

SKIN INFLAMMATION

ROR α -expressing T regulatory cells restrain allergic skin inflammation

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Atopic dermatitis is an allergic inflammatory skin disease characterized by the production of the type 2 cytokines in the skin by type 2 innate lymphoid cells (ILC2s) and T helper 2 (T_H2) cells, and tissue eosinophilia. Using two distinct mouse models of atopic dermatitis, we show that expression of retinoid-related orphan receptor α (ROR α) in skin-resident T regulatory cells (T_{regs}) is important for restraining allergic skin inflammation. In both models, targeted deletion of ROR α in mouse T_{regs} led to exaggerated eosinophilia driven by interleukin-5 (IL-5) production by ILC2s and T_H2 cells. Expression of ROR α in skin-resident T_{regs} suppressed IL-4 expression and enhanced expression of death receptor 3 (DR3), which is the receptor for tumor necrosis factor (TNF) family cytokine, TNF ligand-related molecule 1 (TL1A), which promotes T_{reg} functions. DR3 is expressed on both ILC2s and skin-resident T_{regs}. Upon deletion of ROR α in skin-resident T_{regs}, we found that T_{regs} were no longer able to sequester TL1A, resulting in enhanced ILC2 activation. We also documented higher expression of ROR α in skin-resident T_{regs} than in peripheral blood circulating T_{regs} in humans, suggesting that ROR α and the TL1A-DR3 circuit could be therapeutically targeted in atopic dermatitis.

INTRODUCTION

Atopic dermatitis (AD) is the most common skin inflammatory disease affecting ~17% of children in developed nations (1). AD lesions are characterized by the presence of activated T helper 2 (T_H2) cells, as well as by the expansion of type 2 innate lymphoid cells (ILC2s) (2–4). Both T_H2 cells and ILC2s may contribute to allergic skin inflammation in AD. Cutaneous inflammation elicited by topical application of calcipotriol (MC903), a low-calcemic analog of vitamin D, has been used as a mouse model of acute AD (5, 6). Allergic inflammation in this model is accompanied by expansion of ILC2s driven by epithelial cytokines (2, 4). More importantly, it is dependent on ILC2s; it is preserved in *Rag1*^{−/−} mice and is severely attenuated in *Tslpr*^{−/−} mice, ILC-depleted *Rag1*^{−/−} mice, and ILC2-deficient *Rora*^{sg/sg}/wild-type (WT) bone marrow chimeras (2, 4). Cutaneous inflammation elicited by repeated epicutaneous (EC) application of ovalbumin (OVA) or peanut extract to tape-stripped mouse skin provides an antigen-driven mouse model of acute AD (7–9). Allergic inflammation in this model is dependent on T cells, because it is abolished in *Rag2*^{−/−} mice (9, 10).

CD4⁺FOXP3⁺ T regulatory cells (T_{regs}) constitute a substantial subset of immune cells residing in murine and human skin (11). Lack of T_{regs} in humans and mice results in immune dysregulation associated with allergic skin inflammation (12, 13). T_{reg} numbers are unaltered in AD skin lesions (14). Thus, the role of skin-resident T_{regs} in controlling

allergic skin inflammation is unclear. Here, we have dissected the molecular architecture of skin-resident T_{regs} and identified retinoid-related orphan receptor α (ROR α) as a regulator of genes in T_{regs} responsible for suppressing allergic skin inflammation.

RESULTS

Skin-resident T_{regs} exhibit an activated signature and express the transcription factor ROR α

Specialization of tissue-resident T_{regs} is an important factor in maintaining tissue homeostasis and modulating local immune responses. To investigate whether skin-resident T_{regs} exhibit a specialized phenotype, we compared the phenotype of skin-resident T_{regs} and T_{regs} in skin-draining lymph node (dLN). About 45% of CD4⁺ T cells in ear skin expressed FOXP3 compared with ~20% of CD4⁺ T cells in dLNs (Fig. 1A). Skin T_{regs} localized around dermal blood vessels and interfollicular areas (fig. S1A). We compared the transcriptome of CD3⁺CD4⁺YFP⁺ T_{regs} from the skin and dLN of *Foxp3*^{eyfp-cre} mice. Skin T_{regs} differed from dLN T_{regs} by more than 5000 genes [fold change > 2; false discovery rate (FDR) < 0.05]. Skin T_{regs} were enriched for the expression of genes encoding signaling receptors [*Icos* and *Il1rl1* (ST2)], activation markers (*Cd44* and *Klrg1*), effector molecules (*Il10*, *Ctla4*, and *Areg*), and tissue-homing receptors (*Ccr3*, *Ccr8*, and *Ccr10*) (Fig. 1B). Flow cytometry demonstrated that the percentage of T cells that expressed ST2, ICOS, and CD44 and the expression levels of these markers were significantly higher in skin T_{regs} than in dLN T_{regs} (Fig. 1C). *Rora*, the gene encoding the transcriptional regulator ROR α , was highly up-regulated in skin T_{regs} (Fig. 1B). This was confirmed by quantitative polymerase chain reaction (qPCR) (Fig. 1D). ROR α expression was significantly higher in CD4⁺CD25⁺CD127^{lo} skin T_{regs} than in circulating T_{regs} in humans (Fig. 1E). Human skin T_{regs}, similar to mouse skin T_{regs}, display an activated signature with increased expression of ICOS, CTLA4, and CD44 (15).

To examine and map the fate of ROR α -expressing T_{regs}, we bred *Rora*^{cre} mice to *Rosa26Yfp* (R26Y) mice. In *Rora*^{cre}R26Y mice, yellow fluorescent protein (YFP) marks cells that are expressing or previously

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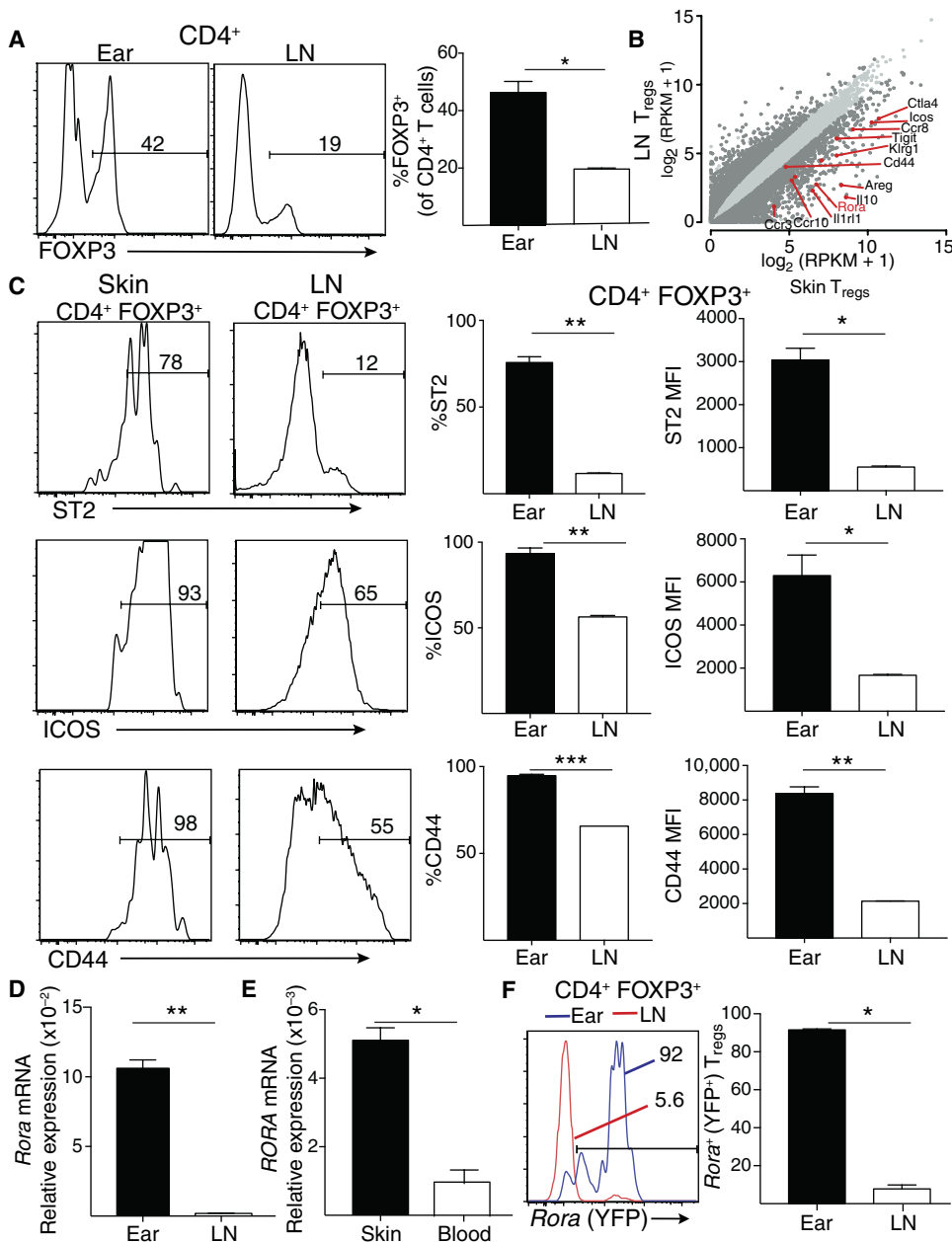


Fig. 1. Skin-resident T_{regs} exhibit an activated signature and express the transcription factor RORα. (A) Representative flow cytometric analysis (left) and quantification (right) of FOXP3⁺ (CD3⁺CD4⁺YFP⁺) cells among CD4⁺ T cells in ear skin compared with dLNs from *Foxp3^{eyfp-cre}* mice (*n* = 3 mice per group). (B) Scatterplot of log₂ (RPKM + 1) values of genes expressed in skin T_{regs} (x axis) compared with LN T_{regs} (y axis) determined by NGS transcriptomic analysis. Genes that differ by more than twofold are shown in dark gray. Select genes are identified. (C) Representative flow cytometric analysis (left) and quantification (right) of CD44⁺, ICOS⁺, and ST2⁺ skin and dLNs T_{regs} and the mean fluorescence intensity (MFI) of these markers (*n* = 3 mice per group). (D) *Rora* expression levels in sorted T_{regs} from skin and dLNs from *Foxp3^{eyfp-cre}* mice (*n* = 3 mice per group). (E) *Rora* expression levels in sorted T_{regs} (CD4⁺CD25⁺CD127^{lo}) from blood and skin of healthy donors (*n* = 2). (F) Representative flow cytometric analysis (left) and quantification (right) of *Rora*⁺(YFP⁺)-expressing T_{regs} in skin and dLN of *Rora^{cre}Rosa^{ylp/yfp}* mice (*n* = 2 mice per group). Columns and bars represent means and SEM. **P* < 0.05, ***P* < 0.01, ****P* < 0.001.

expressed *Rora*. Most of the skin T_{regs} (>90%) in *Rora^{cre}R26Y* mice expressed YFP compared with a small fraction (~5%) of T_{regs} from dLNs (Fig. 1F). *Rora*⁺(YFP⁺) T_{regs} in the skin uniformly expressed the transcription factor HELIOS, but not RORγt (fig. S1B), suggesting that RORα-

expressing skin T_{regs} are natural T_{regs}. The percentage of ICOS⁺ and ST2⁺ T_{regs} and the levels of ICOS and ST2 were significantly higher in *Rora*⁺(YFP⁺) T_{regs} than in *Rora*⁻(YFP⁻) T_{regs} in dLNs (fig. S1C). A negligible subset (<1%) of thymic T_{regs} were *Rora*⁺(YFP⁺) (fig. S1D), suggesting that RORα⁺ T_{regs} expand and/or are induced in peripheral tissues.

We used *Rora^{cre}R26Y* mice to investigate *Rora* expression by cell subpopulations in the skin. In addition to T_{regs}, a fraction of CD3⁺CD4⁺CD25⁻ T cells, CD3⁺CD8⁺ T cells, CD3⁺TCRγδ^{low} dermal γδ T cells, CD3⁺TCRγδ^{high} epidermal γδ T cells, and CD45⁺Lin⁻ ILCs in the skin were YFP⁺ (*Rora*⁺) (fig. S2A). In addition, a fraction of CD45⁻EpCAM⁺ keratinocytes that are mostly derived from the basal layer of the epidermis and a fraction of CD45⁻EpCAM⁻ cells, which contain a mixture of mature keratinocytes and fibroblasts in the skin, were YFP⁺ (*Rora*⁺) (fig. S2A). The percentages of YFP⁺ (*Rora*⁺) cells among skin cell subpopulations were not significantly altered following MC903 treatment (fig. S2, B and C). These results show that RORα expression was not restricted to skin T_{regs}.

RORα deficiency in T_{regs} results in exaggerated allergic skin inflammation in response to topical application of MC903

RORα is necessary for the development of ILC2s (16), promotes T_H17 cell differentiation, and antagonizes FOXP3 in vitro (17), suggesting a potential pro-inflammatory role. To understand how RORα regulates the function and/or maintenance of skin T_{regs}, we generated *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice. Fluorescence-activated cell sorting (FACS) analysis of skin population of cells from *Foxp3^{eyfp}* mice for enhanced green fluorescent protein (eGFP) expression revealed that *Foxp3* expression was restricted to CD4⁺ T cells and was not detected in any other additional skin cell population that expressed *Rora* in the skin, including CD8⁺ T cells, dermal and epidermal γδ T cells, ILCs, and CD45⁻ cells (fig. S3). In addition, none of the *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice had weight loss or developed the staggerer phenotype observed in RORα-deficient *Rora^{sg/sg}*

mice (18). Furthermore, the numbers of ILCs and γδ T cells in the skin were not reduced in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice. These results suggest that *Rora* is deleted specifically in T_{regs} of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice. RNA sequencing (RNA-seq) analysis of T_{regs} revealed complete deletion of

the floxed fourth exon of *Rora* in these mice (fig. S4A). The numbers of YFP⁺ T_{regs} were not altered in the skin or dLNs of these mice (fig. S4B), indicating that ROR α is not required for the accumulation or maintenance of T_{regs} in the skin. The cytokine interleukin-10 (IL-10) is important for T_{reg} function in the gut and lungs (19). There was an increased percentage of IL-10⁺ T_{regs} in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice compared with controls (fig. S4C).

Topical application of MC903 to ear skin of WT mice results in increased dermal thickness and infiltration of CD45⁺ cells that include eosinophils and CD4⁺ T cells (5). There was an increased ear thickness, accompanied with an intense cellular infiltrate, and significantly increased dermal thickness in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice compared with *Foxp3^{eyfp-cre}* controls (Fig. 2, A to C). FACS analysis revealed a threefold increase in dermal infiltration by CD45⁺ cells in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice compared with *Foxp3^{eyfp-cre}* controls (Fig. 2D). Eosinophils accounted for ~45% of CD45⁺ cells in the dermis of MC903-treated *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice, compared with 15% in controls, yielding an eightfold increase in eosinophil numbers (Fig. 2E). The percentages of basophils (ckit⁺IgE⁺), mast cells (ckit⁺IgE⁺), neutrophils (CD11b⁺Gr1^{hi}), T effector cells (T_{eff}) (CD4⁺FOXP3⁻), T_{regs} (CD4⁺FOXP3⁺), and ILCs (Lin⁻CD90⁺) infiltrating MC903-treated skin were comparable in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice and controls. Nevertheless, the numbers of these cell populations were two- to threefold higher in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice (Fig. 2, F and G), reflecting the approximately threefold increase in CD45⁺ cells. MC903-driven allergic inflammation in mice of C56BL/6 background is largely dependent on thymic stromal lymphopoietin (TSLP) (2, 4). The exaggerated cutaneous inflammatory response in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice, which are on a C56BL/6 background, was not due to increased *Tslp* expression (fig. S5A). Serum immunoglobulin E (IgE) levels were higher in MC903-treated *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice than in controls (fig. S5B), indicative of a heightened type 2 response.

ROR α deficiency in T_{regs} results in increased expression of eotaxins and IL-5 in MC903-treated skin

The proportion of eosinophils in blood was comparable in MC903-treated *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice and controls (fig. S6A), suggesting that the exaggerated eosinophilia in MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice resulted from increased eosinophil recruitment. Eotaxins are the major eosinophil chemoattractants (20). There was increased expression of *Ccl11* and *Ccl24*, which encode for eotaxins 1 and 2, in MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice compared with controls (Fig. 3A). IL-5 plays a critical role in tissue eosinophilia by synergizing with eotaxins and promoting eosinophil survival in tissues (21, 22). IL-5 levels were significantly higher in MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice than of controls (Fig. 3B). IL-4 and IL-13 levels were comparable in the two groups (fig. S6B).

IL-5 is predominantly produced by ILC2s and by a subset of activated T_H2 cells (23). ILC2s exist as preactivated resident sentinels in the dermis that rapidly release IL-5 and IL-13 upon stimulation (21). In contrast, T_H2 cells are recruited to the skin at a later stage of allergic inflammation. MC903 treatment up-regulated IL-33/ST2 and CD69 expression, down-regulated CD25 expression, and had negligible effect on KLRG1 expression on skin Lin⁻CD90⁺ ILCs, but the changes were comparable in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice and *Foxp3^{eyfp-cre}* controls. However, in *Il5*, mRNA levels were significantly increased in ILCs from MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice compared with controls (Fig. 3C). There was also a twofold increase in *Il4* and *Il13* mRNA levels in ILCs from MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}*

mice; however, it did not reach statistical significance (fig. S6C). There was a significant increase in CD4⁺FOXP3⁺ IL-5⁺, but not CD4⁺IL-13⁺ or CD4⁺IL-4⁺, T_{eff}s in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice compared with controls (Fig. 3D). The chemokine CCL8 is up-regulated in AD skin lesions (24) and is critical for the recruitment of CCR8-expressing IL-5⁺ T_H2 cells to the skin in a mouse model of AD (24). *Ccl8* expression was strongly increased in MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice (Fig. 3E). In contrast, *Ccl17*, *Ccl22*, *Ccl4*, and *Ccl5* expression was comparable in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice and controls (fig. S6D). *Cxcl1*, *Ccl2*, and *Ccl7* expression demonstrated a trend toward an increase in MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice, which could underlie the increased influx of myeloid cells in these mice (Fig. 2F).

ROR α deficiency in T_{regs} alters the expression of genes involved in T_{reg} cell migration and function and skews T_{regs} to IL-4-producing effectors

To gain an understanding of how ROR α regulates the function of skin T_{regs}, we performed next-generation sequencing (NGS) transcriptomic analysis on T_{regs} isolated from untreated and MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice and *Foxp3^{eyfp-cre}* controls (table S1 and fig. S7). We observed a change in ~1700 genes across the four groups examined (fold change > 2; FDR < 0.05) (Fig. 4A). Expression of the central circadian rhythm genes *Nr1d1* and *Nr1d2* was decreased in skin T_{regs} from *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice compared with controls, consistent with the role of ROR α as a circadian rhythm regulator (25). Genes involved in signaling via transforming growth factor- β (TGF β) (*Smad3*), tumor necrosis factor- α (TNF α) (*Tnfa*), nuclear factor κ B (NF- κ B) (*Irak4* and *Tirap*), and mitogen-activated protein kinase (MAPK) (*Fos* and *Jun*) and in cell adhesion (*Icam2* and *Itga4*) were comparably expressed in T_{regs} from untreated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice and *Foxp3^{eyfp-cre}* controls and underwent comparable changes after MC903 treatment. Genes in the phosphatidylinositol 3-kinase (PI3K)/AKT pathway were down-regulated in ROR α -deficient skin T_{regs}. Dysregulated PI3K/AKT signaling affects *Foxp3* and *Il2ra* expression in T_{regs} and increases their conversion to T_H1 cells (26). We did not observe any effect on *Foxp3*, *Il2ra*, or *Ifng* expression in our transcriptomic and flow cytometric analyses of skin T_{regs} from *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice. Several genes encoding chemokines and chemokine receptors (*Ccl2*, *Ccr3*, and *Ccr5*) were up-regulated upon allergic skin inflammation in all mice, but to greater extent in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice. Up-regulation of these genes is consistent with the increased numbers, and higher velocity, of T_{regs} in MC903-treated skin of *Foxp3^{eyfp-cre}* mice (Fig. 2G; fig. S8, A and B; and movies S1 and S2). Furthermore, T_{regs} in MC903-treated skin showed less directed movement (fig. S8C). Expression of *Ccr6* and CCR6, thought to be important for migration of T_{regs} into neonatal skin (27), was strongly decreased in skin T_{regs} from *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice compared with controls both before and after MC903 treatment (Fig. 4, A to C). However, the numbers of skin T_{regs} in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice were not reduced (Fig. 3G).

T_{reg} suppressive activity is mediated in part by the nucleotides adenosine and cyclic adenosine 3',5'-monophosphate (cAMP) (28). T_{regs} from untreated and MC903-treated skin showed strongly decreased expression of *Nt5e*, which encodes the 5' ectonucleotidase CD73 that metabolizes AMP to adenosine (28), and reduced surface expression of CD73 compared with controls (Fig. 4, B and D), whereas expression of *Pde3b*, which encodes the phosphodiesterase 3B that breaks down cAMP (29), was increased (Fig. 4B). Expression of *Gzmb*, which encodes granzyme B that mediates T_{reg} cytotoxic activity, was up-regulated

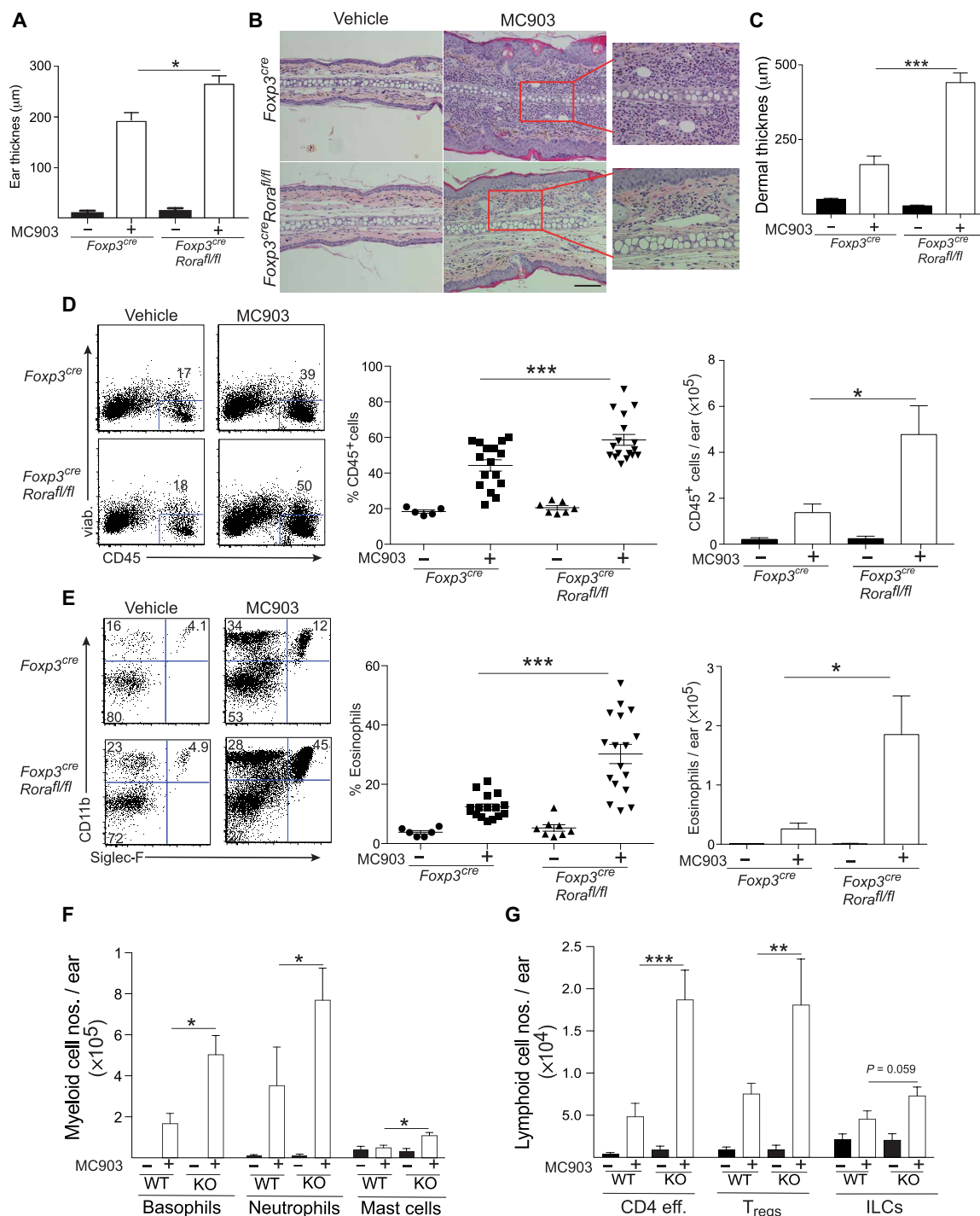


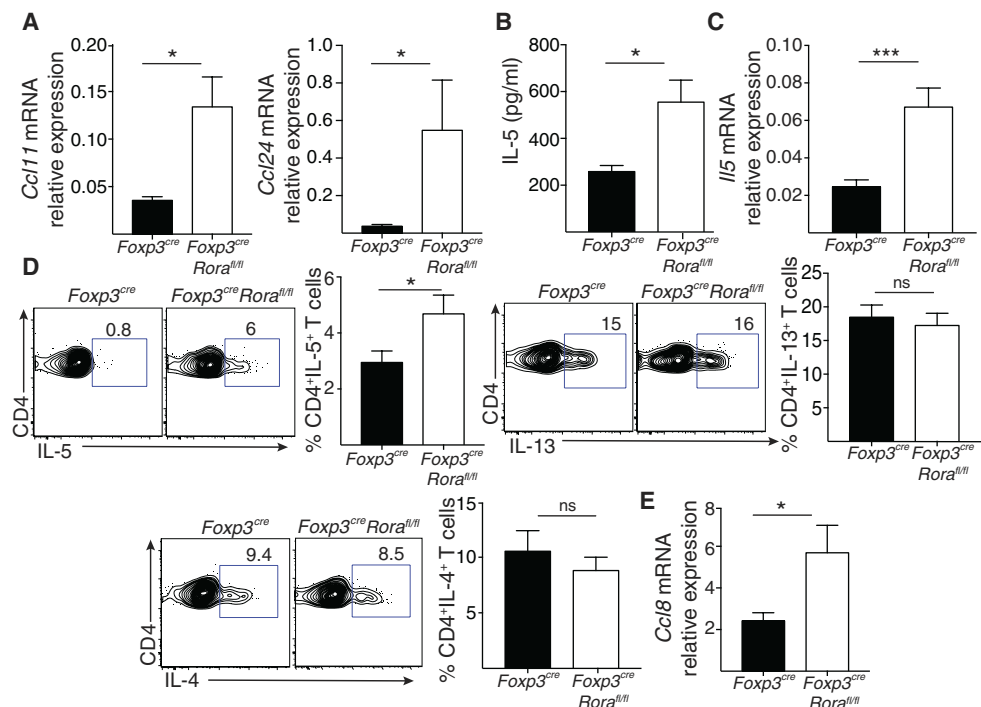
Fig. 2. ROR α deficiency in T_{regs} results in exaggerated skin inflammation in response to topical application of MC903. (A to G) Quantification of ear thickness at day 7 (A); representative H&E-stained sections (B); quantification of dermal thickness (C); representative FACS analysis (left) and quantification of the percentages (middle) and numbers (right) of CD45⁺ cells (D), eosinophils (E), mast cells, neutrophils, and basophils (F), and CD4⁺FOXP3⁺ T_{regs}, CD4⁺FOXP3⁺ T_{effs}, and ILCs (G) in vehicle or MC903-treated ears of *Foxp3^{cre}Rora^{fl/fl}* mice and *Foxp3^{cre}* controls. $n = 3$ to 8 mice per group. Columns and bars represent means and SEM. * $P < 0.05$, *** $P < 0.001$.

(Fig. 4B), indicating that not all genes involved in T_{reg} function were down-regulated in the absence of ROR α .

Expression of IL-4 in T_{regs} inhibits their ability to suppress T_H2 cells and ILC2s (30, 31). *Il4* levels were elevated in skin T_{regs} from *Foxp3^{cre}Rora^{fl/fl}* mice compared with controls (Fig. 4, A and B).

This was confirmed by qPCR (Fig. 4E). Furthermore, the percentage of CD4⁺FOXP3⁺ T_{regs} among IL-4-expressing CD4⁺ cells in MC903-treated skin was significantly higher in *Foxp3^{cre}Rora^{fl/fl}* mice than in controls (Fig. 4F). These results suggest that ROR α expression prevents the conversion of T_{regs} into IL-4-producing effectors. The transcription

Fig. 3. Increased expression of eotaxins and IL-5 in MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice. (A to E) Relative *Ccl11* and *Ccl24* mRNA expression (A); IL-5 levels (B); relative *Il5* expression in sorted Lin[−]CD90⁺ ILCs (C); representative FACS analysis and quantitation of the percentages of CD4⁺IL-5⁺, CD4⁺IL-13⁺, and CD4⁺IL-4⁺ T_{effs} (D); and relative *Ccl8* mRNA expression in MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice and *Foxp3^{eyfp-cre}* controls (E). *n* = 4 to 7 mice per group. Columns and bars represent means and SEM. **P* < 0.05, ****P* < 0.001. ns, not significant.



factor RUNX1 inhibits *Il4* expression in T_{regs} (32). *Runx1* expression was decreased in T_{regs} from *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice (Fig. 5B), suggesting that reduced RUNX1 activity may derepress *Il4* expression in ROR α -deficient T_{regs}. T_{regs} from MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice, but not controls, expressed *Ccl8* and *Ccl24* transcripts (Fig. 4B), suggesting that these T_{regs} contribute to the exaggerated eosinophil-dominated allergic skin inflammation in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice.

ROR α expression in T_{regs} promotes expression of the TL1A ligand DR3 and restrains TL1A-driven allergic inflammation elicited by cutaneous application of MC903

Tnfrsf25 encodes the TNF receptor family member DR3 (death receptor 3), which is expressed constitutively on T cells, including T_{regs} and ILC2s (33, 34). *Tnfrsf25* expression, as determined by RNA-seq analysis, and DR3 surface expression, as determined by FACS analysis, were both strongly reduced in skin T_{regs} from *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice compared with *Foxp3^{eyfp-cre}* controls (Figs. 4B and 5, A and B). In contrast, DR3 surface expression by skin ILC2s was comparable in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice and controls (Fig. 5C). The DR3 ligand TL1A is released by endothelial and myeloid cells. TL1A synergizes with the epithelial cytokines IL-33, IL-7, and IL-25 to enhance IL-5 expression in human and murine ILC2s and to promote allergic inflammation (33, 35, 36). TL1A also acts on T_{regs} to increase their proliferation and their ability to suppress allergic airway inflammation (34). Skin TL1A levels were not altered after MC903 treatment and were comparable in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice and *Foxp3^{eyfp-cre}* controls (Fig. 5D). Given this finding, we tested the hypothesis that selective down-regulation of the TL1A receptor DR3 on T_{regs} from skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice may play an important role in the exaggerated MC903-driven allergic inflammation observed in these mice. MC903-mediated eosinophilia was attenuated in *Tnfrsf25^{-/-}* mice (Fig. 5E), demonstrating a role for TL1A in MC903-driven allergic skin inflammation. Intradermal in-

jection of TL1A into ear skin resulted in a significant increase in the percentage and numbers of eosinophils, but not neutrophils, in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice compared with controls (Fig. 6F). More importantly, local TL1A blockade during MC903 treatment by intradermal injection of neutralizing antibody to TL1A significantly reduced MC903-driven allergic skin inflammation in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice. This was evidenced by a significant decrease in dermal thickness, infiltration by CD45⁺ cells and eosinophils, and expression of *Il5* and *Ccl8* compared with isotype control antibody-treated mice (Fig. 6, G to J). These results suggest that ROR α expression in T_{regs} restrains TL1A-mediated allergic skin inflammation and eosinophilia elicited by cutaneous application of MC903.

ROR α deficiency in T_{regs} results in exaggerated skin inflammation in response to EC sensitization

To investigate whether ROR α deficiency in T_{regs} plays a role in restraining antigen-driven T cell-dependent allergic skin inflammation, we subjected *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice and *Foxp3^{eyfp-cre}* controls to EC sensitization. EC sensitization was elicited by repeated application of the antigen OVA to tape-stripped skin, as illustrated in Fig. 6A. Skin inflammation in this model shares many characteristics of acute AD skin lesions, including epidermal thickening, dermal infiltration by CD45⁺ cells (including eosinophils), and increased expression of type 2 cytokines (8, 9). *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice EC sensitized with OVA exhibited significantly increased epidermal thickness and significantly increased infiltration by CD45⁺ cells compared with *Foxp3^{eyfp-cre}* controls EC sensitized with OVA (Fig. 6, B to D). Furthermore, the numbers of all cell populations analyzed, including eosinophils, basophils, neutrophils, mast cells, CD4⁺Foxp3[−] T cells, T_{regs}, and ILCs, were two- to fourfold higher in OVA-sensitized skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice compared with *Foxp3^{eyfp-cre}* controls (Fig. 6, E and F). *Il4*, but not *Il13*, mRNA levels in OVA-sensitized skin were significantly higher in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice than in controls (Fig. 6G). *Il5* mRNA was not

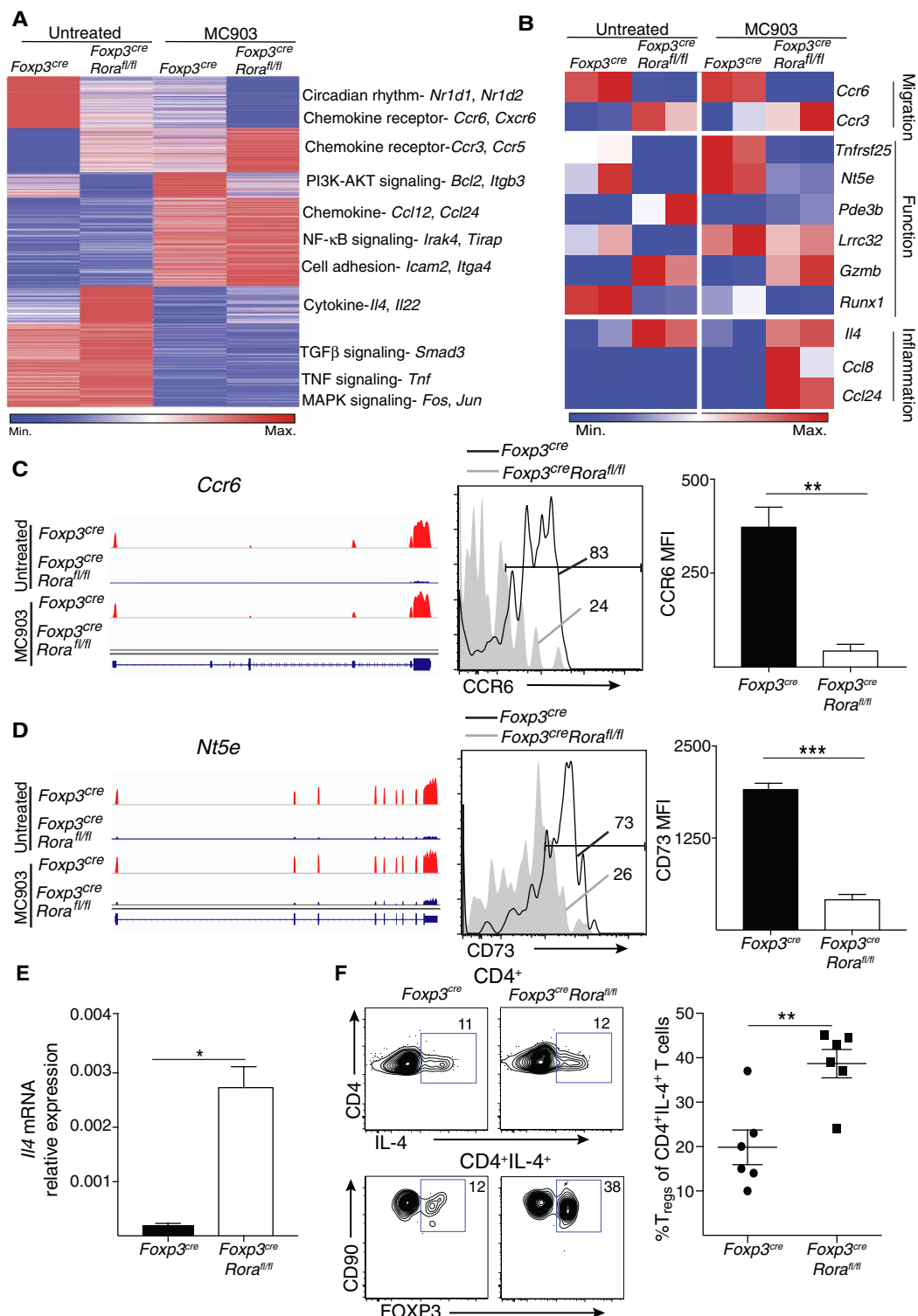


Fig. 4. ROR α deficiency in T_{regs} alters the expression of genes involved in T_{reg} cell migration and function and skews T_{regs} to IL-4-producing effectors. (A) Heat map showing relative expression of genes clustered by K-mean values in skin T_{regs} of *Foxp3^{cre}* and *Foxp3^{cre}Rora^{fl/fl}* mice in the steady state and after MC903 treatment (*n* = 4 to 5 mice per group). **(B)** Heat map showing the relative expression of select chemotaxis, function, and inflammation genes in skin T_{regs} from *Foxp3^{cre}Rora^{fl/fl}* mice and controls (*n* = 4 to 5 mice per group). **(C and D)** RNA-seq tracing of *Ccr6* and *Nt5e* expression (left), representative FACS analysis (middle), and MFIs (right) of CCR6 and CD73 expression in skin T_{regs} of *Foxp3^{cre}* and *Foxp3^{cre}Rora^{fl/fl}* mice (*n* = 4 to 5 mice per group). The numbers in the FACS panels represent the percentage of positive cells relative to fluorescence minus one (FMO) control. **(E)** Relative *Il4* mRNA levels in T_{regs} from MC903-treated skin of *Foxp3^{cre}* and *Foxp3^{cre}Rora^{fl/fl}* mice and controls (*n* = 4 to 5 mice per group). **(F)** Representative FACS analysis of IL-4 expression in CD4⁺ cells and of FOXP3 versus CD90 expression in IL-4⁺CD4⁺ cells (left) and quantitation of the percentage of IL-4⁺CD4⁺FOXP3⁺ cells among IL-4⁺CD4⁺ cells in the skin of MC903-treated *Foxp3^{cre}* and *Foxp3^{cre}Rora^{fl/fl}* mice and controls.

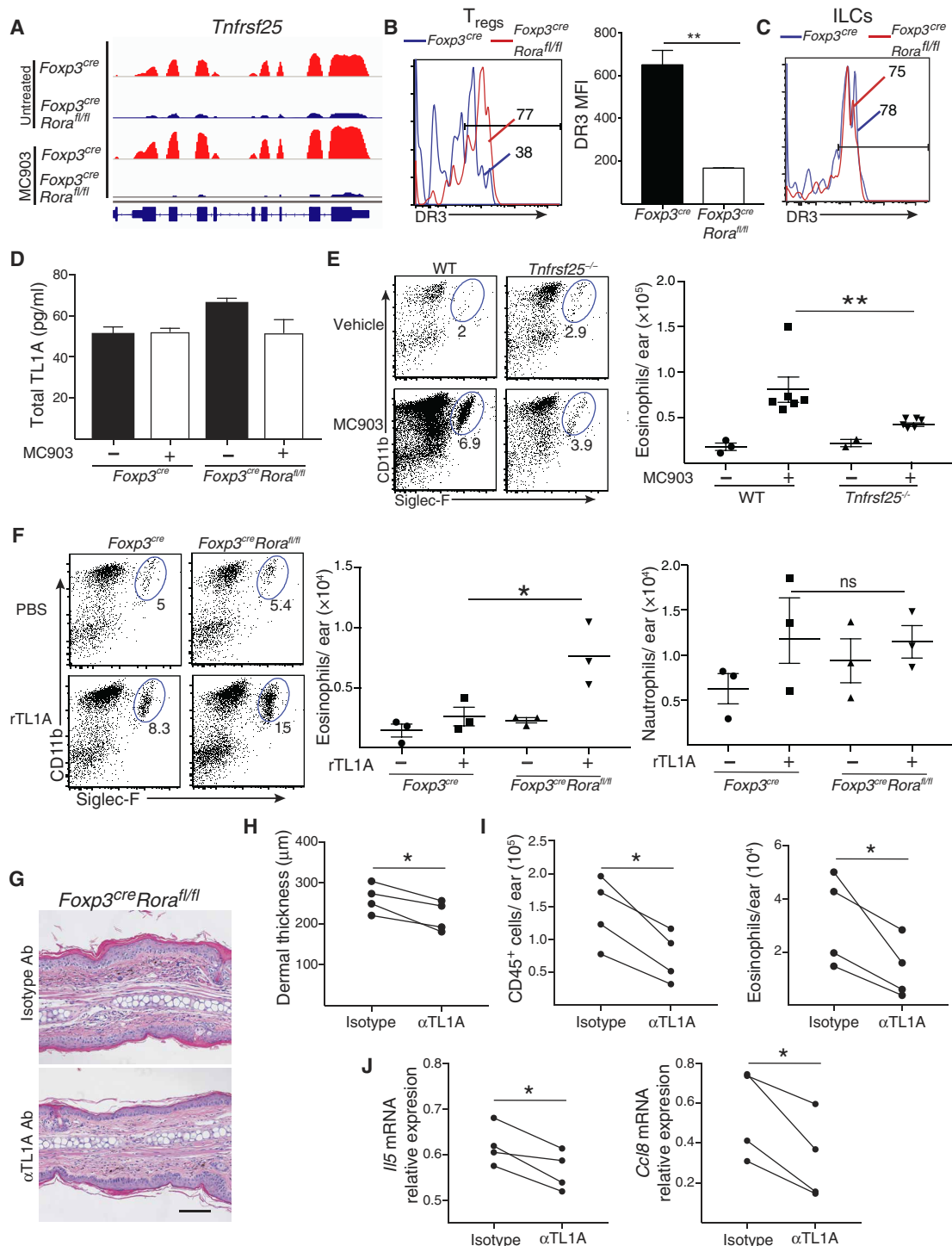


Fig. 5. ROR α expression in T_{reg} s promotes expression of the TL1A receptor DR3 and restrains TL1A-driven allergic inflammation elicited by cutaneous application of MC903. (A) RNA-seq tracing of *Tnfrsf25* expression in skin T_{reg} s from untreated and MC903-treated skin of $Foxp3^{cre}$ and $Foxp3^{cre} Rora^{fl/fl}$ mice. (B) Representative FACS analysis (left) and MFIs (right) of DR3 expression by skin T_{reg} s of $Foxp3^{cre} Rora^{fl/fl}$ mice and $Foxp3^{cre}$ controls ($n = 3$ mice per group). The numbers in the FACS panels represent the percentage of positive cells relative to FMO control. (C) Representative FACS analysis of DR3 expression by ILCs from the skin of $Foxp3^{cre} Rora^{fl/fl}$ mice and $Foxp3^{cre}$ controls. Results are representative of three independent experiments. The numbers in the FACS panels represent the percentage of positive cells relative to FMO control. (D) TL1A levels in vehicle and MC903-treated ear skin of $Foxp3^{cre} Rora^{fl/fl}$ mice and $Foxp3^{cre}$ controls ($n = 4$ mice per group). (E) Representative FACS analysis (left) and quantification (right) of CD11b⁺Siglec-F⁺ eosinophils in MC903-treated ears of *Tnfrsf25*^{-/-} mice and WT controls. (F) Representative FACS analysis (left) and quantification (right) of CD11b⁺Siglec-F⁺ eosinophils and CD11b⁺Gr1^{high} neutrophils in TL1A-injected skin of $Foxp3^{cre} Rora^{fl/fl}$ mice and $Foxp3^{cre}$ controls ($n = 3$ mice per group). (G to J) Representative H&E-stained sections (G), quantification of dermal thickness (H), quantification of CD45⁺ cells (right) and CD11b⁺Siglec-F⁺ eosinophils (left) (I), and relative mRNA expression of *Il5* (right) and *Ccl8* (left) (J) in MC903-treated ears of $Foxp3^{cre} Rora^{fl/fl}$ mice injected with anti-TL1A antibody or isotype control ($n = 4$ mice per group). Columns and bars represent means and SEM. * $P < 0.05$, ** $P < 0.01$.

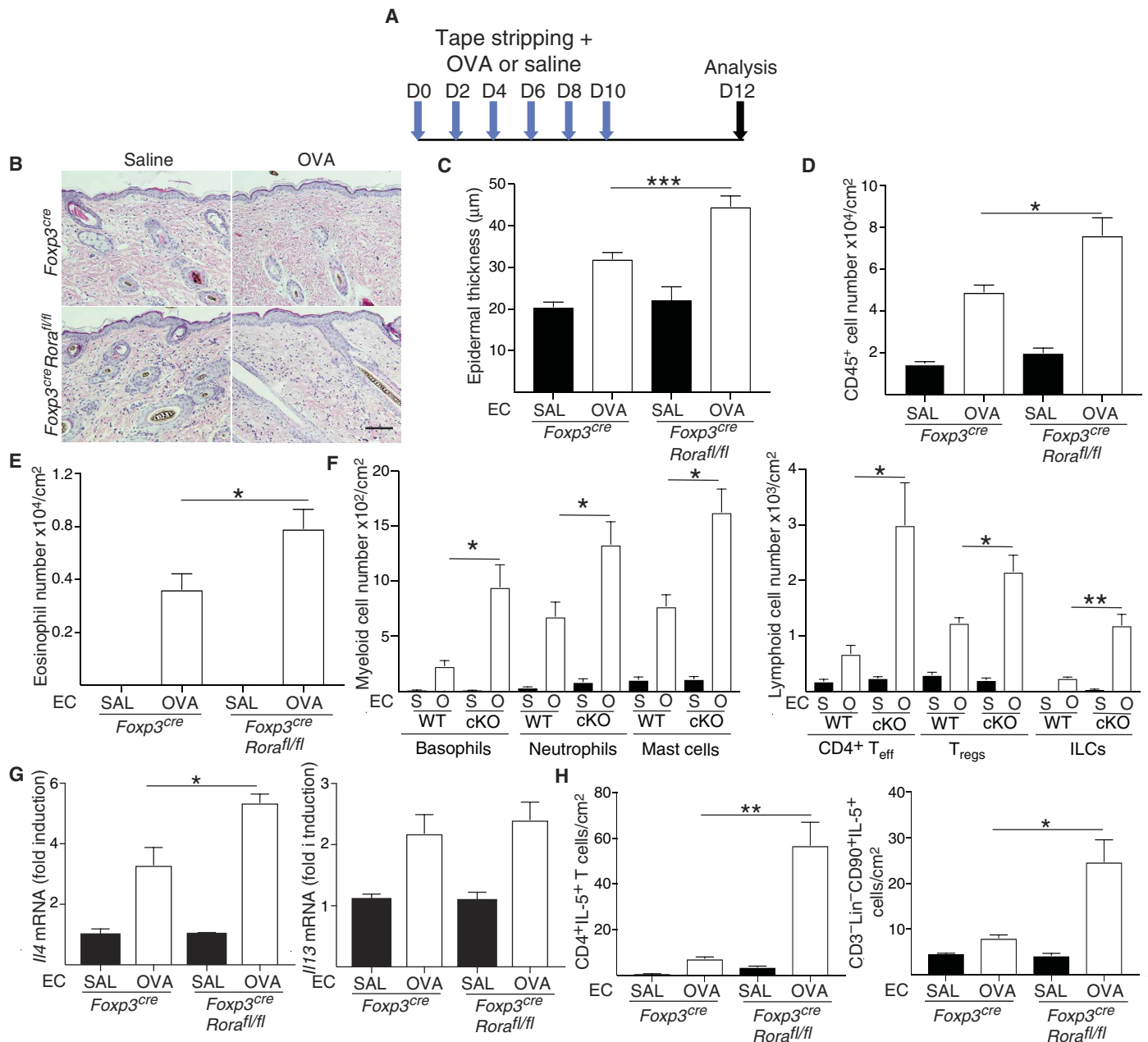


Fig. 6. ROR α deficiency in T_{regs} results in exaggerated skin inflammation in response to EC sensitization. (A) Schematic of the experimental mouse model. (B to H) Representative H&E-stained sections (B); quantification of epidermal thickness (C); number of CD45⁺ cells (D), CD11b⁺Siglec-F⁺ eosinophils (E), mast cells, neutrophils, and basophils (left), and CD4⁺FOXP3⁺ T_{regs}, CD4⁺FOXP3⁺ T_{regs}, and ILCs (right) (F); relative *Il4* (right) and *Il13* (left) mRNA expression (G); and numbers of IL-5⁺ CD4⁺ T cells and ILCs (H) in saline and OVA-sensitized skin of *Fxp3^{cre}Rora^{fl/fl}* mice (also designated as cKO) and *Fxp3^{cre}* controls (also designated as WT). *n* = 3 to 7 mice per group. Columns and bars represent means and SEM. **P* < 0.05, ****P* < 0.001.

detectable in sensitized skin in either group. Nevertheless, intracellular FACS analysis revealed that OVA sensitization caused a small but significant increase in the numbers of IL-5⁺ ILCs and IL-5⁺ CD4⁺FOXP3⁺ T_{regs} in *Fxp3^{cre}* control mice. The numbers of IL-5⁺ ILCs and IL-5⁺ CD4⁺FOXP3⁺ T_{regs} were four- to fivefold higher in OVA-sensitized skin of *Fxp3^{cre}Rora^{fl/fl}* mice than in controls (Fig. 6H). OVA sensitization did not result in significant changes in IL-33R/ST2, CD69, CD25, or KLRG1 expression by skin ILCs in *Fxp3^{cre}Rora^{fl/fl}* or *Fxp3^{cre}* controls. These results suggest that

ROR α ⁺ T_{regs} play an important role in constraining antigen-driven skin inflammation.

DISCUSSION

We show that skin T_{regs} express high levels of the transcription factor ROR α . Deletion of *Rora* in T_{regs} does not alter the number of skin T_{regs} but results in exaggerated type 2 allergic skin inflammation in response to topical application of MC903 or EC sensitization with OVA. Thus,

we have identified ROR α as a regulator of T_{reg} genes responsible for suppressing allergic skin inflammation.

The vast majority of mouse skin T_{regs} expressed ROR α and had an activated phenotype. In contrast, a small minority of T_{regs} in skin dLNs expressed ROR α and had an activated phenotype. It remains to be determined whether circulating ROR α ⁺ T_{regs} are specifically attracted to the skin or whether the skin environment drives ROR α expression in T_{regs}. The numbers of skin T_{regs} are not altered in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice. Furthermore, although the majority of human blood T_{regs} express the skin-homing receptor CLA (37), human blood T_{regs} expressed fivefold less *RORA* mRNA compared with skin T_{regs}. These findings argue for local acquisition of ROR α expression by T_{regs} in the skin.

We demonstrated that ROR α expression in T_{regs} restrains allergic skin inflammation induced by topical application of MC903, an AD model dependent on ILC2s (2, 4). This was evidenced by increased ear swelling and dermal thickness in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice, with a threefold increase in the influx of inflammatory cells that included T cells, basophils, neutrophils, and mast cells, and a selective enrichment in eosinophils that showed an eightfold increase over controls. Type 2 cytokines, such as IL-4, are documented to drive eotaxin expression (21, 38). Increased eosinophilia in MC903-treated skin of *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice may be explained by synergy between increased skin IL-5 expression and increased skin and T_{regs} eotaxin expression, likely driven by increased expression of IL-4 and IL-13. The exaggerated skin inflammation in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice was not caused by increased cutaneous expression of TSLP, the epithelial cytokine essential for MC903-driven skin inflammation in mice on the C57BL/6 background, the background of the *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice we used. ROR α was essential for repressing IL-5 expression in fast-responding ILC2s and for restricting the CCL8-dependent recruitment of IL-5⁺ T_{H2} effector cells to the skin, likely by dampening the expression of *Ccl8* in the skin, and particularly in skin T_{regs}. ROR α also repressed IL-13 and IL-4 expression by skin ILCs, although the effect did not reach statistical significance, but had no effect on IL-4 and IL-13 expression by T cells. We propose that in addition to their role in restraining adaptive immunity, a central function of T_{regs} resident in barrier interfaces, such as skin, is to inhibit the rapid activation of innate lymphocytes. The unrestrained activation of these innate sentinels may contribute to acute flare-ups in allergic diseases.

ROR α regulated the expression of several genes important for T_{reg} migration and function. Changes in chemokine receptor expression may underlie the increased motility of T_{regs} in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice. Our data suggest that decreased expression by ROR α -deficient T_{regs} of *Tnfrsf25* encoding DR3, a gene important for T_{reg} function, contributed to the enhanced skin inflammation in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice. The exaggerated skin inflammation in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice may be a direct effect of decreased TL1A activation of T_{regs} and/or increased availability of TL1A to activate ILC2s. Definitive evidence of the role of DR3 expression on T_{regs} in limiting allergic skin inflammation and its mechanism of action awaits the generation and study of mice with selective deficiency of *Tnfrsf25* in T_{regs} and/or ILC2s. Furthermore, our data indicate that ROR α restrains the conversion of T_{regs} into IL-4–producing effector cells, possibly because ROR α drives the expression of *Runx1*, which inhibits *Il4* gene transcription. Derepression of the T_{H2} proinflammatory genes in ROR α -deficient skin T_{regs} likely contributes to the enhanced allergic skin response in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice. Furthermore, IL-10 expression was increased in ROR α -deficient skin T_{regs}. The transcription factor AhR (aryl hydro-

carbon receptor) enhances IL-10 production in T_{regs} (39), whereas IL-4 suppresses it (40). We observed increased *Ahr* and *Il4* expression in ROR α -deficient skin T_{regs}. Increased expression of AhR and IL-4 may underlie the enhanced IL-10 expression by these cells.

In addition to its role in suppressing ILC2-dependent allergic skin inflammation driven by topical application of MC903, ROR α expression in T_{regs} was important for suppressing T cell–dependent allergic skin inflammation driven by topical application of the antigen OVA to tape-stripped skin, a T cell–dependent mouse model of AD. This was evidenced by increased epidermal thickness, increased dermal infiltration by CD45⁺ inflammatory cells (including eosinophils, mast cells, neutrophils, T cells, and ILC2s), increased cutaneous expression of *Il4*, and increased expression of IL-5 by T cells and ILCs.

We demonstrate significantly higher expression of *RORA* in human skin T_{regs} than in blood T_{regs}, suggesting that our results may be applicable to humans. Our results may be particularly relevant to patients with AD, a disease in which both T_{H2} cells and ILC2s play important roles in allergic skin inflammation. *RORA* polymorphisms in asthma (41) and *Rora* down-regulation in dogs with AD (42) further suggest that ROR α may play a regulatory role in atopic diseases. Moreover, expression of *Rora* in T_{regs} resident in tissues such as the gut (43) may endow them with the ability to dampen allergic inflammation in organs other than skin.

MATERIALS AND METHODS

Mice

Foxp3^{eyfp-cre} (C57BL/6), R26R (C57BL/6), *Rag1*^{−/−} (C57BL/6), and *Rorc^{gfp}* (C57BL/6) mice were purchased from the Jackson Laboratory (Bar Harbor, ME). *Rora^{fl/fl}* (C57BL/6) mice were generated in the laboratory of P. Chambon (France) (44). *Rora^{cre}* (C57BL/6) mice were generated in the laboratory of D. O’Leary (45). *Tnfrsf25*^{−/−} mice were generated by E. Y. Wang and obtained from the laboratory of R. Siegel. *Foxp3^{egfp}* reporter mice were a gift from T. Chatila. All mice were kept in a pathogen-free environment. All procedures performed on the mice were in accordance with the Animal Care and Use Committee of the Children’s Hospital Boston.

Preparation of skin cell homogenates from mice and human skin

Dorsal and ventral ear murine skin was separated using tweezers, chopped, and digested in complete Dulbecco’s modified Eagle’s medium containing Liberase TL (2.5 mg/ml, Roche, Life Technologies) and deoxyribonuclease (DNase) I (20 ng/ml, Sigma) for 90 min at 37°C, with vigorous shaking. Digested tissue was mechanically disrupted with a plunger, filtered, washed, and suspended in media for flow cytometric analysis. Human skin surgical discards of facial skin were obtained from the laboratory of R. Clark (Brigham and Women’s Hospital). To obtain cells from human skin, we removed all the fat using a scalpel and chopped the skin into small pieces and digested for 2 hours at 37°C with vigorous shaking in complete RPMI containing collagenase IV (2 ng/ml, Worthington Pharmaceuticals), hyaluronidase (2 ng/ml, Sigma), and DNase I. Digested tissue was mechanically disrupted using a plunger, filtered, centrifuged, and resuspended for cell sorting.

Flow cytometry

All antibodies were obtained from eBioscience and BioLegend, except anti-mouse Siglec-F, which was purchased from BD Biosciences. Cells were preincubated with Fc γ receptor–specific blocking monoclonal

antibody (2.4G2) and washed before staining. Staining with CD45 and fixable viability dye (eBioscience) was used for FACS analysis of skin cell homogenates. One hundred twenty-three count beads from eBioscience were used for estimating cell counts. Cells were analyzed on LSRFortessa (BD Biosciences), and the data were analyzed with FlowJo software (v9.7).

Intracellular staining analysis for cytokines and transcription factors

LN and skin cell suspensions were incubated with media containing phorbol 12,13-dibutyrate, ionomycin, GolgiPlug, and GolgiStop for 3 hours. Staining for surface markers was performed, followed by fixation and permeabilization using BD Cytofix/Cytoperm buffer. Cells were incubated with antibodies against cytokines, IL-4, IL-5, and IL-13, along with antibodies to FOXP3, overnight in Perm/Wash buffer (BD Biosciences). This protocol was also used to stain cells with anti-FOXP3 and anti-HELIOS markers without quenching the emission of YFP in *Rora*^{cre} R26R mice.

MC903 treatment

MC903 (catalog no. 2700) was purchased from Tocris Biochemicals. The stock was reconstituted in ethanol. MC903 (2 nM) (in a volume of 20 µl) was topically applied on the ears of mice every other day, for a total of four applications. Ethanol (vehicle) was applied on the control ear. Mice were sacrificed 1 day after the last application.

RNA preparation and qPCR

Cells were sorted directly into the lysis buffer of RNA Isolation Micro kit (Zymo Research), and RNA was prepared on the basis of kit instructions. For analysis of transcripts in skin, skin tissue was stored in RNAlater (Ambion) and homogenized using a tissue homogenizer, and RNA was prepared using RNA isolation kits (Zymo Research). Reverse transcription was performed with an iScript cDNA synthesis kit (Bio-Rad Laboratories). PCRs were run on an ABI Prism 7300 (Applied Biosystems) sequence detection system platform. TaqMan primers and probes were obtained from Life Technologies. The house-keeping gene β_2 -microglobulin was used as a control. Relative mRNA expression was quantified using the $2^{-\Delta\Delta Ct}$ method.

RNA-seq and transcriptomic analysis

CD3⁺CD4⁺Foxp3⁺(YFP⁺) T_{regs} from skin and dLNs were sorted on Aria cell sorter into the lysis buffer (PicoPure RNA Isolation kit, Life Technologies). RNA was prepared after DNase treatment (Qiagen) and sent to Dana-Farber Cancer Institute Molecular Biology Core Facility for library preparation and sequencing. Replicates with a minimum RIN (RNA integrity number) score of 7 were processed. Complementary DNA (cDNA) was synthesized using Clontech SMART-Seq v4 reagents from 500 pg of RNA and fragmented to a mean size of 150 base pairs (bp) with a Covaris M220 ultrasonicator. Illumina libraries were prepared from cDNA using Rubicon Genomics ThruPLEX DNA-seq reagents according to the manufacturer's protocol. The finished double-stranded DNA libraries were quantified and sequenced on a single Illumina NextSeq 500 sequencing system run with single-end 75-bp reads by the Dana-Farber Cancer Institute Molecular Biology Core Facility. TopHat was used to align reads to mouse genome [Mm9, National Center for Biotechnology Information (NCBI)], and HTSeq was used to estimate read counts. Read counts from all experiments are listed in table S1. Highly correlated triplicate samples were used for comparative analysis (fig. S6). DESeq2 was used to normalize data and

access differential gene expression with an FDR of <0.05. Expression levels for individual genes are represented as reads per kilobase of transcript per million mapped reads (RPKM). Heat maps were generated using GENE-E software (Broad Institute). RNA-seq raw data can be accessed through accession no. GSE99086.

Intravital two-photon imaging

Foxp3^{egfp} (Balb/c) mice were anesthetized intraperitoneally using ketamine (100 mg/kg) and xylazine (10 mg/kg). One of the ears was gently attached to an aluminum block using double-sided tape. Ear temperature was maintained at 33°C using a heating pad. GenTeal (Novartis) eye gel was spread over the ear to allow immersion of the 20× objective (0.95 numerical aperture). Images were acquired using an upright microscope (Prairie Technologies) coupled to a Mai Tai Ti:Sapphire laser (Spectra-Physics). To visualize vasculature, mice were intravenously injected with Qdot655 (Molecular Probes) diluted in phosphate-buffered saline. Images were acquired with a laser wavelength of 900 nm for optimal GFP excitation and second-harmonic generation. Epidermis and dermis were analyzed by acquisition of ~100-mm optical stacks every 30 to 60 s for 15 to 60 min with 4-mm spacing. Images were transformed into four-dimensional time-lapse movies and analyzed using Imaris software versions 7.4.2 and 8.4.1 (Bitplane). Imaging experiments were performed in the Balb/c background, but similar results were observed using *Foxp3*^{egfp} (C57BL/6) mice. Balb/c mice were preferred to avoid autofluorescence from melanin.

Histology

Tissue samples were stored in 10% formalin and sent to the histology core at Boston Children's Hospital for processing and hematoxylin and eosin (H&E) staining. Slides were analyzed on the 20× objective of bright-field microscope (Nikon), and captured images were analyzed using ImageJ software for enumeration of dermal thickness.

Local treatments by intradermal injection

Recombinant TL1A (0.9 µg/µl; catalog no. 753008, BioLegend) was injected intradermally into the ear of mice in a total volume of 10 µl every day for 3 days. Isotype antibody or anti-human/mouse TL1A antibody (R&D Systems) was injected intradermally into the ears in a total volume of 10 µl every other day for 3 days. Cells from ears were prepared, and flow cytometric analysis was performed as described earlier.

Epicutaneous sensitization

Six- to 8-week-old female mice were epicutaneously sensitized for 2 weeks, as described previously (9). In brief, for each treatment, female mice were anesthetized, and then their back skin was shaved and tape-stripped with a film dressing (Tegaderm, 3M). EC sensitization consisted of applying a 1-cm² gauze containing 200 µg of OVA (Sigma-Aldrich) to the skin after each tape stripping and securing it with a film dressing. Analyses were done at day 12.

Enzyme-linked immunosorbent assays

For detection of total IgE levels, mouse sera were prepared and enzyme-linked immunosorbent assay (ELISA) was performed (88-50460-88, eBioscience) as per the manufacturer's instructions. For quantification of cytokines in the tissue, mouse ears were flash-frozen in liquid nitrogen. Tissue was chopped, lysed, and homogenized in 500 µl of T-PER tissue protein extraction buffer (catalog no. 78510, Thermo Fisher

Scientific) in the presence of complete protease inhibitor and phosphatase inhibitors. Total protein was quantified using a bicinchoninic acid protein assay kit (catalog no. 23227, Pierce), and levels of IL-5 were enumerated after normalizing to the total protein content in the tissue. IL-5 levels in ear skin were measured using Quantikine IL-5 kit (M5000, R&D Systems), and TL1A levels were measured using DuoSet ELISA kit (DY1896-05, R&D Systems).

Statistical analysis

Statistical significance was determined by the Mann-Whitney test or analysis of variance (ANOVA) analysis using GraphPad Prism. $P < 0.05$ was considered statistically significant.

SUPPLEMENTARY MATERIALS

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Fig. S1. ROR α -expressing skin T_{regs} are HELIOS⁺ natural T_{regs} that express high levels of ST2 and ICOS.

Fig. S2. Multiple skin-resident cell types express ROR α .

Fig. S3. CD4⁺ T cells are the only cells that express eGFP in *Foxp3^{egfp}* mice.

Fig. S4. T_{reg}-specific ROR α deficiency does not affect T_{reg} numbers in the skin, nor their ability to produce IL-10.

Fig. S5. Skin TSLP and serum IgE levels in *Foxp3^{eyfp-cre}Rora^{fl/fl}* mice and *Foxp3^{eyfp-cre}* controls.

Fig. S6. Effect of lack of ROR α in T_{regs} on MC903-driven blood eosinophilia and on IL-4, IL-13, and chemokines in MC903-treated skin.

Fig. S7. Analysis of correlation of RNA-seq samples.

Fig. S8. Increased number and motility of T_{regs} in MC903-treated ear skin.

Table S1. Summary of RNA-seq experiments.

Movie S1. Intravital two-photon imaging of the untreated ear dermis of a *Foxp3^{egfp}* animal.

Movie S2. Intravital two-photon imaging of the MC903-treated ear dermis of *Foxp3^{egfp}* mice.

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ROR α -expressing T regulatory cells restrain allergic skin inflammation

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Curbing ILC2 enthusiasm

Atopic dermatitis is an allergic disease driven by type 2 immune responses in the skin. Using mouse models of dermatitis, Malhotra *et al.* have identified tumor necrosis factor (TNF) family cytokine, TNF ligand-related molecule 1 (TL1A), and its receptor death receptor 3 (DR3) as being critical in regulating cross-talk between skin-resident T regulatory cells (T_{regs}) and type 2 innate lymphoid cells (ILC2) that drive skin inflammation. They report that retinoid-related orphan receptor α (ROR α) drives expression of DR3 in T_{regs} and that, upon deletion of ROR α , skin-resident T_{regs} are unable to sequester TL1A that drives effector functions of ILC2s. The studies open up the possibility of targeting the TL1A-DR3 in the context of dermatitis and other skin allergies.

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