

Evidence for a stepwise program of extrathymic T cell development within the human tonsil

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The development of a broad repertoire of T cells, which is essential for effective immune function, occurs in the thymus. Although some data suggest that T cell development can occur extrathymically, many researchers remain skeptical that extrathymic T cell development has an important role in generating the T cell repertoire in healthy individuals. However, it may be important in the setting of poor thymic function or congenital deficit and in the context of autoimmunity, cancer, or regenerative medicine. Here, we report evidence that a stepwise program of T cell development occurs within the human tonsil. We identified 5 tonsillar T cell developmental intermediates: (a) CD34⁺CD38^{dim}Lin⁻ cells, which resemble multipotent progenitors in the bone marrow and thymus; (b) more mature CD34⁺CD38^{bright}Lin⁻ cells; (c) CD34⁺CD1a⁺CD11c⁻ cells, which resemble committed T cell lineage precursors in the thymus; (d) CD34⁻CD1a⁺CD3⁻CD11c⁻ cells, which resemble CD4⁺CD8⁺ double-positive T cells in the thymus; and (e) CD34⁻CD1a⁺CD3⁺CD11c⁻ cells. The phenotype of each subset closely resembled that of its thymic counterpart. The last 4 populations expressed *RAG1* and *PTCRA*, genes required for TCR rearrangement, and all 5 subsets were capable of ex vivo T cell differentiation. TdT⁺ cells found within the tonsillar fibrous scaffold expressed CD34 and/or CD1a, indicating that this distinct anatomic region contributes to pre-T cell development, as does the subcapsular region of the thymus. Thus, we provide evidence of a role for the human tonsil in a comprehensive program of extrathymic T cell development.

Introduction

The development of human T cells can be divided into discrete stages, ranging from that for multipotent progenitors to that for naive T cells. The expression of CD4 and/or CD8 divides thymocytes into the categories of double-negative, immature single-positive, double-positive (DP), and single-positive cells (1, 2). Based on TCR gene rearrangements and differentiation capabilities, double-negative thymocytes are divided into the least mature CD34⁺CD38^{dim}CD1a⁻ cells and the progressively more mature CD34⁺CD38^{bright}CD1a⁻ and CD34⁺CD1a⁺ cells (2, 3). CD34⁺CD1a⁺ thymocytes acquire CD4 and then CD8 as they lose CD34, to become CD34⁻ DP cells. Finally, CD3⁺ DP thymocytes lose either CD4 or CD8 to become single-positive naive T cells (1).

Development of a healthy T cell repertoire is dependent on a functional thymus. However, the potential for human extrathymic lymphoid tissue to augment T cell development is poorly understood but may be important in the setting of poor thymic function or congenital deficit. Previous reports of extrathymic T cell development in humans have relied primarily on the identification of cells expressing genes associated with TCR loci rearrangement in the intestine or bone marrow. Indeed, CD7⁺CD10⁺ cells expressing pre-TCR α (*PTCRA*), recombination activating gene 1 (*RAG1*), and *RAG2* mRNA and varying levels of *TCRB* gene recombination have

been identified in postnatal bone marrow (4). However, recent reports were unable to identify CD34⁺CD45RA⁺CD7⁺CD10⁺ cells in the same tissue (5, 6), indicating that it is unlikely that the CD7⁺CD10⁺ bone marrow cells identified include early CD34⁺ pre-T cells. Similarly, CD5⁺CD7⁺CD3⁻ cells have been identified in neonatal intestine, and *PTCRA*, *TdT* (also known as *DNTT*), and *RAG2* mRNA can be detected in whole lysates of these tissues (7); yet, whether or not CD34⁺ T cell progenitors reside there is unclear. Finally, several reports have identified TdT⁺ cells in extrafollicular zones of the human tonsil near fibrous scaffolding (8–11). However, little is known about the function or phenotype of these cells. Thus, no complete program of human T cell development has been described in an extrathymic tissue.

Human tonsils and lymph nodes contain resident CD34⁺ cells (12, 13). However, to date, the CD34⁺ compartment of human secondary lymphoid tissue has been characterized primarily by its ability to contribute to NK cell development. Specifically, it has been divided into 2 maturational stages of NK cell intermediate, distinguished by the expression of CD117. Stage 1 pro-NK cells are minimally defined as CD34⁺CD117⁻, whereas stage 2 pre-NK cells are defined as CD34⁺CD117⁺. Each of these CD34⁺ populations has the differentiation potential for T cells, DCs, and NK cells (13). However, the minimal definitions of stage 1 and 2 cells do not account for the immense phenotypic heterogeneity within each population (14). Whether stage 1 or 2 NK precursors contain distinct subpopulations, each capable of giving rise to other hematopoietic lineages, is unknown.

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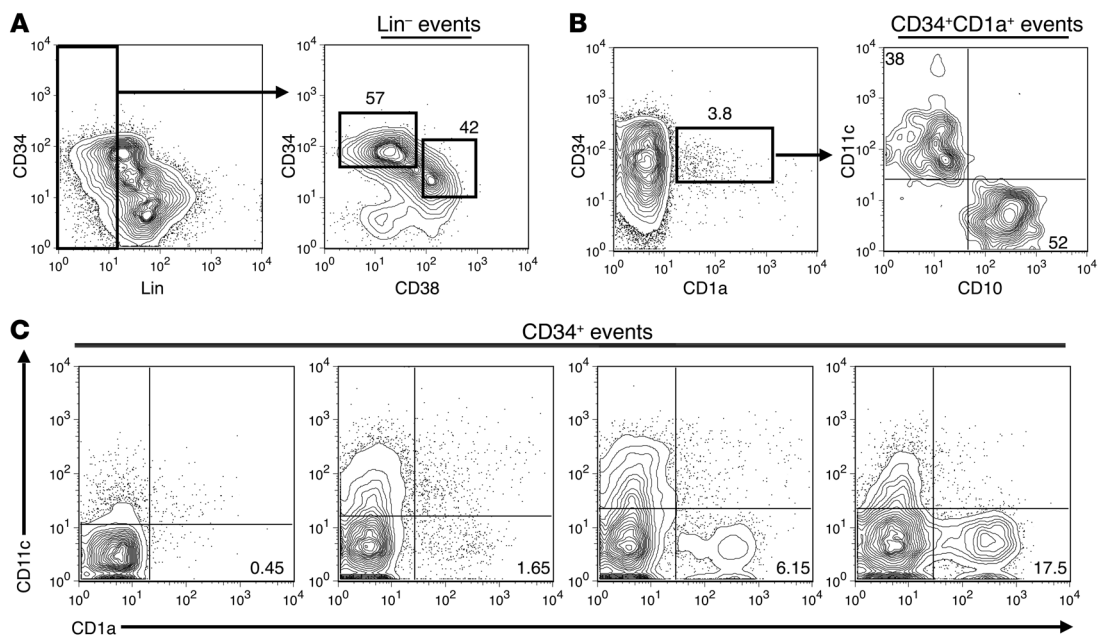


Figure 1

The human tonsil contains $CD34^+CD38^{dim}Lin^-$, $CD34^+CD38^{bright}Lin^-$, and $CD34^+CD1a^+CD11c^-$ cells. **(A)** Expression of CD34 and lineage markers (CD11c, BDCA-2, CD117, CD161, CD19, CD3, CD1a) on CD34-enriched tonsillar cells (left). Total lymphocytes were gated on Lin^- events and analyzed for their expression of CD34 and CD38 (right). The number above each gate indicates the mean percentage of CD34⁺ cells falling within that gate. **(B)** Expression of CD1a on CD34-enriched tonsillar cells (left). The number above each gate indicates the mean percentage of CD34⁺ cells falling within that gate. Events were gated on $CD34^+CD1a^+$ cells and analyzed for their coexpression of CD10 and CD11c (right). The number within the top left and bottom right quadrants represents the mean percentage of $CD34^+CD1a^+$ cells that are defined as either $CD11c^+$ or $CD10^+$, respectively. **(C)** The percentage of CD34⁺ cells that are defined as $CD34^+CD1a^+CD11c^-$ in 4 representative human pediatric tonsils. Plots are gated on total CD34⁺ cells. Numbers in the bottom right quadrant indicate the percentage of CD34⁺ cells that are $CD1a^+CD11c^-$ in that particular donor. All data are from a representative tonsil of **(A)** 3, **(B)** 4, or **(C)** 24 individual donors.

Here, we provide evidence for a role of human tonsil in a step-wise program of extrathymic T cell development. We identify 5 T cell developmental intermediates within the tonsil, each one with T cell differentiation potential, as well as a phenotype closely resembling the corresponding populations in the thymus. Furthermore, we discovered that the TdT^+ cells found near the fibrous scaffold of the tonsil express CD34 and/or the pre-T cell marker CD1a, indicating that distinct anatomical regions of the tonsil contribute to the earliest stages of extrathymic T cell development. Finally, we demonstrate that $CD34^{+/-}CD1a^+CD11c^-$ tonsillar and thymic cells retain both T cell and NK cell developmental potential.

Results

CD34⁺CD38^{dim}Lin⁻, CD34⁺CD38^{bright}Lin⁻, and CD34⁺CD1a⁺CD11c⁻ cells reside within the human tonsil. Dim expression of CD38 on CD34⁺ cells identifies multipotent progenitor populations in the bone marrow, thymus, and umbilical cord blood, and increased expression of CD38 on CD34⁺ cells is associated with differentiation (1, 3, 15–17). Analyzing magnetically enriched CD34⁺ tonsillar cells for the expression of CD34 and CD38, together with the lineage (Lin) antigens CD3, CD19, CD117, CD161, BDCA-2, CD11c, and CD1a, revealed that the human pediatric tonsil contains a population of $CD34^+CD38^{dim}Lin^-$ cells as well as a $CD34^+CD38^{bright}Lin^-$ subset. In general, the $CD34^+CD38^{dim}$ subset tended to express higher surface density of CD34 than the $CD34^+CD38^{bright}$ population (Figure 1A), suggesting that the

human tonsil may contain a multipotent hematopoietic progenitor minimally defined as $CD34^+CD38^{dim}Lin^-$ as well as a more mature $CD34^+CD38^{bright}Lin^-$ population.

Commitment to the T cell lineage in the human thymus is associated with the acquisition of CD1a on $CD34^+CD10^+CD7^+$ thymocytes (1). However, CD1a expression is also canonically associated with DC populations (18, 19). Within the human tonsil there was a small but consistent population of $CD34^+CD1a^+$ lymphocytes (Figure 1B), which lacked CD117, CD161, BDCA-2, CD14, CD19, and CD3 (data not shown). However, when analyzed for the expression of CD10 and the DC marker CD11c, $CD34^+CD1a^+$ tonsillar cells expressed either CD11c or CD10 (Figure 1B). Substantial donor-to-donor variation exists in the relative size of the $CD34^+CD1a^+CD11c^-$ subset in the tonsil (Figure 1C). In 24 pediatric donors, $CD34^+CD1a^+CD11c^-$ tonsillar cells constituted between 0.15%–19.4% of the total CD34⁺ population (mean, $3.82\% \pm 1.06\%$; median, 1.47%). No statistical correlation was found between the percentage of $CD34^+CD1a^+CD11c^-$ cells and the age (1–16 years) or gender of the donor.

Tonsillar CD34⁺CD38^{dim}Lin⁻, CD34⁺CD38^{bright}Lin⁻, and CD34⁺CD1a⁺CD11c⁻ cells phenotypically resemble the earliest stages of thymocyte development. To determine whether or not the $CD34^+CD38^{dim}Lin^-$, $CD34^+CD38^{bright}Lin^-$, and $CD34^+CD1a^+CD11c^-$ cells identified in the tonsil represent extrathymic T cell precursors similar to those found in the thymus, each subset from tonsil and thymus was extensively pheno-



typed by flow cytometry (Figure 2). In both the tonsil and thymus, CD34⁺CD38^{dim}Lin⁻ cells expressed CD10, CD33, CD45RA, and HLA-DR as well as variable levels of CD25 and intracellular TdT. Similarly, CD34⁺CD38^{dim}Lin⁻ cells from both tissues lacked expression of CD4, CD8, CD56, and CD116 (Figure 2). However, there are distinct differences between tonsillar and thymic CD34⁺CD38^{dim}Lin⁻ cells. Thymic CD34⁺CD38^{dim}Lin⁻ cells expressed CD2 and most expressed CD5 and CD7, while a small fraction expressed the IL-7 receptor α (CD127) (Figure 2B). The same cells in the tonsil lacked CD2 and CD127, and only a small fraction expressed CD5 or CD7 (Figure 2).

CD34⁺CD38^{bright}Lin⁻ cells from the tonsil and thymus are similar in that both expressed CD2, CD10, and low levels of CD45RA. Similarly, both populations lacked expression of CD4, CD8, CD56, and CD116. However, CD34⁺CD38^{bright}Lin⁻ cells from the tonsil expressed variable levels of CD7 and CD5, and only small amounts of intracellular CD3 ϵ (icCD3 ϵ), whereas those from the thymus were all CD7⁺CD5⁺icCD3 ϵ ⁺. Furthermore, tonsil CD34⁺CD38^{bright}Lin⁻ cells expressed CD33 and variable levels of HLA-DR, which were largely absent on the same cells from the thymus (Figure 2). Despite the differences seen in CD34⁺CD38^{bright}Lin⁻ cells, each subset represents a phenotype that is in many ways intermediate between CD34⁺CD38^{dim}Lin⁻ cells and CD34⁺CD1a⁺CD11c⁻ cells from the respective tissues. In the tonsil, intermediate levels of CD5, CD7, CD25, CD127, HLA-DR, and TdT on CD34⁺CD38^{bright}Lin⁻ exemplify this.

The CD34⁺CD1a⁺CD11c⁻ populations in the thymus and tonsil are remarkably similar. Both expressed the early T cell markers CD2, CD5, CD7, CD10, and icCD3 ϵ . Furthermore, both demonstrated moderate levels of CD127 and intracellular TdT, an enzyme necessary for TCR gene rearrangement. Portions of the CD34⁺CD1a⁺CD11c⁻ subset in both the tonsil and thymus expressed CD4, suggesting that some CD34⁺CD1a⁺ cells may be progressing to the immature single-positive stage, characteristic of normal thymocyte development (1). Like their thymic counterparts, tonsillar CD34⁺CD1a⁺CD11c⁻ cells did not express the GM-CSF receptor (CD116) or HLA-DR, consistent with the hypothesis that they are not DCs (Figure 2).

CD4⁺CD8⁺CD1a⁺ DP pre-T cells reside within the human tonsil. If the human tonsil supports complete T cell development, it should contain DP pre-T cells. However, Nascimbeni et al. have identified peripheral blood CD3⁺ DP T cells that have a mature memory function (20), indicating that CD4 and CD8 alone are not enough to distinguish DP lymphocytes as pre-T cells. CD34-depleted tonsillar or thymic cells and total mononuclear peripheral blood cells were analyzed for expression of CD4, CD8, CD3, and CD1a. Indeed, a population of CD4⁺CD8⁺ cells resides within the human tonsil, and it was divided into CD1a⁺ and CD1a⁻ cells. The tonsillar CD1a⁻ DP cells uniformly expressed CD3, whereas the CD1a⁺ DP cells variably expressed CD3, suggesting they contain CD3⁻ T cell precursors (Figure 3A). Furthermore, thymic DP cells expressed CD1a with similar CD3⁺ expression as seen on the tonsillar CD1a⁺ subset (Figure 3A). Conversely, blood DP cells were CD1a⁻CD3⁺ (Figure 3A), consistent with the findings of Nascimbeni et al., who found that DP memory T cells lack CD1a (20). These findings suggest that expression of CD1a can distinguish among distinct subsets of DP lymphocytes. Furthermore, its expression on a fraction of CD4⁺CD8⁺CD3^{+/+} tonsillar cells, similar to that in the thymus, supports the hypothesis that pre-T cells reside within the tonsil.

In the thymus, CD1a is expressed throughout T cell development until its loss on naive CD3⁺ T cells (1). To investigate the relationship between CD34⁺CD1a⁺CD11c⁻ tonsillar cells and potential downstream T cell intermediates, CD19⁻ tonsillar cells were enriched for CD1a-expressing cells. Like the CD34⁺CD1a⁺ subset described in Figure 1, the CD34⁺CD1a⁺ fraction of the human tonsil can be divided into 2 distinct populations based on its expression of the DC marker CD11c (Figure 3B). CD34⁺CD11c⁻CD1a⁺ tonsillar cells displayed low forward and side scatter properties, suggesting they are small, agranular lymphocytes, whereas the CD11c⁺CD1a⁺ subset displayed higher forward and side scatter (Figure 3B). In contrast to those in the tonsil, CD34⁺CD1a⁺ cells in the human thymus were largely CD11c⁻ and displayed low forward and side scatter (Figure 3B). These data suggest that CD11c expression in the human tonsil is useful for distinguishing large CD1a⁺ DCs from what we believe to be a novel population of small CD11c⁻CD1a⁺ lymphocytes.

Tonsillar cells were next enriched for cells expressing CD34, CD1a, or a combination of both and analyzed for the expression of CD11c, CD34, CD1a, and CD3. It appears that both CD1a⁺ thymic and tonsillar cells lose CD34 expression as they increase CD1a surface density (Figure 3C, left). Furthermore, when the enriched cells were gated on total CD34⁺CD11c⁻ events, a similar progression from CD1a⁺CD3⁻ to CD1a⁺CD3⁺ cells was seen in both tissues (Figure 3C, right). As CD1a expression is seen on thymocytes, spanning from CD34⁺ precursors all the way to CD3⁺ near-mature T cells (1), these data suggest that a similar pattern describes putative T cell developmental intermediates of the tonsil.

Tonsillar CD34-CD1a⁺CD11c⁻ cells express T cell antigens in patterns similar to those of human thymocytes. The relationship of CD34, CD1a, and CD3 as well as the identification of CD1a⁺CD4⁺CD8⁺ DP cells suggest that the human tonsil can support T cell development. After gating on CD34⁺CD11c⁻ tonsil or thymic cells, 3 populations were identified: (a) CD1a⁺CD3⁻, (b) CD1a⁺CD3⁺, and (c) CD1a⁻CD3⁺ cells (Supplemental Figure 1A; supplemental material available online with this article; doi:10.1172/JCI46125DS1). Expression of thymocyte-associated antigens was analyzed on these 3 populations from the tonsil and thymus (Figure 4). In both tissues, the expression of early thymocyte markers, such as CD10 and intracellular TdT, decreases as cells progressed from CD1a⁺CD3⁻ to CD1a⁺CD3⁺, whereas the CD1a⁻CD3⁺ cells did not express these markers in either tissue. In both the tonsil and thymus, CD5 and CD7 increased as cells progress through the 3 identified populations. CD1a⁺CD3⁻ and CD1a⁺CD3⁺ cells from both tissues expressed CD4, and many coexpressed CD8, although donor variation in CD8 expression was observed (data not shown). Finally, cells from all 3 subsets expressed icCD3 ϵ , and surface expression of TCR $\gamma\delta$ and TCR $\alpha\beta$ began at the CD1a⁺CD3⁺ stage in both the thymus and tonsil. Total tonsillar and thymic CD34⁺CD11c⁻CD1a⁺ cells did not express the lineage markers CD56, CD117, CD94, CD16, CD19, CD14, BDCA-2, or CD116, but a small subset expressed CD161 (Supplemental Figure 1, B and C). Thus, total tonsillar CD1a⁺CD11c⁻ cells phenotypically resemble CD1a⁺ cells in the human thymus. Furthermore, early thymocyte antigen loss and the acquisition of mature T cell-associated antigens suggest that CD34⁺CD11c⁻ tonsillar precursors progress from CD1a⁺CD3⁻ cells, which acquire CD3 and then lose CD1a, to become mature CD1a⁻CD3⁺ T cells.

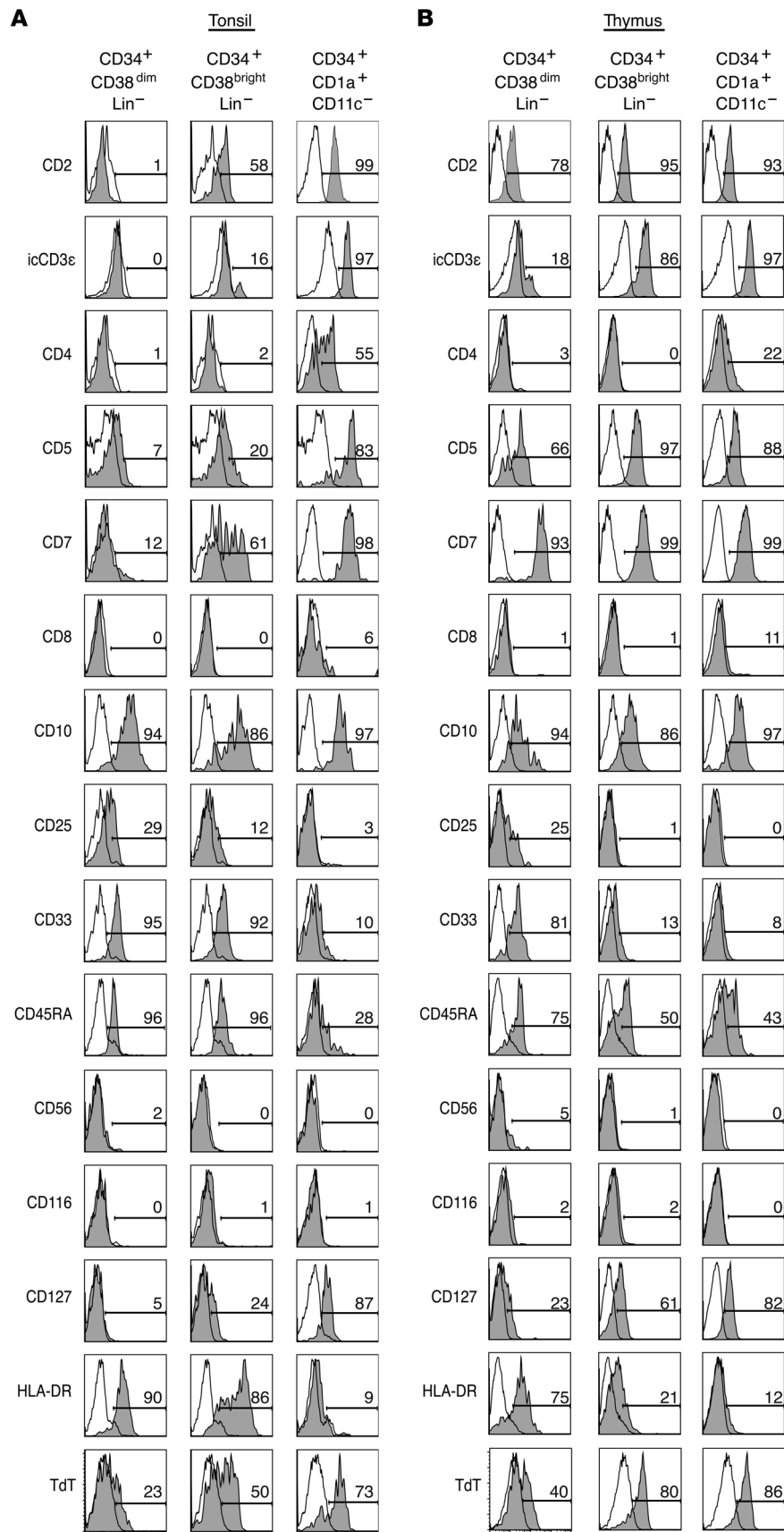


Figure 2

Phenotypic comparison of CD34⁺CD38^{dim}Lin⁻, CD34⁺CD38^{bright}Lin⁻, and CD34⁺CD1a⁺CD11c⁻ cells in human tonsil and thymus. (A) Tonsillar cells were magnetically depleted of CD3- and CD19-expressing cells and were then enriched for CD34⁺ cells. Enriched cells are gated on CD34⁺CD38^{dim}Lin⁻, CD34⁺CD38^{bright}Lin⁻, or CD34⁺CD1a⁺CD11c⁻ events. (B) Thymic cells were magnetically enriched for CD34⁺ cells and then gated on the same 3 populations as in A. Gray shading indicates staining with the indicated antibody, whereas no shading indicates staining with an isotype-matched control antibody. The number above each gate indicates the percentage of events falling within that gate. Data in each histogram is from a representative donor, where $n \geq 3$ for tonsil and $n \geq 2$ for thymic data.

Expression of T cell development-associated genes in putative tonsillar T cell precursors. Specific gene expression patterns in the human thymus can be used to track the development of T cell precursors as they progress from multipotent cells toward mature T cells. For example, *RAG1* and *PTCRA*, both of which are required for successful TCR rearrangement, are expressed early in T cell development and peak at the CD34⁺CD1a⁺ and DP pre-T cell stages (3, 6). Similarly, expression of *BCL-X* has been detected in both murine and human CD4⁺CD8⁺ DP thymocytes, and the *BCL-X_L* (also known as *BCL2L1*) isoform has been shown to be critical for providing necessary antiapoptosis signals in murine DP cells (21, 22). In contrast, *THPOK* is a transcription factor that is essential for development of mature CD4⁺ cells, and its expression in the thymus and periphery is largely restricted to CD3⁺ cells (23).

Thus, we used real-time RT-PCR to assess expression of these 4 genes within putative T cell precursor populations in the human tonsil. Six distinct populations were sorted from human tonsil and thymus, as shown in Figure 5A: (a) CD34⁺CD38^{dim}Lin⁻, (b) CD34⁺CD38^{bright}Lin⁻, (c) CD34⁺CD1a⁺CD3⁻, (d) CD34⁺CD1a⁺CD3⁺, (e) CD34⁺CD1a⁺CD3⁺, and (f) CD34⁺CD1a⁺CD3⁺ cells. The complete gating strategy and representative sort purities are shown in Supplemental Figure 2, and the relative frequencies of each tonsillar subset are shown in Supplemental Table 1. In duplicate experiments, the levels of *RAG1* (Figure 5B), *PTCRA* (Figure 5C), and *BCL2L1* (Figure 5D) mRNA were highest in either population 3 or 4 of the tonsil. Expression of *THPOK* mRNA, on the other hand, was largely restricted

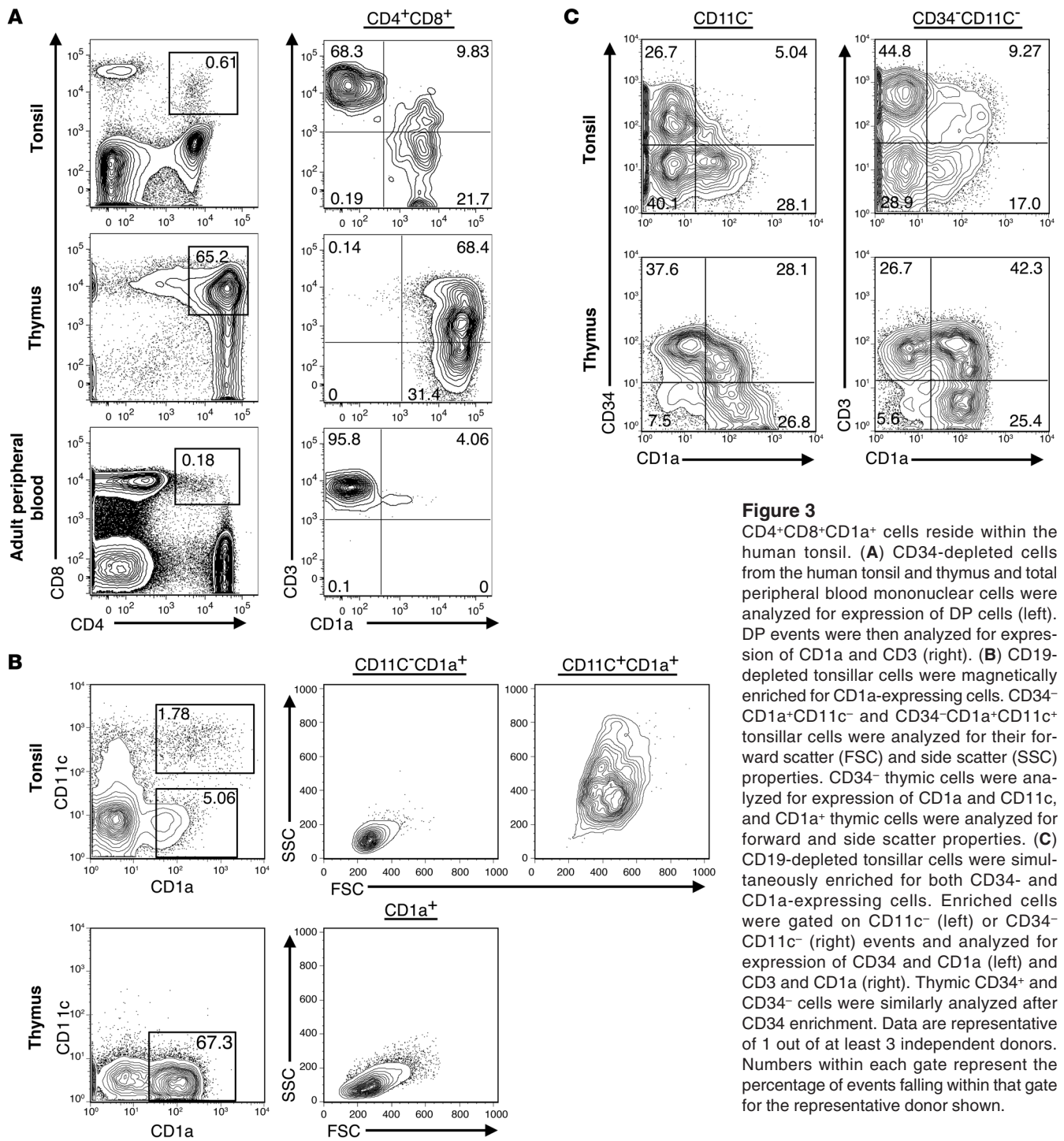


Figure 3

CD4⁺CD8⁺CD1a⁺ cells reside within the human tonsil. **(A)** CD34-depleted cells from the human tonsil and thymus and total peripheral blood mononuclear cells were analyzed for expression of DP cells (left). DP events were then analyzed for expression of CD1a and CD3 (right). **(B)** CD19-depleted tonsillar cells were magnetically enriched for CD1a-expressing cells. CD34⁻CD1a⁺CD11c⁻ and CD34⁻CD1a⁺CD11c⁺ tonsillar cells were analyzed for their forward scatter (FSC) and side scatter (SSC) properties. CD34⁻ thymic cells were analyzed for expression of CD1a and CD11c, and CD1a⁺ thymic cells were analyzed for forward and side scatter properties. **(C)** CD19-depleted tonsillar cells were simultaneously enriched for both CD34- and CD1a-expressing cells. Enriched cells were gated on CD11c⁻ (left) or CD34⁻CD11c⁻ (right) events and analyzed for expression of CD34 and CD1a (left) and CD3 and CD1a (right). Thymic CD34⁻ and CD34⁻ cells were similarly analyzed after CD34 enrichment. Data are representative of 1 out of at least 3 independent donors. Numbers within each gate represent the percentage of events falling within that gate for the representative donor shown.

to the CD3⁺ populations 5 and 6 (Figure 5E). These patterns of gene expression were found to be similar to the patterns noted in the comparable thymic populations (Figure 5, B–E).

In situ localization of tonsillar CD34⁺TdT⁺ and CD1a⁺TdT⁺ cells. TdT⁺ cells have been identified in the human tonsil near the fibrous scaffold (8–11). However, they have yet to be characterized with regard to origin, hematopoietic lineage, or function. Our identification of CD1a⁺CD11c⁻TdT⁺ cells in the tonsil suggests that the previously identified TdT⁺ cells may be extrathymic T cell

precursors. Paraffin-embedded pediatric tonsils were analyzed by immunohistochemistry for the presence of TdT, CD34, CD1a, CD11c, and Rag1. We confirmed that TdT⁺ cells are largely restricted to the extrafollicular regions near the fibrous scaffold of the tonsil (Figure 6A). Further, these cells did not express CD11c (data not shown) but did variably coexpress CD34, CD1a (Figure 6B), and/or nuclear Rag1 (Supplemental Figure 3). Furthermore, CD34⁺CD1a⁺ cells were easily identified in the same region of the tonsil fibrous scaffold (Figure 6B). TdT⁺ cells were

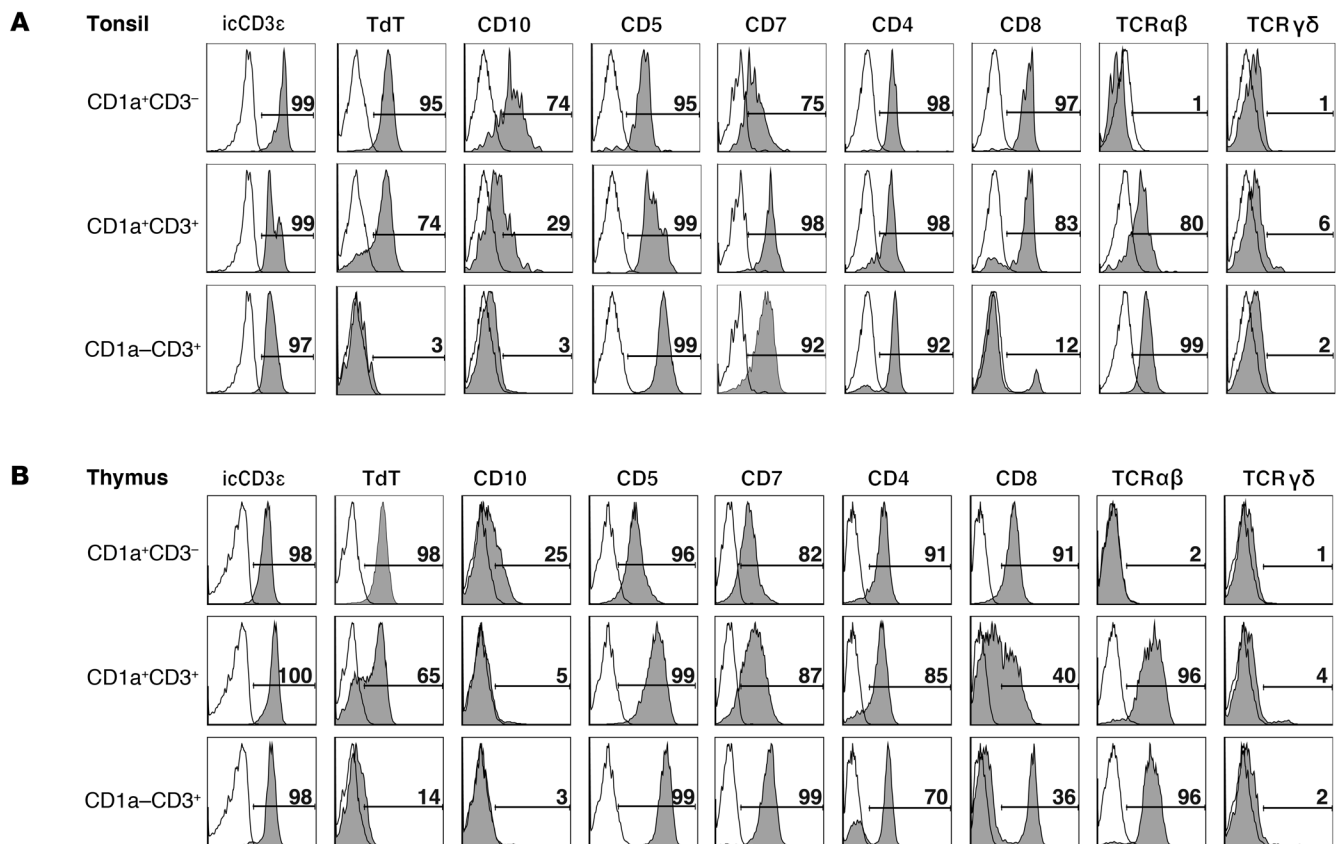


Figure 4

Putative T cell precursors in the human tonsil acquire T-associated antigens in a fashion similar to that of those in the thymus. (A) CD1a-enriched tonsillar cells were gated on CD34⁻11c⁻ events and analyzed for the expression of antigens on 3 populations: CD1a⁺CD3⁻, CD1a⁺CD3⁺, and CD1a⁻CD3⁺. (B) CD34-depleted thymic cells were gated on the same 3 populations and analyzed as in A. Gray shading indicates staining with the indicated antibody, whereas no shading indicates staining with an isotype-matched control antibody. The number listed above each gate indicates the percentage of events falling within that gate. Data in each histogram are from a representative donor, where $n \geq 3$ for tonsil and $n \geq 2$ for thymic data.

identified in both pediatric and adult tonsil. Analyses performed in adult reactive lymph nodes revealed that rare TdT⁺ cells could be identified, but no coexpression of CD1a or CD34 was observed (data not shown). To further examine the microenvironment of tonsillar TdT⁺ cells, we used immunohistochemistry to analyze the expression of the notch ligands delta-like 1 (DL1) and delta-like 4 (DL4) within the tonsil fibrous regions, as notch ligands are essential for complete T cell development (24–27). Both proteins were easily identified (Figure 6C), further supporting the hypothesis that the tonsil may contribute to extrathymic T cell development. Taken together, these data suggest that the TdT⁺ cells previously identified in the tonsil belong to a program of extrathymic T cell differentiation and localize to a specific micro-environment near the tonsillar fibrous scaffold.

Ex vivo T cell differentiation potential of putative extrathymic T cell precursors. To determine whether the putative extrathymic T cell precursors have the capacity to differentiate into mature T cells, the 6 populations were sorted from the tonsil and thymus, as shown in Figure 5A and Supplemental Figure 2, and were cultured on the OP9-DL1 cell line with flt3 ligand (FL) and IL-7. After 26 days, the cells were harvested and analyzed by flow cytometry. To exclude the GFP⁺ OP9-DL1 cells, harvested cells were gated on GFP-CD45⁺

events. Populations 1–3 from the tonsil and thymus expanded 93–270 fold, whereas populations 4–6 only expanded 1–6 fold (Supplemental Table 2). All 6 populations from the tonsil and thymus gave rise to CD3⁺ cells. In the tonsil and thymus, the percentage of GFP⁺CD45⁺ cells that coexpress CD4 and CD8 increased as the cells progressed from population 1 to population 4 (Figure 7). Furthermore, in both the tonsil and thymus, the percentage of GFP-CD45⁺ cells that express CD3 increased as one progressed from population 1 to population 6, as did the percentage of CD3⁺ cells that coexpress a TCR (Figure 7 and Supplemental Figure 4, A and B). When we compared the relative ability of each tonsillar population to generate mature T cells with that of the same populations in the thymus, we found that population 2 and 3 cells from the thymus generated a significantly higher proportion of CD3⁺ cells than did populations 2 and 3 from the human tonsil (population 2, 51.9% ± 7.0% from the thymus vs. 4.1% ± 1.12% from the tonsil, $P < 0.0001$; population 3, 57.23% ± 4.8% from the thymus vs. 32.36% ± 8.1% from the tonsil, $P = 0.0044$; Supplemental Figure 4A). Similarly, we found that thymic populations 2 and 3 generated significantly more CD3⁺ cells that coexpressed a TCR (either TCRαβ or TCRγδ) than did the same populations in the tonsil (population 2, 39.3% ± 8.5% from the thymus vs. 1.41% ± 0.6% from the tonsil, $P < 0.0001$;

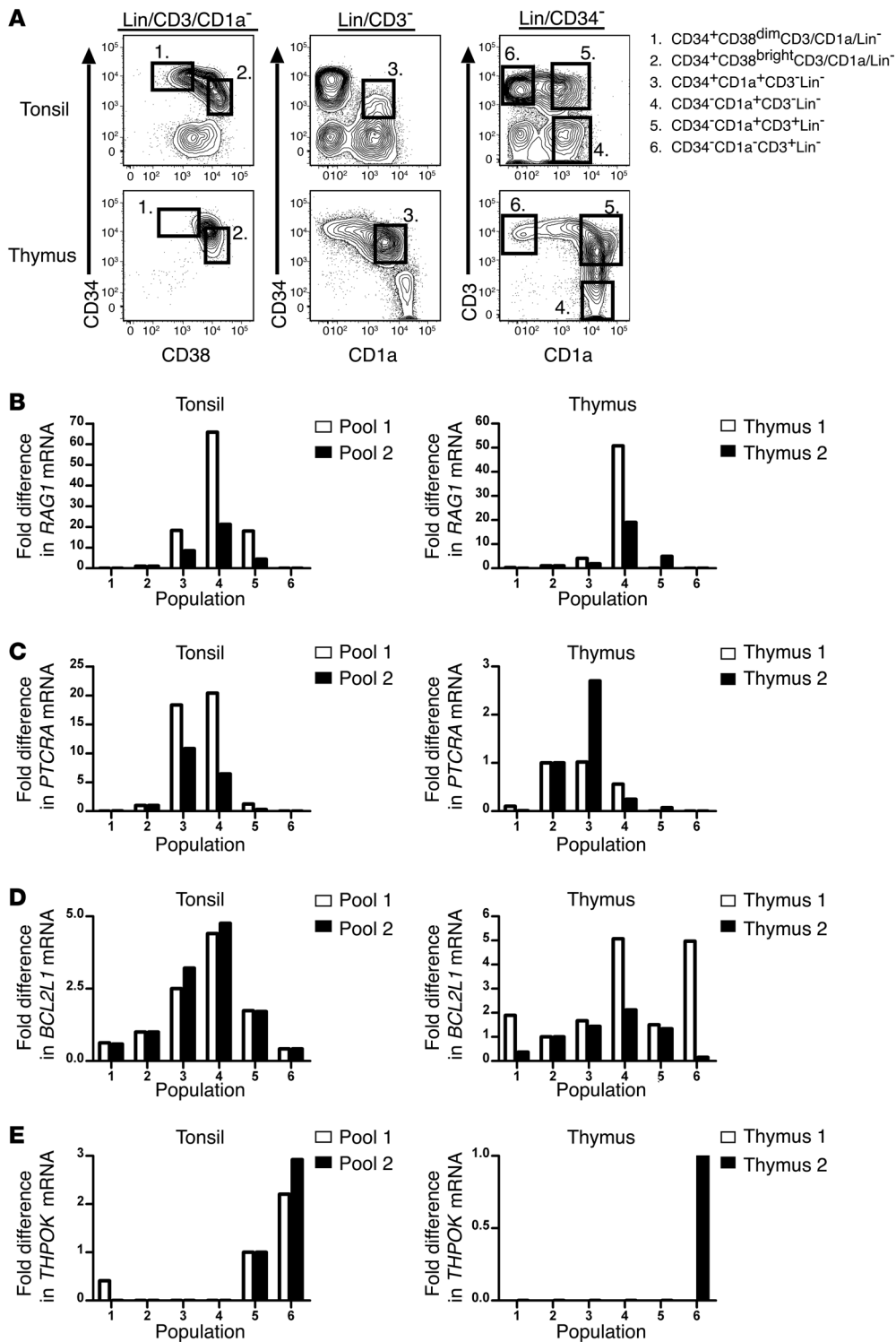


Figure 5 Quantification of gene expression in tonsillar precursor cells by real-time RT-PCR. **(A)** Tonsillar cells were depleted of CD19⁺ cells, and gated on the 6 populations shown. **(B–E)** Expression of the genes *RAG1*, *PTCRA*, *BCL2L1*, and *THPOK* in 6 populations of human tonsillar and thymic cells. Cells were sorted from 8 tonsil donors, and 2 pools were generated for each population consisting of cells from 4 of the 8 donors (see Methods). Cells were sorted from 2 thymic donors. **(B–D)** For analysis of *RAG1*, *PTCRA*, and *BCL2L1*, expression is displayed relative to that of population 2, which was arbitrarily set at 1. **(E)** For *THPOK*, expression is displayed relative to that of population 5 for the tonsil and population 6 for the thymus, each of which was arbitrarily set at 1.

population 3, 44.1% ± 7.2% from the thymus vs. 20.3% ± 5.7% from the tonsil, $P = 0.0041$; Supplemental Figure 4B). No statistical difference was observed in the percentage of GFP-CD45⁺ cells that were CD3⁺ or CD3⁺TCR⁺ when we compared tonsillar populations 1, 4, 5, or 6 with those same populations in the human thymus. Several controls were performed to rule out contamination from mature tonsillar T cells as being responsible for the observed out-

growth of CD3⁺ cells from cultures of populations 1–4. For example, CD19⁺ tonsillar cells were similarly sorted and cultured on the OP9-DL1 cell line. After 26 days, no CD3⁺ cells were generated from the CD19⁺ tonsillar cells. Thus, these data demonstrate that CD34⁺CD38^{dim}Lin⁻, CD34⁺CD38^{bright}Lin⁻, CD34⁺CD1a⁺CD3⁻, and CD34⁻CD1a⁺CD3⁻ tonsillar cells (populations 1–4) contain differentiation potential for T cell development ex vivo.

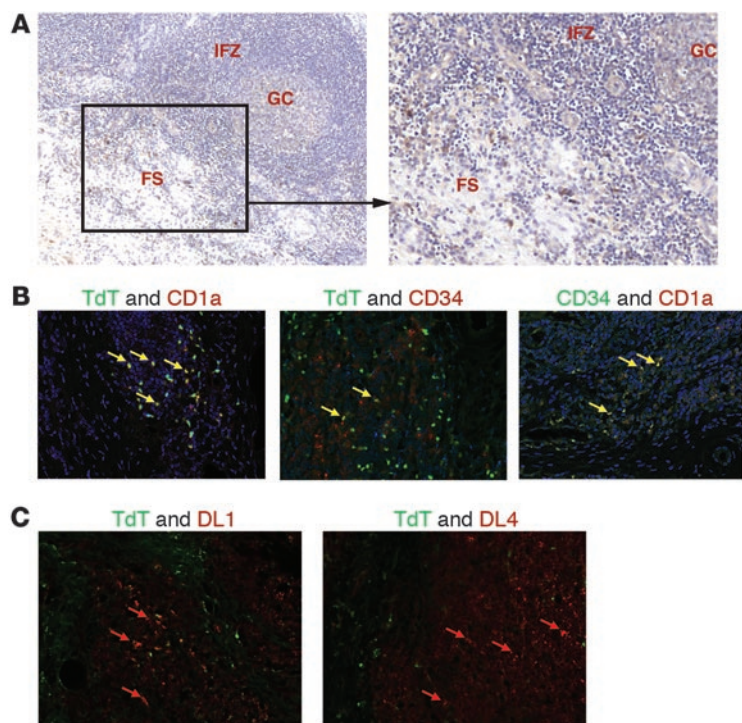


Figure 6

CD1a⁺TdT⁺ cells reside near the fibrous scaffold of the human tonsil. **(A)** Immunohistochemical staining of paraffin-embedded tonsillar sections shows TdT⁺ cells (dark brown) localized to the fibrous scaffold region. The image on the left (original magnification, ×50) is of a representative section, including a germinal center (GC), the surrounding interfollicular zone (IFZ), and an adjacent fibrous scaffold region (FS). The image on the right is of the same tonsil section, but at an original magnification of ×100. **(B)** Cells coexpressing TdT, CD1a, and CD34 are found within the scaffold region of the tonsil. TdT (green) and CD1a (red) coexpression is shown on the left, TdT (green) and CD34 (red) coexpression is shown in the middle, and CD34 (green) and CD1a (red) are shown on the right. Note the yellow cells in each section, which represent cells coexpressing both proteins of interest (yellow arrows). Blue staining indicates hematoxylin counterstain. **(C)** TdT (green), DL1 (red), and DL4 (red) proteins are expressed in the same geographical region of the human tonsil. Original magnification, ×400 (**B** and **C**).

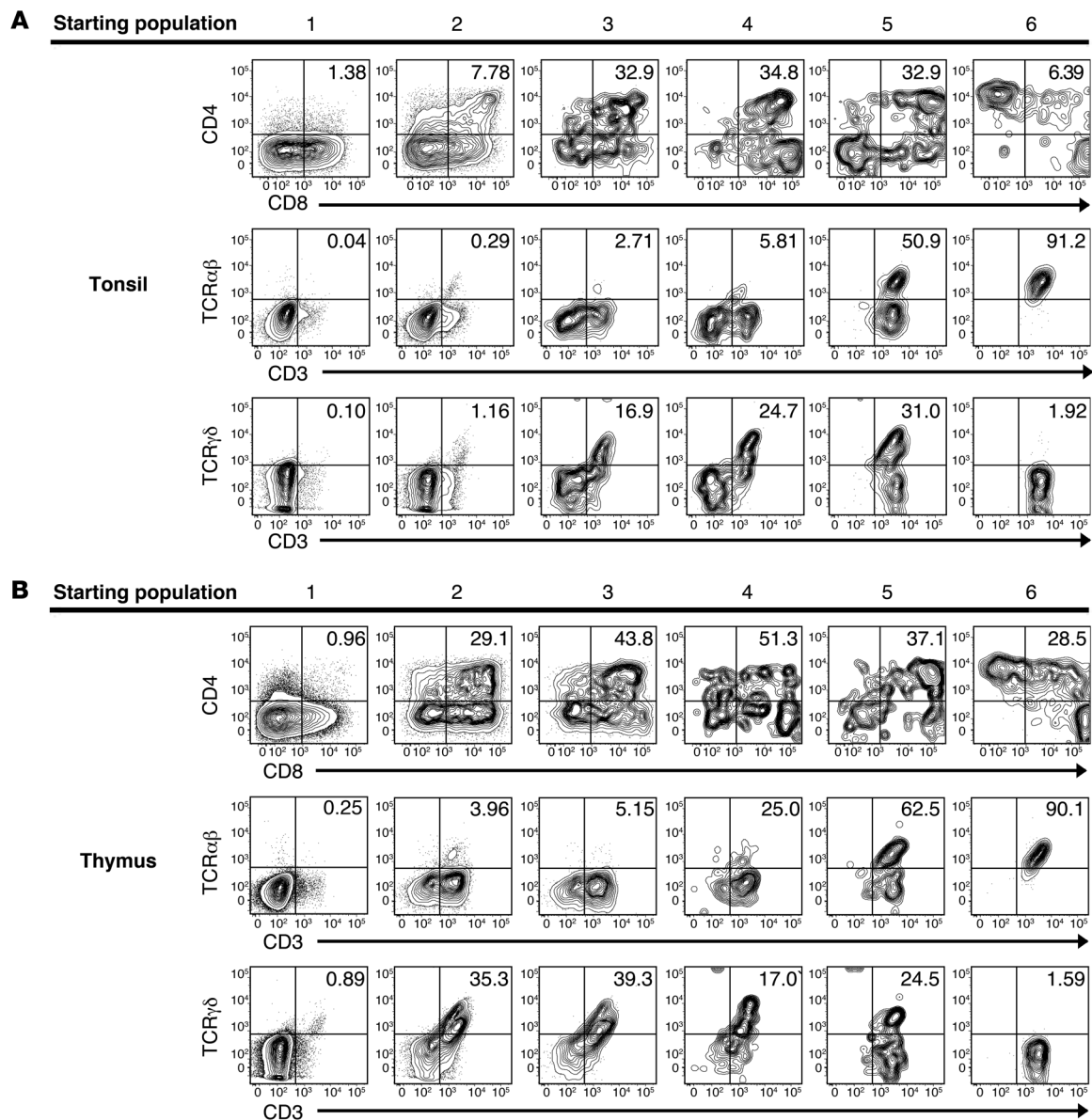
To gain further insight into the kinetics of T cell development from tonsillar precursor cells, we sorted populations 1–6 from the tonsil and cultured them on the OP9-DL1 cell line with exogenous FL and IL-7 for either 7 or 14 days. Whereas a few of the GFP-CD45⁺ progeny were CD3⁺ at days 7 or 14, only a very small percentage of these CD3⁺ cells coexpressed a TCR (Supplemental Figure 4, C and D). This suggests that TCR expression is an event that occurs later in T cell development and confirms that the appearance of mature T cells at 26 days is not simply an artifact of contamination by tonsillar CD3⁺TCR⁺ cells. Furthermore, in 26-day cultures with FL and IL-7 on the OP9 stromal cell line, which lacks notch ligand expression, populations 1–4 from the tonsil failed to generate CD3⁺TCR⁺ T cells. Tonsillar populations 5 and 6, on the other hand, maintained a population of CD3⁺TCR⁺ cells in these conditions (data not shown).

Ex vivo NK cell differentiation potential of putative extrathymic T cell precursors. Human NK cell development has been studied extensively in human tonsils and lymph nodes (12–14, 28–31). To examine whether extrathymic T cell precursors could contribute to tonsillar NK cell development as well, the 6 populations identified in Figure 5A were cultured with FL, c-kit ligand (KL), IL-3, IL-7, and IL-15 on the OP9 stromal cell line. Whereas cultures of populations 5 and 6 from both tissues remained exclusively CD3⁺ (Figure 8A), populations 1–3 from both the tonsil and thymus gave rise to CD56⁺CD3[−] NK cells. Population 4 cells from the tonsil gave rise to CD56⁺CD3[−] cells in 3 out of 3 experiments, whereas thymic population 4 cells gave rise to CD56⁺CD3[−] cells in 1 out of 5 experiments, although no population 4 cells survived in 3 out of these 5 thymic experiments. The resulting CD56⁺ cells derived from tonsil and thymus expressed CD161 and low levels of NKp46 (Figure 8, B and C), along with variable expression of CD5, the latter of which increased from less than 10% in tonsillar populations 1 and 2 to more than 50% in populations 3 and 4.

To further investigate the T and NK cell differentiation capabilities of putative extrathymic T cell precursors in the tonsil, single cells were sorted from tonsillar populations 1–4 into wells containing OP9-DL1 cells and the cytokines FL, IL-7, and IL-15. The individual cells were cultured in this system for 14 days. After culture, each well was harvested and examined for the presence of CD5⁺CD3[−] T cell precursors, CD3⁺ T cells, or CD56⁺CD3[−] NK cells (Supplemental Table 3). Between 16% and 67% of the wells plated sustained CD45⁺ cells at the time of harvest. This assay demonstrates that all 4 populations have the potential to differentiate into CD5⁺CD3[−] T cell precursors and CD56⁺CD3[−] NK cells and that there was an increasing tendency for a single cell to generate CD3⁺ cells as one progressed from population 1 through population 4. Occasionally, single cells from populations 2 and 3 gave rise to wells containing both CD3⁺ T cells and CD56⁺CD3[−] NK cells.

Discussion

We have identified 5 subsets of lymphocytes which we believe to be novel within the human tonsil: (a) CD34⁺CD38^{dim}Lin[−], (b) CD34⁺CD38^{bright}Lin[−], (c) CD34⁺CD1a⁺CD11c[−], (d) CD34⁺CD1a⁺CD3[−]CD11c[−], and (e) CD34⁺CD1a⁺CD3⁺CD11c[−] cells. We have provided phenotypic, gene expression, and functional evidence that these subsets represent intermediates in a program of extrathymic T cell development. Tonsil CD34⁺CD38^{dim}Lin[−] and CD34⁺CD38^{bright}Lin[−] cells share many phenotypic properties of the same cells identified in the thymus and have differentiation potential to develop into T cells and NK cells. Remarkably, the CD34⁺CD1a⁺CD11c[−] cells identified in the tonsil display a phenotype that is virtually identical to that of thymic CD34⁺CD1a⁺ T cell precursors and display consistent T cell developmental capabilities. Similarly, CD34⁺CD1a⁺CD11c[−] cells of the tonsil highly resemble CD34⁺CD1a⁺ cells of the thymus, in that they can be divided into 2 populations based on the expression of CD3,

**Figure 7**

T cell differentiation potential of putative extrathymic T cell precursors in the human tonsil. (**A** and **B**) Cells were sorted from the human (**A**) tonsil or (**B**) thymus, as indicated in Figure 5A. All 6 populations were cultured independently on OP9-DL1 cells with FL and IL-7 for 26 days. After harvest, cells were analyzed for the expression of CD3, CD4, CD8, TCR $\alpha\beta$, and TCR $\gamma\delta$. Data are gated on GFP-CD45⁺ events to exclude OP9-DL1 stromal cells from analysis. Data shown are representative of experiments performed with 7 individual tonsils or 4 thymus donors. Numbers represent the mean percentages of GFP-CD45⁺ cells that stained positive for both antigens indicated on the dot plot.

they coexpress CD4 and CD8, and they retain T cell developmental potential. Furthermore, in addition to identifying CD1a⁺TdT⁺ extrathymic T cell precursors in human tonsil, we have localized the bulk of this population near fibrous scaffolding, suggesting that it resides in a distinct microenvironment. Finally, we have demonstrated that CD34⁺/CD1a⁺CD11c⁻ cells from the thymus and tonsil are not committed to the T cell lineage but retain *ex vivo* differentiation potential to also develop into NK cells.

While previous reports have identified a role for murine intestinal lymphoid tissue in T cell development (32–35), the few studies investigating human extrathymic T cell differentiation

are difficult to interpret. It is known that uncommitted hematopoietic progenitors with T cell differentiation capabilities reside within the bone marrow (6, 36, 37) and cord blood (24, 38). Furthermore, tonsillar TdT⁺ cells of unclear etiology and pre-T α ⁺ cells within bone marrow and neonatal intestine have been identified (4, 7–9). However, a complete description of T cell differentiation in an extrathymic human tissue has not been provided. Our work suggests that T cell development occurs in the human tonsil, and future studies should address what proportion of the total CD3⁺ tonsillar pool is derived within the tonsil compared with that from the thymus.

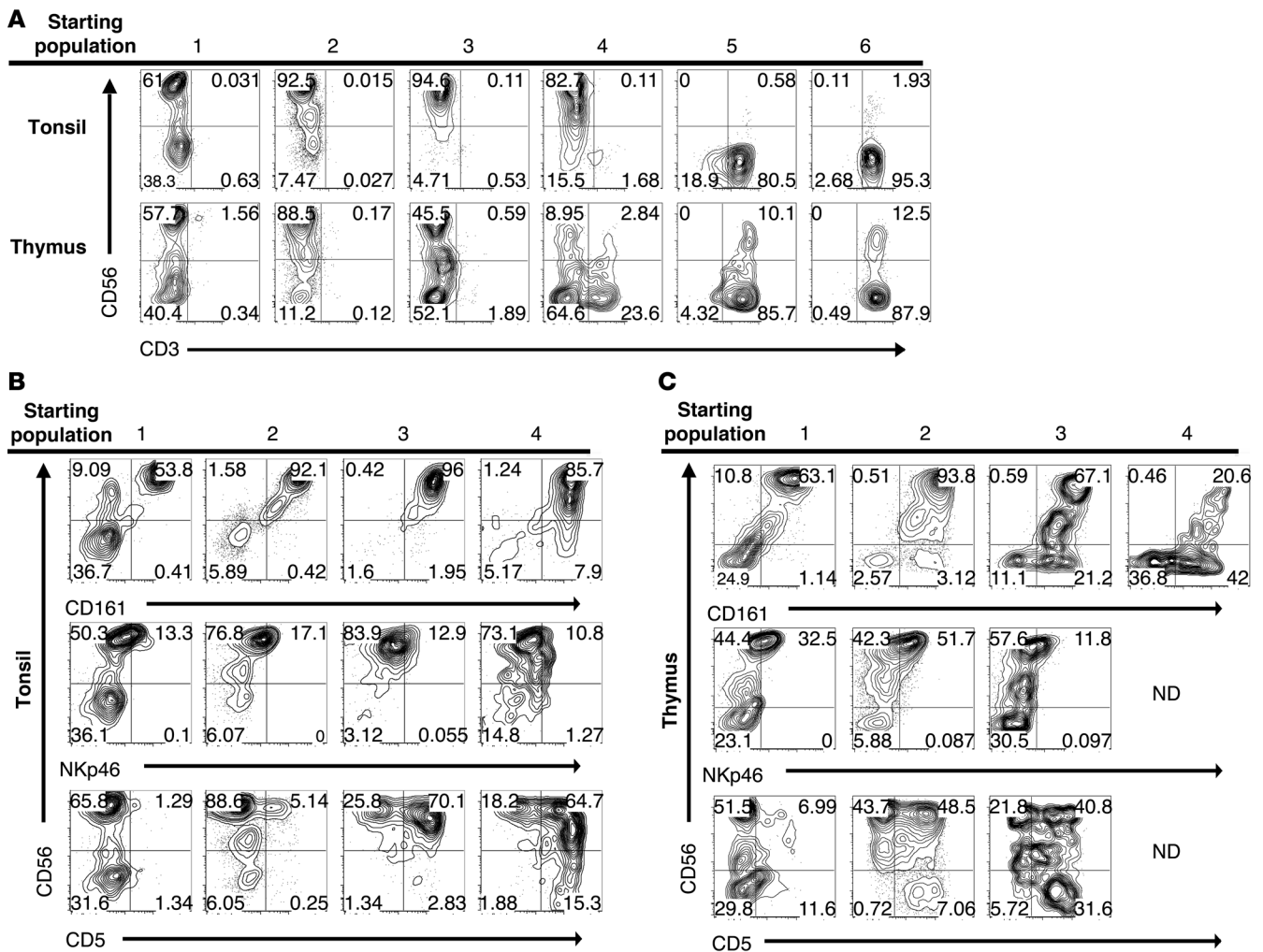


Figure 8

NK cell differentiation potential of putative extrathymic T cell precursors in the human tonsil. (A) Tonsil and thymus cells were sorted as described in Figure 5A. Sorted cells were cultured on OP9-GFP cells and cultured with FL, KL, IL-3, IL-7, and IL-15 for 18 to 19 days. Harvested cells were gated on GFP-CD45⁺ events and analyzed for expression of CD3 and CD56. (B and C) GFP-CD45⁺ progeny from populations 1–4 were gated on CD3⁺ events and analyzed for expression of CD56, CD161, NKp46, and CD5. CD3, CD56, CD161, and CD5 data are representative of independent experiments performed with 3 tonsil or 5 thymus donors. NKp46 data are representative of 2 experiments performed with tonsil donors or 3 experiments performed with thymic donors. No data (ND) are available for NKp46 or CD5 expression on progeny of thymic population 4 cells, due to the low numbers of harvested cells. All dot plots and gate frequencies are from a representative experiment. Numbers within each quadrant represent the percentage of events falling within that gate for the representative experiment shown.

It is intriguing to think of a scenario in which both the thymus and tonsil are seeded by the same bone marrow-derived CD34⁺CD38^{dim}CD45RA⁺ progenitor that gives rise to NK cells and T cells in each tissue. However, of the 5 populations identified, we found the most substantial differences when comparing the CD34⁺CD38^{dim}Lin⁻ and CD34⁺CD38^{bright}Lin⁻ populations from the tonsil and thymus. Specifically, thymic CD34⁺CD38^{dim}Lin⁻ cells largely expressed the T cell-associated antigens CD2, CD7, and CD5, whereas tonsillar CD34⁺CD38^{dim}Lin⁻ cells did not. Thus, it is possible that these populations represent distinct progenitors that independently seed the thymus and tonsil. Alternatively, it is possible that the thymus and tonsil are seeded by similar CD34⁺CD38^{dim} progenitors but that the phenotype of these entering cells is rapidly altered by the thymic or tonsillar microenviron-

ment, as is seen in the mouse (39, 40). As demonstrated in Figure 2B, while a majority of thymic CD34⁺CD38^{dim}Lin⁻ cells are CD7⁺, a few lack CD7 (4%–17%). Indeed, recent reports suggest that a CD34⁺CD45RA⁺CD10⁺CD7⁻ cell seeds the thymus after birth, and a CD34⁺CD45RA⁺CD10⁺CD7⁺ precursor has yet to be identified in postnatal bone marrow or peripheral blood (5, 6). Thus, it is possible that both tissues are seeded by the same circulating CD34⁺CD38^{dim}CD10⁺CD7⁻ hematopoietic progenitor and that this cell is capable of giving rise to downstream T cell developmental intermediates in each tissue.

Regardless of whether the 2 tissues are seeded by the same progenitor, it is intriguing that TdT⁺ cells within the tonsil aggregate in foci, surrounding fibrous scaffolding. Were the trafficking of T cell precursors to the tonsil a random event associated with a



“leaky” thymus, one would assume that the cells would perfuse the tonsil in a scattered arrangement in the interfollicular zone near mature T cells. However, the consistent aggregation of T cell precursors near the fibrous regions of the tonsil suggests that this space serves a unique anatomical function in supporting extrathymic T cell precursor homing and/or maturation. This hypothesis is further supported by the identification of Notch ligands, which are required for complete T cell differentiation, within the same regions of the human tonsil.

The capsule of the tonsil is highly invaginated into crypts, which create a large surface area important for foreign antigen exposure. Thus, the localization of T cell precursors to this fibrous region may play a role in mucosal immunity. Alternatively, the thymus is segregated to support different phases of thymocyte development in anatomically distinct microenvironments (41–44), and our finding of tonsillar CD1a⁺TdT⁺ cells near the fibrous scaffolding, but not in the T cell-rich interfollicular zones, suggests that perhaps extrathymic T cell development in the tonsil is also anatomically divided to provide the proper microenvironment for each phase of maturation. In the thymus, the least mature progenitors enter the thymus at the junction between the outer cortex and the inner medulla. These cells migrate outward until they reach the subcapsular zone, which marks the division between the fibrous thymic capsule and the cortex, and then they travel inward toward the medulla to complete differentiation (43, 44). Future studies should investigate whether the microenvironment of the tonsillar fibrous scaffold is analogous to the subcapsular zone of the thymus, in terms of supporting T cell development, or whether it provides a unique signal directing the homing of extrathymic precursors to this region of the tonsil.

This report also provides evidence for the expression of key genes involved in TCR rearrangement within these tonsillar populations. Specifically, we found that populations 2–5 express mRNA for *RAG1*, which is necessary for initiation of TCR rearrangement (45), and for *PTCRA*, which is a molecular indicator that successful TCR β rearrangement has occurred (3). The expression of these 2 genes, along with TdT, suggests that somatic rearrangement of the TCR may occur in the human tonsil.

In 1998, Spits et al. reported that CD34⁺CD1a⁺ thymocytes generate only small numbers of NK cells (46). Since then it has generally been held that loss of CD34 and acquisition of CD1a are associated with T cell commitment (1, 46, 47). Our findings confirm and extend their work in the thymus, in that we found that CD34⁺CD38^{dim}, CD34⁺CD38^{bright}, and CD34⁺CD1a⁺ thymocytes and tonsil cells readily generate NK cells and that tonsillar CD34⁺CD1a⁺CD11c⁻ cells also retain NK cell potential. Furthermore, some CD56⁺CD3⁻ NK cells generated in our ex vivo cultures coexpressed CD5, and there was a relative increase in CD56⁺CD5⁺CD3⁻ progeny generated as one progressed from CD34⁺CD38^{dim}Lin⁻ cells to CD34⁺CD1a⁺CD11c⁻ cells. This suggests that the generated NK cells are not merely a result of starting populations that were impure at sorting but that a unique CD5⁺ progenitor may be giving rise to the CD56⁺CD3⁻ cells in cultures initiated with the CD34⁺CD1a⁺CD11c⁻ and CD34⁺CD1a⁺CD11c⁻ subsets. Our finding that all 4 populations from the tonsil generate CD56⁺CD3⁻ NK cells at the single cell level as well substantiated these results. It has become evident that the human tonsil is a significant site of NK cell development (12–14, 28–31). However, whether or not differentiating CD34⁺/CD1a⁺CD11c⁻ cells that also express TdT contribute to the physiological NK cell differentiation within this tissue remains unclear.

That the tonsil does support T cell development could have significant clinical and scientific implications. First, the role of tonsillar T cell development in human disease is unknown. That T cells can develop extrathymically in this tissue raises the possibility that the tonsil may actually support the generation of autoreactive T lymphocytes or even contribute to malignant transformation. Furthermore, the role of extrathymic T cell development in the tonsil during inflammation should be of future interest. In this study, we have obtained pediatric tonsils from patients undergoing routine tonsillectomy. While it can be presumed that the patients are afebrile at the time of surgery, the possibility that repeat or sustained inflammation contributes to the presence of extrathymic T cell development should be further explored.

Alternatively, the identification of an extrathymic reservoir of human T cell precursors in normal individuals could suggest clinical opportunities to enhance T cell regeneration in situations of thymic defect. For example, T cell deficiency is characteristic in patients with poor thymic function due to complete DiGeorge syndrome, chemoablation or radioablation, or surgical thymectomy (48–52). Indeed, the efficient regeneration of viable thymic tissue after total body irradiation for bone marrow transplant continues to be a significant clinical challenge (49). In the future it may be possible to harvest tonsillar T cell precursors or healthy tonsil stroma via tonsillectomy in order to augment T cell regeneration after transplant in the same individual. While these possibilities remain speculative at the moment, the identification of a program of extrathymic T cell development in the human tonsil is a unique finding with potential implications for human health and disease.

Methods

Human samples. Human tonsils and thymuses were obtained from Nationwide Children’s Hospital (Columbus, Ohio, USA). Tonsils were obtained from pediatric patients undergoing tonsillectomies, whereas thymuses were obtained from children undergoing thoracic surgery. Paraffin-embedded adult tonsils and reactive lymph nodes were obtained from the Biospecimen Shared Resource at The Ohio State University, and adult peripheral blood was obtained from the American Red Cross (Columbus, Ohio, USA). Mononuclear cells were obtained from tonsils and thymuses after manual disaggregation and Ficoll-Paque centrifugation. For experiments wherein tonsillar cells were enriched for CD34-expressing cells only, total cells were depleted of CD19⁺ and CD3⁺ cells by magnetic depletion and were enriched for CD34⁺ cells using an indirect CD34 Selection Kit (Miltenyi Biotec). For experiments in which tonsillar cells were enriched for CD1a⁺ cells only, total cells were depleted of CD19⁺ cells and then magnetically enriched for CD1a-expressing cells (CD1a magnetic beads, Miltenyi Biotec). To simultaneously enrich for CD34- and CD1a-expressing tonsillar cells, CD19-depleted mononuclear cells were subjected to positive selection using the indirect CD34 Selection Kit (Miltenyi Biotec) and CD1a magnetic beads (Miltenyi Biotec) concurrently. To isolate thymic T cell progenitors, total thymic cells were subjected to positive CD34 selection using the indirect CD34 Isolation Kit (Miltenyi Biotec).

Flow cytometry and cell sorting. Antibodies used for flow cytometry were purchased from BD Biosciences, except those for CD56, CD1a, CD127, Nkp46, CD16, and CD161-PE (all from Beckman Coulter) and those for CD161-APC, CD161-FITC, CD11c-FITC, and BDCA-2 (all from Miltenyi Biotec). Intracellular staining for TdT and icCD3 ϵ was performed after surface staining using the BD Cytofix/Cytoperm Plus Kit (BD). Flow cytometry was performed on a FACSVantage (BD) or LSR II (BD), and data were



analyzed using FlowJo software (Treestar Inc). Cell sorting was performed on a FACSAria (BD). The complete gating strategy and a representative purity assessment are shown in Supplemental Figure 2. Purities of sorted populations were routinely more than 95% pure.

Immunohistochemistry. Immunohistochemistry was performed as described previously (12, 31, 53) using the UltraView Universal system (Ventana Medical). Anti-TdT (1:100), α -CD1a (1:10), α -CD34 (1:100), α -CD11c (1:100), α -DL1 (1:300), and α -DL4 (1:300) were purchased from Abcam, and α -Rag1 (1:10) was purchased from Santa Cruz Biotechnology Inc. For immunohistochemistry involving α -DL1 and α -DL4, antigen retrieval was performed for 30 minutes prior to staining. For immunohistochemistry involving α -Rag1, antigen retrieval was performed for 30 minutes, along with digestion in Ventana Protease 1 (1:100) for 4 minutes. Images were obtained using a DP 12 camera, a BX50 microscope, and UPLANF1 objectives (Olympus). DAB- and fast-red-stained sections were digitally converted to fluorescent green and red, respectively, using the Nuance FX system (Cambridge Research & Instrumentation).

Differentiation assays. Cultures were performed in α -MEM plus L-glutamine (Gibco) supplemented with 20% fetal bovine serum and penicillin G plus streptomycin (100 μ g/ml; Gibco), with 500–5,000 cells per well. Medium was supplemented with the following cytokines as indicated: human recombinant FL (100 ng/ml; Miltenyi Biotec), KL (100 ng/ml; Amgen), IL-3 and IL-7 (10 ng/ml; Miltenyi Biotec), and IL-15 (1 nM; Miltenyi Biotec). T cell differentiation assays were performed as described previously (13, 24) on OP9-DL1 cells (a gift from J.C. Zúñiga-Pflücker, University of Toronto, Toronto, Ontario, Canada) with IL-7 and FL. NK cell differentiation assays were performed on OP9 cells (a gift from J.C. Zúñiga-Pflücker) with FL, KL, IL-7, IL-3, and IL-15. For single cell cultures, individual cells from populations 1–4 of the human tonsil were sorted directly into 59 to 60 wells containing OP9-DL1 cells, FL, IL-7, and IL-15 for 14 days. Medium was changed every 4 to 5 days.

Real-time RT-PCR. Tonsillar populations 1–6 were sorted from 8 tonsil donors, as shown in Figure 5A and Supplemental Figure 2. From these, we generated 2 pools for each population, with each pool containing cells from 4 of the 8 donors and approximately 3,500 cells. 5,000 cells from populations 1–6 were sorted from 2 thymus donors for comparison. RNA was extracted using the RNeasy Micro Kit (Qiagen), and real-time RT-PCR was

performed as previously described (30), using ABI TaqMan primer/probe sets for *PTCRA* (probe Hs00300125_m1), *RAG1* (probe Hs00172121_m1), *BCL2L1* (probe Hs00236329_m1), and *THPOK* (probe_Hs01035470_m1). Expression levels were normalized to *18S* expression and were analyzed using the comparative $\Delta\Delta$ CT method (54).

Statistics. To investigate whether age or gender correlates with the percentage of CD34⁺CD1a⁺CD11c⁻ cells, a Spearman correlation and a 2-sample, 2-tailed *t* test were used. In reporting the percentages of CD3⁺ T cells generated in T cell differentiation assays, the mean percentage \pm SEM of GFP⁻CD45⁺ cells was listed. For comparison of CD3⁺ cells generated from populations in the human thymus and those from the tonsil, the linear mixed effects models were used to take account of the correlation among observations from the same donor. After Holm's adjustment for multiple comparisons, *P* values of equal to or less than 0.005 were considered significant.

Study approval. Tonsillar, thymic, and peripheral blood samples were obtained with approval of these experiments from The Ohio State University Comprehensive Cancer Center Institutional Review Board. As all human samples were obtained as deidentified tissue, informed consent was not required.

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