**Foxn1** Is Dynamically Regulated in Thymic Epithelial Cells during Embryogenesis and at the Onset of Thymic Involution

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**Abstract**

Thymus function requires extensive cross-talk between developing T-cells and the thymic epithelium, which consists of cortical and medullary TEC. The transcription factor **Foxn1** is the master regulator of TEC differentiation and function, and declining **Foxn1** expression with age results in stereotypical thymic involution. Understanding of the dynamics of **Foxn1** expression is, however, limited by a lack of single cell resolution data. We have generated a novel reporter of **Foxn1** expression, **Foxn1G**, to monitor changes in **Foxn1** expression during embryogenesis and involution. Our data reveal that early differentiation and maturation of cTEC functionality, and proportional expansion of FOXN1-negative TEC in both cortical and medullary sub-lineages. Cortex-specific down-regulation of **Foxn1** between 1 and 3 months of age may therefore be a key driver of the early stages of age-related thymic involution.

**Introduction**

The thymus is required for T cell commitment, differentiation and repertoire selection. These functions are mediated by the epithelial component of the thymic stroma, which is composed of cortical (c) and medullary (m) thymic epithelial cells (TECs). cTECs and mTECs are spatially separated and functionally distinct. The early stages of T cell development, from commitment to positive selection, occur in the cortex, while central tolerance—including negative selection and T regulatory cell induction—is imposed in the medulla [1].
The transcription factor FOXN1 is required throughout ontogeny for TEC production and function. Null mutations in Foxn1 result in the nude phenotype, characterised by hairlessness and athymia in mice, rats and humans [2–6]. The nude thymic primordium forms normally, but TEC development is blocked at the thymic epithelial progenitor cell (TEPC) stage [7, 8]. FOXN1 is also required to maintain the postnatal thymus [9–11]. Age-related thymic involution, a stereotypical process of age-related degeneration, is associated with a progressive decline in Foxn1 expression [9, 10, 12]. Overexpression of Foxn1 in young mice is sufficient to delay involution [10], and TEC-specific up-regulation of Foxn1 in the fully involuted thymus is sufficient to drive robust regeneration that returns the architecture, gene expression and function of the organ to a pre-involution state [13]. Regulation of Foxn1 mRNA expression is thus a primary component of the mechanism of age-related thymic involution [13].

Orchestration by FOXN1 of the gene expression program that controls TEC differentiation and function depends on precise control of Foxn1 expression levels [9, 14]. During development, lineage progression of both TEC sub-lineages requires an increasing amount of FOXN1, such that each successive stage of development occurs at a certain threshold dose [14]. Low-level Foxn1 expression is permissive for exit from the TEPC state and initiation of TEC differentiation, but insufficient to support acquisition of specialist TEC functions including the attraction of lymphoid progenitors [14]. Maintenance of TEC in the adult thymus is similarly sensitive to FOXN1 level, such that reduction of Foxn1 expression to 30% of wild-type (WT) levels results in premature involution [9]. Despite its importance in establishing and maintaining functional TEC, understanding of the dynamics of Foxn1 expression level remains limited due to a lack of single cell data.

To interrogate Foxn1 expression at single cell resolution, we generated and analyzed a novel transgenic mouse line in which activation of the Foxn1 promoter is faithfully reported by GFP expression. Our data reveal that Foxn1 is differentially expressed and dynamically regulated in both the fetal and adult thymus. In particular, initiation of age-related thymic involution occurs concurrently with cTEC-specific down-regulation of Foxn1 and the rapid proportional expansion of a population of Foxn1-negative TEC. Collectively, our data implicate dynamic regulation of Foxn1 as a requirement for both TEC sub-lineage development and the initiation of age-related thymic involution, and suggest that reduced cTEC functionality resulting from decreased Foxn1 expression is a primary instigator of the age-associated decline in thymic cellularity and output.

Materials and Methods

Animal Research

Ethics statement. This study was carried out in strict accordance with UK Home Office guidelines, as established in the ANIMALS (SCIENTIFIC PROCEDURES) ACT 1986. The experimental protocol was approved by the UK Home Office under Project Licence PPLL60/4435.

Mice. Male C57BL/6 or Foxn1G/− mice were used for isolation of postnatal TEC. For timed matings, Foxn1G/− females were housed with CBA males and noon of the day of the vaginal plug was assumed to be E0.5. All animals were housed and bred at University of Edinburgh animal facilities.

Genotyping. Mice were genotyped by PCR using the following primers (shown 5’ to 3’):
Forward: CAAGTCCTCGTTCAGCATCA; Reverse: GCTTCTCGTACTTGCCGTTC.

Flow cytometry
Fetal and postnatal thymi were prepared for flow cytometric analysis and sorting as previously described [13–15]. Briefly, postnatal thymi were dissociated in RPMI-1640 HEPES medium.
(Life Technologies) with 1.25mg/ml collagenase D (Roche) and 0.05mg/ml DNase I (Roche) for 3x15 minutes at 37°C. The remaining tissue was digested in RPMI-1640 HEPES medium with 1.25mg/ml collagenase/dispare (Roche) and 0.05mg/ml DNase I for 30 minutes at 37°C. CD45+ cells were depleted using α-CD45 magnetic beads (Miltenyi Biotech) in conjunction with LS columns (Miltenyi Biotech) and a QuadroMACS (Miltenyi Biotech) according to the manufacturer’s instructions. CD45+ cells were not depleted for analyses requiring absolute numbers of TEC. Antibodies used in flow cytometry are shown in S1 Table. Isotype controls were included in analyses where appropriate. DAPI was used to discriminate dead cells in experiments not involving intracellular staining. Lineage+ cells (Lineage = CD3, CD4, CD8, Ter119, CD11c, CD31) were excluded (except for Fig 1C). For Ki67 and active caspase-3 analyses, GFP-positive and GFP-negative TEC were isolated by FACS prior to analysis by flow cytometry. Fixable viability dye eFluor450 (eBioscience) was used to identify dead cells. Samples were fixed with BD CytoFix/CytoPerm (BD Biosciences) and permeabilized using BD Perm/Wash (BD Biosciences) according to the manufacturer’s instructions. All flow cytometry data were analyzed using FlowJo Version 9.7.7 (Tree Star, Inc).

**RT-qPCR**

100 cells of the required phenotype were sorted by flow cytometry into 10μl CellsDirect 2x Reaction Mix (Life Technologies) and 4U SUPERase In RNase inhibitor (Ambion). cDNA was synthesized and pre-amplified in one step by addition of 1μl CellsDirect SuperScript III reverse transcriptase / Platinum Taq mix (Life Technologies) and target-specific primers at a final concentration of 50nM. Primer sequences are shown in S2 Table. Thermal cycling conditions were as follows: 50°C for 15 minutes; 95°C for 2 minutes; 18 cycles of 95°C for 15 seconds, 60°C for 4 minutes. qPCR was conducted using the Roche Universal Probe Library (Roche) and LightCycler 480 real-time PCR instrument (Roche) according to the manufacturer’s instructions. Amplification conditions were as follows: 95°C for 5 minutes; 40 cycles of 95°C for 10 seconds, 60°C for 20 seconds. Data were normalised to the geometric mean of three housekeepers (Hprt, Ywhaz, Hmbs). All samples were run in triplicate and no RT and no template controls were included in all experiments. Data were analyzed using LightCycler 1.5 software (Roche) and the ΔCt method.

**Statistical analysis**

Statistical analysis was performed using Student’s T-test. p-values < 0.05 were considered significant. Errors shown are standard deviations throughout. Sample sizes of at least n = 3 independent biological experiments were used for statistical analyses. For adult analyses, n is a single mouse or a single group of mice. For fetal analyses, n is a pool of genetically identical TEC isolated from a single litter.

**Results**

**Generation and validation of a Foxn1-eGFP reporter mouse line**

To determine Foxn1 expression levels in individual TEC, we generated a transgenic mouse line in which a LoxP site-flanked cassette containing the eGFP cDNA, an IRES element, the cDNA encoding puromycin resistance and a transcriptional stop sequence, was knocked into intron 1b of the Foxn1 locus (NCBI Gene ID: 15218) by homologous recombination in ES cells generating a revertible functionally null allele, Foxn1G (Fig 1A and S1 Fig). Preliminary analysis demonstrated GFP expression in the fetal thymus and adult hair shafts of Foxn1G/G and Foxn1G/G mice (not shown). To test fidelity of reporter expression, we analyzed fetal TEC at
day 13.5 of embryonic development (E13.5). At this stage, 99.6±0.3% of EpCAM+ epithelial cells were GFP+ (Fig 1B), consistent with earlier reporter and expression analyses [16, 17] and with the finding that all TEC in the adult thymus are derived from a Foxn1+ lineage [11]. As expected, GFP was not expressed in T-cells or endothelial cells at any developmental stage (shown for 1 month old in Fig 1C). We next compared Foxn1 mRNA and eGFP levels in four
subsets of postnatal TEC: cTEC MHC Class II<sup>hi</sup> (cTEC<sub>hi</sub>), cTEC MHC Class II<sup>lo</sup> (cTEC<sub>lo</sub>), mTEC MHC Class II<sup>hi</sup> (mTEC<sub>hi</sub>) and mTEC MHC Class II<sup>lo</sup> (mTEC<sub>lo</sub>), defined using the cell-surface markers UEA1 (mTEC), Ly-51 (cTEC) and MHC Class II (MHCII). As previously described [14, 18], MHCII<sup>hi</sup> TEC expressed more Foxn1 mRNA than MHCII<sup>lo</sup> TEC in both cortex and medulla, and cTEC expressed Foxn1 more highly than mTEC of equivalent MHCII status (Fig 1D and 1E). GFP expression in each TEC subset correlated closely with Foxn1 mRNA levels (Fig 1E and 1F). These data validated the Foxn1<sup>G/+</sup> reporter line and demonstrated its capacity to reliably detect differences in Foxn1 expression of as little as 1.7-fold (Fig 1F).

**Heterogeneity of Foxn1 expression is established during fetal development**

To investigate the origin of the differential Foxn1 expression characteristic of different postnatal TEC sub-lineages, Foxn1<sup>G/+</sup> thymi were analyzed at E13.5, E15.5 and E17.5 (Figs1B and 2). By E13.5, the thymus and parathyroid domains of the common primordium have resolved into...
two discrete organs, preventing contamination of TEC analyses by Foxn1-negative parathyroid cells. At each stage, more than 99% of EpCAM+ TEC were GFP+ (Figs 1B and 2A, [20]) and MHCIIhi cells expressed more GFP than MHCIIlo cells (Fig 2B). At E17.5, when an appreciable UEA1+ population is first apparent in Foxn1G/+ embryos, GFP expression was higher in CD205+ cTEC than in UEA1+ mTEC with equivalent MHCII levels (Fig 2C and 2D). Downregulation of Foxn1 expression in Foxn1hi TEP may therefore be required for mTEC divergence or expansion, or alternatively, UEA1+ TEC may be derived from a Foxn1lo lineage. In sum, the graded expression of Foxn1 observed in the adult thymic epithelium is present from the earliest stages of TEC differentiation, linking Foxn1 dosage to the early differentiation and maturation of cortical and medullary TEC.

**Foxn1 negative TEC in the postnatal thymus**

The above analyses revealed a GFP-negative subpopulation in postnatal Foxn1G/+ TEC, representing 24.8±7.6% of total TEC in 3 month-old mice (Fig 3A). These Foxn1-negative cells were present in the MHCIIhi and MHCIIlo fractions of both cTEC and mTEC, but were predominately MHCIIlo (90.7±3.1% of total GFP-negative cells were MHCIIlo) (Fig 3B). Downregulation of Foxn1 is a primary component of age-related thymic involution [13], and the presence of Foxn1-negative postnatal thymic epithelial stem cell (TESC), derived from a Foxn1-negative lineage has also been suggested [21]. The GFP-negative TEC observed in Foxn1G/+ mice may therefore result from down-regulation of Foxn1 in Foxn1-positive TEC or expansion of a rare Foxn1-negative TEC population (>99% of EpCAM+ TEC are GFP+ throughout fetal thymic development, Figs 1B and 2A). Previous lineage tracing of Foxn1-negative TEC using a Foxn1::Cre mouse line suggested that these cells arise from Foxn1-positive precursors [11]. However, Cre was driven from a Foxn1 promoter fragment, which may not precisely recapitulate endogenous Foxn1 expression. Therefore, to test the origins of the postnatal GFP(Foxn1)-negative TEC identified here, Foxn1Cre mice were crossed with Gt(ROSA)26Sorwatm4(ACTB-tdTomato,EGFP)Luo/J reporter mice (mT/mG mice) [22] in which tdTomato expression is extinguished and eGFP expression activated upon Cre-mediated recombination of the reporter locus. Of TEC isolated from 9 week to 3 month-old Foxn1Cre;tmT/mG mice, 0.5±0.3% were GFP-negative (shown for 3 month-old in Fig 3C). In comparison, 24.8±7.6% of TEC in 3 month-old Foxn1G/+ mice are GFP negative (Fig 3A). These data establish that expansion of a minor TEC subpopulation that never expresses Foxn1 cannot account for the GFP(Foxn1)-negative TEC population present in the

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**Fig 3. Foxn1flkGFP TEC subpopulations emerge postnataally in both cTEC and mTEC compartments.**

(A-C) Flow cytometric analysis of thymi from 12 week-old adult Foxn1G/+ (A,B) or Foxn1Cre;tmT/mG (C) mice, for the markers shown. Data are shown after gating against Lineage- and on total EpCAM+ cells (A-C) and further gating on GFPneg cells (B). Absolute number of GFPneg cells in 3 month-old mice, 1.31±10^4±3.01±10^3. WT, wild type. Red line in (A) shows FMO. (A,B) n = 6, (C) n = 3 independent biological experiments.

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adult $\text{Foxn1}^{G/+}$ thymus: if this was the case, the proportion of GFP$^+$ cells in the $\text{Foxn1}^{C/G+};\text{mT/mG}$ reporter mice and $\text{Foxn1}^{G/+}$ mice would be equivalent. Thus, the GFP-negative TEC in the adult $\text{Foxn1}^{G/+}$ thymus must be generated by down-regulation of $\text{Foxn1}$ in previously $\text{Foxn1}$-positive TEC.

Proportional expansion of $\text{Foxn1}$ negative TEC is associated with thymic involution

Down-regulation of $\text{Foxn1}$ has been implicated as a cause of thymic involution [9, 10, 12, 13], but the dynamics of $\text{Foxn1}$ expression during ageing have not been reported at the single cell level in defined TEC subsets. Age-related decline in thymus weight and cellularity occurred with the same kinetics in $\text{Foxn1}^{G/+}$ as in WT mice (Fig 4A). Therefore, to investigate the relationship between the $\text{Foxn1}$-negative TEC population and postnatal thymus function, we interrogated GFP ($\text{Foxn1}$) expression at a range of ages spanning the initiation and progression of thymic involution. The absolute number of both GFP$^+$ and GFP$^-$ TEC decreased in $\text{Foxn1}^{G/+}$ mice between 1 month and 2 years old ($p = 0.0007$, $p = 0.0003$ for GFP$^+$ and GFP$^-$ TEC respectively) (Fig 4B). However the proportion of GFP-negative TEC increased; at 1 month, 3 months, 12 months and 24 months old, 18.0±.4.1%, 24.8±7.6%, 28.7±3.2% and 29.3±7.9% of

**Fig 4.** Proportional expansion of $\text{Foxn1}^{\text{neg}}$ TEC occurs at the onset of age-related thymic involution. (A) Thymus involution in $\text{Foxn1}^{G/+}$ mice occurs with normal kinetics. (C, D, E) Flow cytometric analysis of $\text{Foxn1}^{G/+}$ thymi at the ages shown; the proportion of GFP$^-$ TEC increases with age. Red (C, D) and grey (E) lines show FMO. (B) Absolute number of GFP$^+$ and GFP$^-$ TEC isolated from $\text{Foxn1}^{G/+}$ thymi at the timepoints shown. Absolute numbers are as follows: 1 month; GFP$^+$ $4.94 \times 10^5 \pm 1.81 \times 10^4$, GFP$^-$ $1.06 \times 10^3 \pm 3.07 \times 10^3$. 3 months; GFP$^+$ $4.03 \times 10^4 \pm 8.86 \times 10^3$, GFP$^-$ $1.31 \times 10^4 \pm 3.01 \times 10^3$. 12 months; GFP$^+$ $1.55 \times 10^4 \pm 2.36 \times 10^3$, GFP$^-$ $6.55 \times 10^3 \pm 1.88 \times 10^3$. 24 months; GFP$^+$ $4.52 \times 10^3 \pm 2.75 \times 10^3$, GFP$^-$ $1.81 \times 10^3 \pm 1.10 \times 10^3$. (A, B, C) $n = 3$, (D, E) $n = 2$ independent biological experiments.

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TEC were GFP-negative respectively (p = 0.05, p = 0.43, p = 0.90 for 1 month v 3 months, 3 months v 1 year, 1 year v 2 years respectively)(Fig 4C). Thus, the proportion of GFP-negative TEC increases rapidly between 1 month and 3 months postnatal and is stable thereafter, consistent with a recent report based on antibody staining [20]. These two phases correlate inversely with the initially fast and then slower decrease in thymus size characteristic of the onset and progression of thymic involution (Fig 4A), suggesting that involution is a consequence of a proportional increase in subfunctional, Foxn1-negative TEC. However, our data demonstrate that age-related down-regulation of Foxn1 is not uniform in all TEC; some TEC continue to express high Foxn1 (GFP) levels even in very old mice (Fig 4D and 4E).

Initiation of thymic involution correlates with decreased Foxn1 expression in cTEC

The rapid proportional expansion of GFP-negative TEC between 1 month and 3 months of age suggested a link between this process and the initiation of age-related thymic involution. We therefore investigated whether age-associated changes in GFP expression occurred with the same kinetics in cTEC and mTEC. The proportion of GFP-negative TEC was equivalent in the cortex and medulla at 1 month and 3 months, and increased in both subsets between 1 and 3 months of age (p = 0.0031, p = 0.0004 for cTEC and mTEC respectively) (Fig 5A). In mTEC, this proportional change reflected a significantly decreased number of GFP+ mTEC in 3- compared to 1-month old mice (4.13x10^4±1.59x10^4 and 2.47x10^4±2.77x10^3 GFP+ mTEC at 1 month and 3 months respectively, p = 0.018). In cTEC, both GFP+ and GFP- cells increased numerically between 1 and 3 months of age (p = 0.004, p = 0.0001 respectively) (Fig 5B).

To investigate how thymic epithelial composition is altered during early involution, we analysed cell death in GFP+ and GFP- cTEC and mTEC at 1 month and 3 months, and also determined proliferation rate in the same TEC subsets. At 1 month, a higher proportion of GFP+ than GFP- mTEC was apoptotic (51.7%±2.5% and 25.0%±3.0% of GFP+ and GFP- mTEC respectively, p = 0.0003), whereas at 3 months the frequency of apoptosis was equivalent in GFP+ and GFP- mTEC (21.6%±10.6% and 19.9%±3.3% of GFP+ and GFP- mTEC respectively, p = 0.803) (Fig 5C and 5D). Additionally, GFP+ mTEC were less proliferative than GFP- mTEC in 6 week-old mice (p = 0.05) (Fig 5D). These data indicate that apoptosis of GFP+ mTEC contributes to the proportional expansion of GFP- mTEC between 1 and 3 months of age, and suggest that a difference in proliferative index may also contribute to the increased proportion of GFP- mTEC at 3 months old.

In cTEC, the proportion of apoptotic cells was not significantly different in the GFP+ and GFP- subsets at 1 month (13.9%±9.6% and 9.6%±3.1% of GFP+ and GFP- cTEC respectively, p = 0.50), but was increased in GFP+ compared to GFP- cTEC at 3 months (2.65%±0.6% and 7.31%±1.2% of GFP+ and GFP- cTEC respectively, p = 0.004). Additionally, GFP+ cTEC were more proliferative than GFP- cTEC at 6 weeks old (p = 0.009, Fig 5D). Thus in contrast to mTEC, the increased proportion of GFP+ cTEC in 3 month old mice cannot be explained by apoptosis of GFP+ cells and differences in proliferation rate between GFP+ and GFP- TEC. Further analysis indicated that the median level of GFP expression in GFP+ cTEC declines between 1 and 3 months (p = 0.008, Fig 5E and 5F). These data suggest that the accumulation of GFP cTEC in 3 month old mice primarily results from specific downregulation of Foxn1 in a GFP+ cTEC subpopulation that subsequently persists in the epithelium. This broad Foxn1 down-regulation was not observed in GFP+ mTEC. Thus, our data collectively indicate that the earliest stages of involution are driven by different mechanisms in the cortex and medulla.
Down-regulation of Foxn1 results in loss of cTEC function at the onset of involution

In keeping with the observed decrease in median GFP expression levels in GFP⁺ cTEC, RT-qPCR analysis of WT mice showed significantly reduced Foxn1 mRNA levels in cTEC (p = 0.008) but not in mTEC (p = 0.485) between 1 and 3 months of age (Fig 6A). FOXN1 in cTEC regulates the expression of several factors required for attraction, specification and proliferation of thymocytes, including Dll4, Ccl25 and KitL [13, 14]. RT-qPCR analysis showed a significant reduction in expression of Dll4 and Ccl25 in WT cTEC between 1 and 3 months old (p = 0.013 and p = 0.001 respectively) (Fig 6B and 6C). KitL and Ccr11, in contrast, were not
significantly down-regulated ($p = 0.224$ and $p = 0.362$ respectively) (Fig 6D and 6E). In the medulla, Foxn1 expression was maintained between 1 and 3 months of age but the expression of Aire and Cd80 decreased ($p = 0.029$ and $p = 0.017$ respectively) (Fig 6F and 6G). Cd40 was significantly down-regulated in cTEC ($p = 0.041$), but not in mTEC ($p = 0.128$) (Fig 6H). Thus cortex-specific down-regulation of Foxn1 between 1 and 3 months of age may be a key driver of the early stages of involution.

Discussion

Using the Foxn1$^G$ reporter mouse, we have conducted a single cell resolution analysis of the level and dynamics of Foxn1 expression in defined subsets of fetal and postnatal TEC. Our data reveal that early fetal mTEC express lower levels of Foxn1 than their cTEC counterparts, suggesting either that downregulation of Foxn1 expression in FOXN1-high TECPC is required for mTEC divergence or expansion, or that a Foxn1-low mTEC progenitor is present from the earliest stages of thymus organogenesis. We further show that the onset of thymic involution correlates with rapid proportional expansion of Foxn1-negative TEC in both TEC sublineages, and broad down-regulation of Foxn1 in cTEC. Finally, we show that c-TEC specific downregulation of Foxn1 is associated with reduced expression of genes required to promote thymocyte commitment, proliferation and differentiation, suggesting that age-related thymic involution is initiated by reduced cTEC function.

Lineage progression of postnatal mTEC correlates with regulation of Foxn1 expression

The lineage relationships between the different subpopulations of postnatal mTEC have not yet been clearly defined in vivo. However adult mTEC MHCII$^{lo}$ can give rise to mTEC MHCII$^{hi}$
when combined with single positive thymocytes in reaggregate thymic organ culture [23]. Similarly, 70% of postnatal Cldn3/4hi SSEA1+ TEC, which have some mTEC progenitor activity, are MHCIIlo/neg in 4 week old mice [24]. Our single cell resolution data show that almost all MHCIIlo mTEC express less Foxn1 than MHCIIhi mTEC, and therefore indicate that the transition from MHCIIhi mTEC to MHCIIlo mTEC is associated with up-regulation of Foxn1. Recently, it was also shown that MHCIIlo cTEC can give rise to both cTEC and mTEC in ectopically grafted reaggregates. Consistent with mTEClo being precursors of mTECchi TEC, UEA1+ cells appeared in these grafts before MHCII was up-regulated in either sub-lineage [25]. Our data show that, per cell, Foxn1 is expressed at an equivalent or higher level in Ly-51+MHCIIlo TEC compared to UEA1+MHCIIlo TEC, demonstrating that differentiation of UEA1-positive cells from their UEA1-negative precursors does not require up-regulation of Foxn1. This observation is consistent with the Foxn1-independent divergence of the mTEC lineage in embryogenesis [14], and demonstrates that lineage progression of postnatal mTEC proceeds via two distinct stages, which are respectively independent of and dependent on up-regulation of Foxn1 expression.

AIRE+ mTEC in the postnatal thymus are MHCIIhi and CD80hi and are usually considered to be terminally differentiated [23]. However recent lineage tracing has revealed the existence of a post-AIRE population in the thymic medulla [26]. These cells previously expressed AIRE and maintain intermediate levels of tissue specific antigen (TSA) expression, but are MHCIIhi and CD80lo [26, 27]. Involucrin, a marker of terminally differentiated epidermis, is enriched in post-AIRE mTEC, and Hassall’s corpuscles contain cells that previously expressed AIRE [27]. Post-AIRE TEC are therefore thought to be a distinct stage of terminal mTEC development. However, our data suggest an alternative possibility, that at least some of these cells are sub-functional TEC produced as a result of involution-associated down-regulation of Foxn1 and MHCII.

Thymic involution is initiated in cTEC

The total cellularity of the postnatal thymus is dependent on the number and function of TEC [14, 28]. TEC functionality is particularly important, as increased epithelial proliferation cannot fully compensate for sub-optimal gene expression [14]. Our analyses show that at the onset of involution, down-regulation of Foxn1 occurs more broadly in cTEC than in mTEC. cTEC-specific targets of FOXN1 include Ccl25 and Dll4, which are required for attraction and specification of thymocytes respectively. Reduced expression of FOXN1 targets in cTEC may thus be the primary driver of declining total cellularity at the initiation of involution. This is consistent with the observation that cortical thinning occurs early in involution and precedes CMJ disorganisation and altered mTECcTEC ratio when Foxn1 is downregulated postnatally [9]. Identification of factors able to modulate FOXN1 expression or function specifically in cTEC could thus provide a novel approach to preventing or reversing thymic involution.

Rescue of Foxn1 negative cTEC may facilitate thymus regeneration

The size of the involuted thymus can be at least transiently restored by several approaches, including sex steroid ablation (SSA) and up-regulation of FOXN1. SSA results in thymic rebound through relieving the direct repression of Dll4 by androgen receptor (AR). Although Dll4 is expressed in both cTEC and endothelial cells, AR represses Dll4 specifically in cTEC [29]. SSA-mediated rebound therefore occurs, at least in part, by modulating cTEC activity. cTEC function may also be an important feature of FOXN1-mediated thymic regeneration. When FOXN1 activity is up-regulated in the thymic epithelium of old mice, all TEC populations (cTEC MHC\textsuperscript{hi}, cTEC MHC\textsuperscript{lo}, mTEC MHC\textsuperscript{hi}, mTEC MHC\textsuperscript{lo}) expand. However, there is
no increase in proliferation of cTEC MHCIIhi cells. One possible source of new MHCIIhi cTEC is a postnatal progenitor, either bipotent or restricted to the cTEC lineage. Alternatively, some new cTEC MHCIIhi could be derived from cTEC that have down-regulated Foxn1 and MHCII. These cells accumulate in the thymic epithelium as involution progresses, suggesting that functional rescue of Foxn1-negative TEC may contribute to thymic rebound. Thus, competence to restore optimal gene expression to sub-functional cTEC may partly explain both the notable regenerative capacity of the thymus and the transient nature of some thymus-regenerating interventions.

Taken together, our single cell-level analyses show that Foxn1 is tightly and differentially regulated at two stages of thymus ontogeny: first as FOXN1-low mTEC emerge during early thymus organogenesis; and second at the onset of age-related involution. These data suggest that precise manipulation of Foxn1 expression levels will be necessary for both directed differentiation of the TEC sub-lineages, and sustained endogenous regeneration of the thymus.

**Supporting Information**

S1 Fig. Characterization of the Foxn1<sup>GFP</sup> allele.

(PDF)

S1 Table. Antibodies used for flow cytometry

(PDF)

S2 Table. Primers used for RT-qPCR

(PDF)

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**Author Contributions**

Conceived and designed the experiments: KON CCB. Performed the experiments: KON NB CT CSN HJV CDP FHS TG. Analyzed the data: KON CT CSN NB HJV TG CCB. Wrote the paper: KON CSN NB CCB.

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