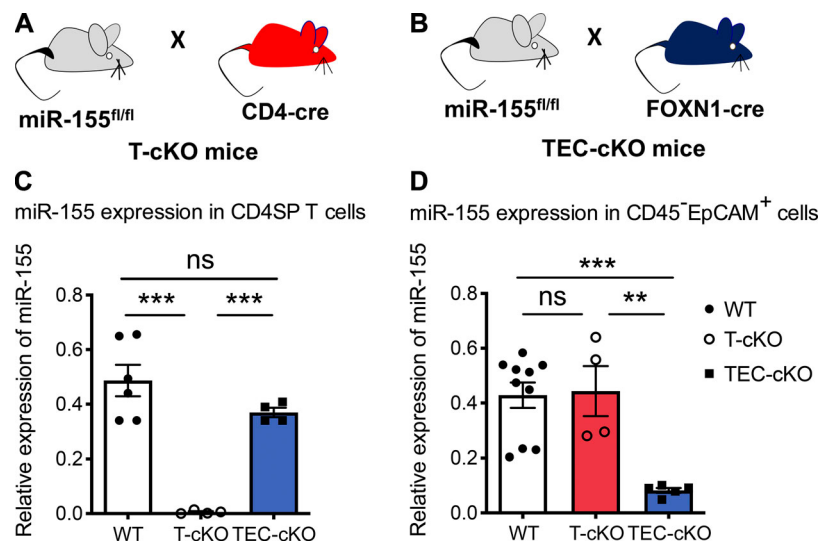


Figure S1. **Generation of T-cKO and TEC-cKO mice. (A and B)** Schematic of the generation of T-cKO (A) and TEC-cKO (B) mice. **(C and D)** qPCR analyses of the expression of miR-155 in CD4SP T cells (CD4⁺CD8⁻; C) and TECs (CD45⁺EpCAM⁺; D) isolated from the thymus of 5–6-wk-old WT, T-cKO, and TEC-cKO mice. Each symbol represents an individual mouse, and the bar represents the mean. Data are pooled from at least three independent experiments. Results of Student's *t* test: **, *P* < 0.01; ***, *P* < 0.001; ns, not significant.

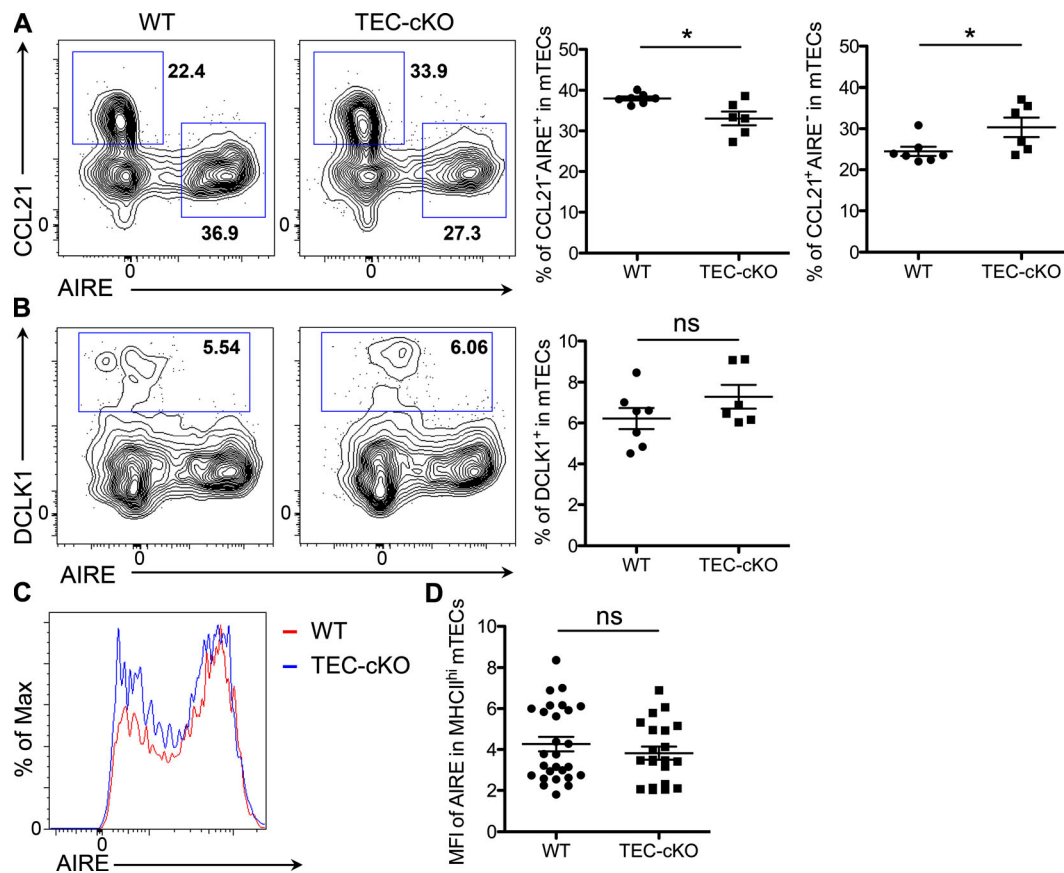


Figure S2. **Effects of miR-155 deficiency on different mTEC subsets.** (A and B) FACS analysis and frequencies of CCL21⁺AIRE⁺ and CCL21⁺AIRE⁻ mTECs (A) as well as DCLK1⁺ tTuft cells (B) from 5–6-wk-old WT and TEC-cKO mice. (C and D) FACS analyses (C) and mean fluorescence intensity (MFI; D) of AIRE in MHCII^{hi} mature mTECs of TEC-cKO mice compared with WT controls. Each symbol represents an individual mouse, and the bar represents the mean. Data are pooled from at least three independent experiments. Results of Student's *t* test: *, *P* < 0.05; ns, not significant.

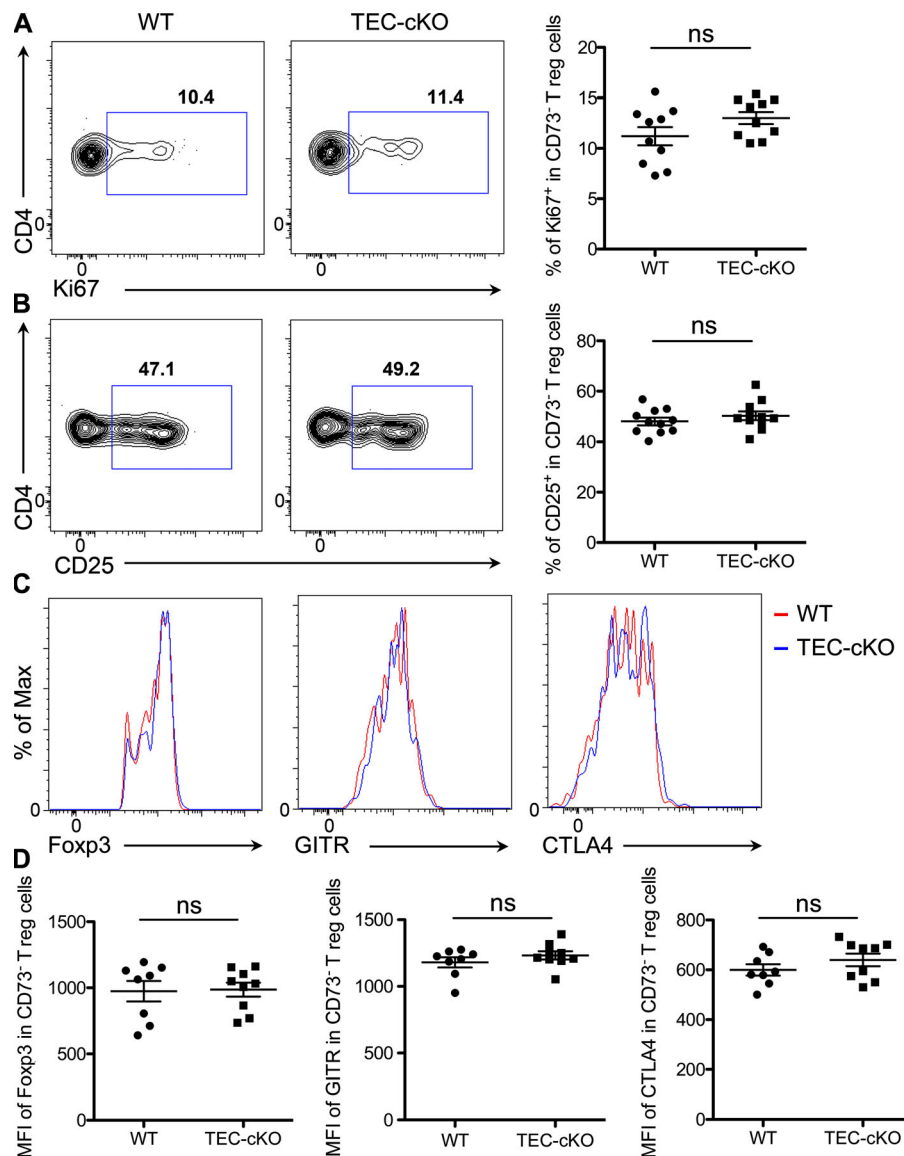


Figure S3. **Despite reduced frequencies, tT reg cells developed in TEC-cKO mice exhibited normal phenotype.** (A and B) FACS analysis and frequencies of Ki67⁺ (A) and CD25⁺ (B) in thymic CD73⁻ Fop3⁺ T reg cells from 5–6-wk-old WT and TEC-cKO mice. (C and D) FACS analyses (C) and mean fluorescence intensity (MFI; D) of Foxp3, GITR, and CTLA4 in thymic CD73⁻ Fop3⁺ T reg cells from 5–6-wk-old WT and TEC-cKO mice. Each symbol represents an individual mouse, and the bar represents the mean. Data are pooled from at least three independent experiments. Results of Student's *t* test: ns, not significant.