The autoimmune regulator (Aire) serves an essential function for T cell tolerance by promoting the "promiscuous" expression of tissue antigens in thymic epithelial cells. Aire is also detected in rare cells in peripheral lymphoid organs, but the identity of these cells is poorly understood. Here, we report that Aire protein–expressing cells in lymph nodes exhibit typical group 3 innate lymphoid cell (ILC3) characteristics such as lymphoid morphology, absence of "classical" hematopoietic lineage markers, and dependence on RORγt. Aire+ cells are more frequent among lineage-negative RORγt+ cells of peripheral lymph nodes as compared with mucosa-draining lymph nodes, display a unique Aire-dependent transcriptional signature, express high surface levels of MHCII and costimulatory molecules, and efficiently present an endogenously expressed model antigen to CD4+ T cells. These findings define a novel type of ILC3-like cells with potent APC features, suggesting that these cells serve a function in the control of T cell responses.

Introduction

The autoimmune regulator (Aire)’s crucial function in the promotion of promiscuous gene expression in medullary thymic epithelial cells (mTECs) is well established. mTECs express thousands of tissue-restricted antigens (TRAs) and present these on MHCII and II (Kryewski and Klein, 2006; Mathis and Benoist, 2007; Peterson et al., 2008; Klein et al., 2014). Aire deficiency strongly diminishes TRA expression in mTECs, offering an explanation how Aire mutations cause the human autoimmune polyendocrine syndrome type 1 (Husebye et al., 2018) and sim-
Here, we aimed to clarify the identity of Aire-expressing cells in lymph nodes. We identified three phenotypically distinct subsets of hematopoietic cells that expressed endogenous Aire mRNA, including the previously described EpCAM−CD11c+ eTACs. However, Aire protein was exclusively found in an EpCAM−CD11c− innate lymphoid cell (ILC) 3-like cell type with potent APC features.

Results and discussion

We confirmed in two independent mouse strains that Aire-reporter expression in LNs was confined to MHCII+ cells (Gardner et al., 2013). Surprisingly, Aire-reporter+ cells not only contained cells with the reported EpCAM−CD11c+ “eTAC” phenotype, but also similar proportions of EpCAM−CD11c− and EpCAM−CD11c− cells (Figs. 1 A and S1 A). Endogenous Aire mRNA was highest in EpCAM−CD11c− cells (Figs. 1 B and S1 B). Aire-protein was detectable by intracellular staining (ICS) in 10–20% of Aire-reporter−EpCAM−CD11c− cells, but not in the other two subsets (Figs. 1 C and S1 C). Aire was localized in nuclear dots, akin to its subcellular distribution in mTECs (Fig. 1 D).

Independent of reporter systems, Aire+ LN cells were detectable by ICS, and these cells were uniformly CD45−EpCAM−CD11c− (Fig. 1 E). Aire−ICS+ cells were found at similar frequencies in non-adaptive ILCs, such as as LTi-like cells (Fig. S1 D). Their abundance was very low in newborns and increased with age (Jones et al., 2018; Fig. 1 F and 2 K). Thus, Aire+ LN cells resembled LTi-like ILCs, yet displayed distinctive features, and from here on will be referred to as Aire+ ILC3-like cells.

Aire, besides its role in TRA expression, has been suggested to control the development and/or survival of mTECs (Gray et al., 2007; Hikosaka et al., 2008; Wang et al., 2012). To address whether a lack of Aire in cells that otherwise express it would perturb their differentiation, we generated mixed [Aire−/−; Aire-reporter: Aire+−/−, Aire-reporter1] → WT BM-chimeras. Aire-reporter+ cells that had emerged from either Aire−/− or Aire+−/− precursors were present at equal frequencies. Thus, Aire is unlikely to be a developmental or homeostatic regulator of Aire+ ILC3-like cells (Fig. 3 A).

Aire expression in mTECs is orchestrated by NF-κB signals that emanate from receptors of the TNF superfamily (Akiyama et al., 2012) and converge on an essential enhancer-element (CNSI) in the Aire gene (Haljasorg et al., 2015; Lam Fl et al., 2015). Aire+ LN cells were absent in Aire−CNSI−/− mice, indicating that the gene-proximal requirements for Aire expression in Aire+ ILC3-like cells resemble those in mTECs (Fig. 3 B). MHCII−dependent cognate interactions of mTECs and thymic B cells with developing CD4 T cells provide a crucial platform for Aire-inducing Rank or CD40 signals (Rossi et al., 2007; Akiyama et al., 2008; Hikosaka et al., 2008; Irla et al., 2008; Roberts et al., 2012; Yamano et al., 2015). We asked whether a similar scenario applies to Aire+ LN cells. In mixed [MHCII−/−; MHCII−/−] → WT BM-chimeras, MHCII-deficient precursor cells efficiently gave rise to Aire+ LN cells (Fig. 3 C). Moreover, Aire+ LN cells were similarly abundant in Rag2−/− mice and WT controls (Fig. 3 D). Thus, in contrast to what has been established for mTECs and thymic B cells, Aire+ LN cells are independent of cross-talk with T cells, indicating that an “innate program” governs their cellular identity.

Given the critical role of Aire’s NF-κB response element in Aire+ LN cells, it was likely that extrinsic TNF receptor family signals of “nonadaptive” origin were crucial for their differentiation. Aire+ LN cells emerged with equal efficacy from either CD40−/− or CD40+/+ precursors (Fig. 3 E). By contrast, Rank
Figure 1. **Phenotype of Aire-expressing cells in LNs.**

(A) Expression of GFP and MHCII in LN cells from Adig Aire-reporter mice and WT controls and staining for CD11c and EpCAM on gated Aire-GFP+MHCII+ cells (representative of \( n \geq 4 \) each). (B) Aire mRNA in medullary and cortical thymic epithelial cells (mTECs and cTECs, respectively) and in Aire-GFP+ LN cells sorted according to expression of CD11c and EpCAM. Data are mean values ± SEM of triplicates. AU, arbitrary units. (C) ICS for Aire protein in subsets of Aire-GFP+ LN cells. The average frequency ± SEM of Aire-ICS+ cells is indicated (\( n = 4 \)). (D) Nuclear localization of Aire protein and surface marker expression in Aire-expressing LN cells or mTECs visualized by imaging flow cytometry. (E) ICS for Aire protein and surface expression of MHCII in total LN cells from WT and Aire-/- mice. Histograms on the right show CD11c, EpCAM, and CD45 on gated Aire+MHCII+ cells (representative of \( n = 5 \)). (F) Number of Aire+ LN cells per 10^6 total LN cells in mice of the indicated age (\( n \geq 3 \) each; 0 wk = 4 d old). (G) Aire and MHCII expression in total LN cells from Aire-/- → Aire-/-, Aire-/- → WT or WT → Aire-/- BM chimeras (representative of \( n \geq 4 \) each).
Figure 2. Aire+ LN cells display ILC3 characteristics. (A) Expression of hematopoietic lineage markers on Aire-GFP+MHCII+CD11c−EpCAM−LN cells from Adig mice (representative of n ≥ 4). (B) RNA expression of hematopoietic lineage-specific signature genes in Aire+ LN cells. (C) RNA expression of ILC signature genes in enriched Aire+ LN cells. (D) ICS for RORγt and Gata3 in MHCII+Aire+ LN cells (blue dots) back-gated on total Lin− (CD3, CD19, B220, Gr-1, CD11c, and CD11b) LN cells (gray dots; representative of n = 3). (E) Morphology of representative Aire-GFP+MHCII+CD11c−EpCAM− LN cells, ‘canonical’ LN ILC3s (Lin−MHCII−RORγt+IL7Rα+), and eTACs (Aire-GFP+MHCII+EpCAM+CD11c+). (F) Contribution of WT and Rag2−/− precursor cells to Aire-ICS+ LN cells or T cells in 1:1 mixed [Rag2−/−:WT] → WT BM-chimeras (n = 6). (G) Contribution of Rorc+ and Rorc− precursor cells to Aire-ICS+ cells in LN or spleen or to DCs in 1:1 mixed [Rorc+/−:Rorc−/−] → WT BM-chimeras (n = 7). (H) Gating strategy for Lin− (Lin1: CD3, CD19, B220, Gr-1; Lin 2: CD11c and CD11b) RORγt+ LN ILC3s and expression of Aire by ICS. (I) Intracellular staining for Id2 protein and surface expression of c-kit, CCR6, Nkp46, IL7Rα, CD90, and CD4 in Aire+ (filled blue histogram) and Aire− (open orange histogram) Lin−RORγt+ cells (representative of n = 3). FMO, fluorescence minus one. (K) Number of ‘canonical’ LN ILC3s (orange) and Aire+ LN cells (blue) in 4-d- and 6-wk-old animals (n = 4). Data are mean ± SEM.
deficiency resulted in a severe cell-intrinsic defect of hematopoietic precursors to contribute to Aire+ LN cells in mixed chimeras (Fig. 3 F). Intriguingly, concomitant to being virtually absent from Aire+ ILC3-like cells, Rank-deficient cells were strongly over-represented among canonical IL7Rα+ ILC3s. A possible explanation for this was that LN ILC3s under steady-state conditions differentiate into Aire+ ILC3-like cells upon Rank stimulation, yet accumulate when their capacity to signal via Rank is genetically ablated. In support of such a precursor/progeny relationship, in vitro culture of canonical LN ILC3s with Rank-ligand–expressing stromal cells induced Aire expression in a fraction of cells (Fig. 3 G).

RNA sequencing (RNA-seq) of Aire-reporter+ LN cells from WT and Aire-deficient mice revealed 707 differentially expressed transcripts (Fig. 4 A). Of these, 334 were Aire-induced, and 373 were repressed (Tables S1 and S2). Quantitative PCR analysis of selected Aire-induced or Aire-repressed genes confirmed their Aire-regulated expression (Fig. 4 B). Among Aire-induced transcripts, only 60 were classified as TRAs (Sansom et al., 2014; Fig. 4 A). Thus, in contrast to what is known for...
Figure 4. Aire orchestrates a distinct genetic program in Aire+ ILC3-like cells. (A) RNA-seq data from Lin−MHCIIhiCD80+ cells from Aire+/+ and Aire−/− mice. Genes encoding for TRAs are colored in red. Fold-change cutoff: 1.5; P value: 0.05 (indicated by a dashed line). (B) Quantitative PCR analysis of mRNA expression of selected differentially expressed genes. Data are mean ± SD of three independent experiments. ns, not significant. (C) Comparison of Aire-dependent gene expression in Aire+ LN cells and mTECs. Genes that are up-regulated (depicted in red; 1,732 genes) or down-regulated (depicted in blue; 423 genes) by Aire in mTECs by at least 1.5-fold were projected onto the volcano plot shown in A. Student's t test was used to calculate P values.
mTECs (Anderson et al., 2002; Kyewski and Klein, 2006), the Aire-induced transcriptome of Aire+ LN cells is not biased toward TRAs. Interestingly, Aire-regulated genes in Aire+ LN cells and mTECs were largely nonoverlapping, that is, only 90 of 707 Aire-regulated transcripts in Aire+ LN cells were also affected by Aire deficiency in mTECs. Moreover, many of these genes displayed opposing effects in mTECs or Aire+ LN cells (Fig. 4 C). Thus, Aire controls a distinct transcriptional program in Aire+ ILC3-like cells.

Aire+ LN cells had high surface levels of MHCII and strongly expressed mRNAs encoding for costimulatory or coinhibitory molecules (Fig. 5 A). To relate this to canonical LN ILC3s, we directly compared the surface phenotype of Aire+IL7Rα- or Aire+IL7Rα- Lin+CD45+RORγt+ LN cells (Figs. 5 B and S3 A). Aire+ ILC3-like pLN cells had homogeneously high surface levels of MHCII, CD60, CD86, CD40, and Icos-L and detectable amounts of PD-L1. By contrast, with the exception of CD86, these molecules were substantially lower or absent on Aire+IL7Rα+ ILC3s (Fig. 5 C). We confirmed and extended these data through gating on MHCIIhiIL7Rα- or MHCIIα/αIL7Rα- cells among Lin+CD45+RORγt+ cells, which clearly separated Aire+ ILC3-like cells and Aire- canonical ILC3s (Fig. S3, A and B). Consistent with our previous observations, MHCIIhiIL7Rα- Aire+ ILC3-like cells were far more abundant in pLNs as compared with mesenteric LNs (Fig. S3, A–C), were slightly lower for CD45 (Fig. S3 D), and had much higher surface levels of APC-associated molecules as compared with MHCIIα/αIL7Rα- cells (Fig. S3 E). Moreover, MHCIIhiIL7Rα- Aire+ ILC3-like cells were negative for CD4, CD90, and CD25, whereas canonical ILC3s were heterogeneous for CD4 and CD90 and homogenously positive for CD25 (Fig. S3 F).

Given the potent APC features of Aire+ ILC3-like cells, we asked whether a model antigen expressed by these cells was visible to CD4 T cells. Influenza hemagglutinin (HA)-specific CD4 T cells and polyclonal “reference” T cells were adoptively transferred into Aire-HCO mice, which express HA from a bacterial artificial chromosome-transgenic Aire gene (Hinterberger et al., 2010). 14 d after transfer, HA-specific CD4 T cells were strongly diminished in Aire-HCO recipients but not in WT controls (Fig. 5 D), indicating that peripheral antigen encounter resulted in the deletion of these cells. Similar results were obtained when Aire-HCO → WT BM-chimeras were used as recipients, indicating that deletion of TCR-HA cells resulted from HA expression in the hematopoietic compartment (Fig. 5 E). To directly assess presentation of endogenously expressed HA, Aire+ ILC3-like cells from Aire-HCO mice were cultured with HA-specific CD4 T cell hybridoma cells carrying a GFP IL-2 reporter (Aschenbrenner et al., 2007). For comparison, we performed these assays also with mTECs and EpCAM+CD11c+ eTACs. Aire+ ILC3-like cells presented HA with similar efficacy as mTECs. By contrast, no direct presentation of HA was measurable with eTACs, despite similar expression of the Aire-reporter (Figs. 5 F and S1 A). When loaded with exogenous HA-peptide, all three cell types elicited similar responses, excluding that differential expression of MHCII accounted for these differences (Fig. 5 F).

In sum, our data suggest that Aire protein-expressing cells in mouse pLNs may represent a hitherto unknown ILC3 subset. In distinction from the majority of Lin+RORγt+ LN ILC3s, they do not express the IL7Rα, which, besides the absence of Aire+ ILC3s from the intestine and their low frequency in mesenteric LNs, may explain why Aire expression has not been noticed in transcriptomic analyses of ILC subsets (Robinette et al., 2015; Gury-BenAri et al., 2016). It will be interesting to clarify the potential precursor progeny relationship between canonical ILC3s and Aire+ LN cells and how the Rank pathway is involved. It was recently shown that Rank signaling negatively regulates the abundance and effector functions of intestinal ILC3s (Bando et al., 2018). Together with our findings, this suggests that both in pLNs as well as in the intestine, ILC3s receive tonic Rank stimulation, yet may differ with regards to the ensuing biological response.

Aire+ ILC3-like cells display potent APC features, consistent with accruing evidence that ILC3s can orchestrate CD4 T cell responses. Selective ablation of MHCII molecules on RORγt+ cells resulted in impaired tolerance to commensals and intestinal inflammation (Hepworth et al., 2013, 2015). Other reports linked MHCII deficiency in RORγt+ cells to impaired T and B cell responses to “foreign” antigens (von Burg et al., 2014). How this relates to Aire+ ILC3-like cells and under which circumstances antigen recognition on these cells may result in tolerance versus immunity remains to be seen. Along these lines, it will be interesting to see whether Aire expression in Aire+ ILC3-like cells is of true physiological significance, as these cells do not recapitulate bona fide “promiscuous” TRA expression as it is seen in mTECs. It remains possible that Aire influences features of Aire+ LN cells that are unrelated to TRA expression.

Materials and methods

Mice

Adig (Gardner et al., 2008), Aire-HCO (Hinterberger et al., 2010), Aire−/− (Ramsey et al., 2002), Aire-CNS1−/− (Haljasorg et al., 2015), TCR-HA (Kirberg et al., 1994), Cd40−/− (Kawabe et al., 1994), Rag2−/− (Hao and Rajewsky, 2001), Rorc−/− (Eberl et al., 2004) Rocr-EGFP (Lochner et al., 2008), and Tfnsr11a−/− (Li et al., 2000) mice have been described previously. Aire−/− and Adig mice were backcrossed onto a BALB/c background for ≥10 generations. Animal studies were approved by local authorities (Regierung von Oberbayern Az. 02-17-193 and Ethical Committee of the Czech Academy of Sciences).

Preparation of LN single-cell suspension

Unless stated otherwise, experiments were performed with pooled pLNs (axillary, brachial, inguinal, and cervical). LNs were pierced with a needle and enzymatically digested with 0.1 mg/ml Dispase (Gibco) in RPMI. After incubation at 37°C for 10 min, the supernatant was adjusted to 3% FCS and 2 mM EDTA and gently homogenized by gently pipetting up and down. To stop the digestion, the supernatant was replenished, and cell suspensions were homogenously positive for CD25 (Fig. S3 F).

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Flow cytometry

Single-cell suspensions of digested LNs were surface stained according to standard procedures. For intracellular Aire or RORγt staining, cells were fixed and permeabilized using reagents from the Foxp3 staining kit (eBioscience) and stained with anti-Aire mAb 5H12 (eBioscience) conjugated to Alexa Fluor 660, Alexa Fluor 488, or FITC- or PE-conjugated RORγt mAb B2D (eBioscience). Cells were analyzed or sorted using a BD FACSCANTOII or LSRII flow cytometer or BD FACSAriaFusion or BD Influx cell sorters.

Figure 5. Aire+ ILC3-like cells display potent APC features and directly present endogenous antigen on MHCII. (A) RNA expression of genes involved in antigen presentation in Aire+ ILC3s. (B) Frequency of Aire+IL7Rα- and Aire-IL7Rα+ cells among Lin- RORγt-ICS+CD45+ cells in pLNs (Lin1: CD3, CD19, B220, Gr-1; Lin 2: CD11c and CD11b). (C) Surface expression of MHCII and costimulatory molecules on Aire+IL7Rα- (filled blue) or Aire-IL7Rα+ (open black) Lin- RORγt-ICS+CD45+ pLN cells (representative of n ≥ 3). (D) Peripheral deletion of HA-specific CD4 T cells upon adoptive transfer into Aire-HCO mice (n = 6 each). (E) Peripheral deletion of HA-specific CD4 T cells upon adoptive transfer into Aire-HCO→WT BM chimeric mice (n = 6 each). (F) GFP expression in HA-specific CD4+ NFAT-GFP-reporter hybridoma cells after 16 h co-culture with mTECs, Aire+ ILC3-like cells (Aire-hCD2+MHCII+CD11c+EpCAM+), or eTACS (Aire-hCD2+MHCII+CD11c+EpCAM+) from Aire–HCO mice without (upper row) or with (lower row) exogenous HA-peptide. Filled gray histograms are from A5 cells that were cultured alone (representative of three experimental replicates). Data are mean ± SEM. Student’s t test was used to calculate P values.
Imaging flow cytometry
LN cells were prepared by enzymatic digestion. Thymic stromal cells were prepared by enzymatic digestion and density fractionation. Cells were stained for surface markers, fixed and permeabilized, and stained with Alexa Fluor 488-conjugated anti-Aire mAb 5HI2 (eBioscience). DAPI was added immediately before analysis. Images were acquired using the ImageStream imaging flow cytometer (Amnis), and data were analyzed with the Ideas 6.0 software.

In vitro stimulation of ILC3s
LN cells from Aire-HCO mice were prepared by enzymatic digestion. B cells and T cells were depleted with biotin-conjugated anti-B220, anti-TCRβ, and anti-biotin MicroBeads (Miltenyi Biotech). Lineage-negative (CD3, CD11c, and CD19), hiCD2-negative, IL7Rα+ MHCII+ -positive cells were sorted. Sorted cells (4 × 10^6) were cultured for 3 d together with 2 × 10^4 irradiated (3,000 rad) ST2 or ST2-RankL (Nutt et al., 1999) mouse BM stroma cells in flat-bottom 96-well plates. Where indicated, agonistic anti-CD40 mAb (10 μg/ml; FGK45; Bio X Cell) was added.

RNA-seq
Total mRNA was extracted from Lin CD45+ MHCIIhiCD80+ cells using the RNeasy Plus Microkit (Qiagen). The quality of the isolated RNA was controlled by Bioanalyzer 2100 electrophoresis (Agilent). RNA-seq libraries were prepared by the European Molecular Biology Laboratory Genomics core facility in Heidelberg, Germany using a NEBNext rapid illumina RNA-Seq library prep kit (Bibo Scientific) after polyA enrichment with NEBNext Poly(A) Beads (Bibo Scientific), starting with ~1–5 pg of total RNA. Sequencing was performed on an Illumina NextSeq 500 sequencer (Illumina). Low quality reads and adaptor sequences were trimmed out using cutadapt. Reads mapping to ribosomal RNA were filtered out using SortMeRNA. Preprocessed mRNA reads were mapped to the mouse genome (BALB/cJ) from Ensembl version 86 (Mus musculus_balbcj.BALB_cJ_v1.86.gff3) using GSNAP (version 2017–02-25). Gene annotations were downloaded from Ensembl (Mus_musculus_balbcj.BALB_cj_v1.dna_sm.toplevel.fa) using GSNAP. DESeq2 version 1.14.1 was used for feature counting, data normalization, and comparison of the different groups. Differentially expressed genes were selected based on Storey’s q-value < 0.05 and at least 1.5-fold change in transcription activity. For the construction of heat maps, fragments per kilobase of transcript per million mapped reads-normalized counts were used. The raw sequencing data were deposited at the ArrayExpress database under accession no. E-MTAB-7088.

Quantitative PCR
Total mRNA was extracted from cells using the RNeasy Plus Microkit (Qiagen). Reverse transcription was performed using random primers (Thermo Fisher Scientific) and RevertAid reverse transcription (Thermo Fisher Scientific). Gene expression was determined by quantitative PCR reaction using Sybr green and LC480-II cycler (Roche) and quantified using the relative quantification method (Pfaffl, 2001).

Antigen-presentation assay
10^4 HA-TCR hybridoma cells (A5 cells) were co-cultured with 2 × 10^3 APCs in 200 μl IMDM supplemented with 1% FCS. After 17 h, cells were harvested, and GFP expression of A5 cells was measured by flow cytometry.

BM and fetal liver chimeras
BM was depleted of B and T cells using biotinylated mAbs to B220 and TCRβ and streptavidin magnetic activated cell sorting beads (Miltenyi Biotech). Recipient mice were irradiated with 2 × 450 rad and reconstituted with 1 × 10^7 BM cells. For the generation of mixed chimeras, congenically marked BM or fetal liver cells from Rag2−/− (CD45.1+/-), MHCII−/+, RORc−/−, CD40−/−, or Tnfrsf11a−/− mice (CD45.2−/−), and BM or fetal liver cells from WT mice of matching congenic genotype, were mixed at a ratio of 1:1 (5 × 10^6 each) and intravenously injected into lethally irradiated recipient mice (CD45.1−/− CD45.2−/−).

Adoptive T cell transfer
Single cell suspensions of LN cells from HA-TCR-RAG2−/− (CD45.1) and WT mice (CD45.2) were prepared. LN T cells from WT mice were enriched with biotin-conjugated anti-CD4 and streptavidin MicroBeads (Miltenyi Biotech) according to standard procedures. LN T cells were mixed (5 × 10^6) at a ratio of 1:1 i.v. injected into recipient mice (CD45.1−/− CD45.2−/−). After 14 d, the LNs were collected and single cell suspensions were analyzed by flow cytometry.

Statistical analysis
Unless indicated otherwise, statistical significance was assessed using the two-tailed Student’s t test.

Online supplemental material
Fig. S1 shows the phenotype of Aire-expressing LN cells. Fig. S2 shows gating strategies. Fig. S3 shows that MHCIIhiIL-7Rα−/ILC3-like cells express Aire− and costimulatory molecules. Table S1 shows Aire-induced genes in Aire− ILC3-like cells. Table S2 shows Aire-repressed genes in Aire+ ILC3-like cells.

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