Essential functions of Runx/Cbfβ in gut conventional dendritic cells for priming RorγT+ T cells

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Acquired immune responses are initiated by activation of CD4+ helper T (Th) cells via recognition of antigens presented by conventional dendritic cells (cDCs). DCs instruct Th-cell polarization program into specific effector Th subset, which will dictate the type of immune responses. Hence, it is important to unravel how differentiation and/or activation of DC are linked with Th-cell–intrinsic mechanism that directs differentiation toward a specific effector Th subset. Here, we show that loss of Runx/Cbfβ transcription factors complexes during DC development leads to loss of CD103+CD11b+ cDC2s and alters characteristics of CD103−CD11b+ cDCs in the intestine, which was accompanied with impaired differentiation of RorγT+ Th17 cells and type 3 RorγT regulatory T cells. We also show that a Runx-binding enhancer in the Rorc gene is essential for T cells to integrate cDC-derived signals to induce RorγT expression. These findings reveal that Runx/Cbfβ complexes play crucial and complementary roles in cDCs and Th cells to shape converging type 3 immune responses.

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Introduction

Conventional dendritic cells (cDCs) are specialized antigen-presenting cells of the immune system. DCs in the intestine lamina propria (ILP) sense diverse antigens and migrates to draining lymph nodes where they instruct CD4+ T helper (Th) cells to differentiate into several types of effector Th cells, such as Rorγ+ Th17 and Foxp3+ peripherally induced regulatory T (iTreg) cells (Durai & Murphy, 2016; Honda & Litman, 2016). Gut cDCs are composed of two main subsets named cDC1 and cDC2 (Guilliams et al, 2014), with specialized polarizing Th functions. Gut CD103+ DCs were initially reported to induce Foxp3+ Treg cells (Coombes et al, 2007; Sun et al, 2007). However, gut CD103− DCs are now subdivided into CD103−CD11b+ cDC2 and CD103+CD11b+ cDC1. Although the functions of CD103+CD11b− cDC2 are not fully understood, previous studies have suggested that CD103+CD11b− cDC2 have the capacity to induce both Th17 cells (Lewis et al, 2011; Persson et al, 2013; Schlitzer et al, 2013) and iTreg cells (Bain et al, 2017). On the other hand, Foxp3+ iTreg cells can be divided into RorγT+Foxp3+ iTreg and RorγT+Foxp3− Treg, the latter is designated as type 3 Treg (Park & Eberl, 2018). Although the exact roles of RorγT+ type 3 Treg cells have not yet been unraveled, they are involved in suppressing exaggerated Th2 responses (Ohnmacht et al, 2015), Th17 and Th1 responses (Sefik et al, 2015). However, it remains elusive which cDC subset(s) regulates the differentiation of RorγT− Th17 and RorγT+Foxp3− Treg cells and how T cells integrate signals from cDCs to activate Rorc gene to induce RorγT expression.

Runx transcription factor family proteins function as heterodimers with Cbfβ and regulate many types of hematopoietic cells (de Bruijn & Speck, 2004; Ebihara et al, 2017). Among three mammal Runx proteins Runx1, Runx2, and Runx3, loss of Runx in hematopoietic cells leads to spontaneous development of colitis (Brenner et al, 2004) and airway inflammation in part by altering DCs function (Fainaru et al, 2004). In this study, we show that Runx/Cbfβ functions in DCs are essential not only for the differentiation of intestinal CD103+CD11b+ cDC2 but also for the priming of RorγT-expressing T cells to maintain gut homeostasis.

Results

Runx/Cbfβ complexes are essential for the differentiation of gut CD103+CD11b+ cDC2s

Runx/Cbfβ complexes regulate differentiation of Langerhans cells, epidermal–specific antigen–presenting cells, at least by transmitting TGFβ receptor signaling (Tenno et al, 2017). During DC differentiation in the gut, TGFβ receptor signaling was shown to be
essential for the differentiation of CD103−CD11b+ cDC2s (Bain et al., 2017). We thereby addressed the roles of Runx/Cbfβ complexes by inactivating the Cbfβ gene during DC development using a CD11c-Cre transgene (CbfβF/F; CD11c-Cre mice). We defined gut DCs as CD45+CD64−CD11c+MHC-II+ cells and examined CD103 and CD11b expression. Although the differentiation of CD103+CD11b− cDC1s was not affected by loss of Cbfβ, percentage and absolute cell numbers of CD103+CD11b− cDC1s were dramatically decreased in the small intestine, which was accompanied with increased relative numbers of CD103+CD11b+ cDC2s (Fig 1A). In the mesenteric lymph nodes, migratory gut DCs were defined as CD45+MHC-II+CD11clow cells. As we observed in the small intestine, CD103+CD11b− cDC2s in the migratory DC fraction were decreased in both relative and absolute cell numbers upon loss of Cbfβ (Fig 1B). CD103+CD11b− cDC2s also tended to be decreased also in the large intestine of CbfβF/F; CD11c-Cre mice (Fig S1A).

CD101 is another marker that is predominantly expressed on CD103+CD11b+ cDC2s (Bain et al., 2017), and we confirmed this observation (Fig S2A). In the small intestine of CbfβF/F; CD11c-Cre mice, CD101+CD11b+ cDCs were also decreased and remaining CD101+CD11b− cDCs were CD103+CD11b− cells, whereas most CD101+CD11b+ cells in the control mice expressed CD103 (Fig S2A). Along with no accumulation of CD101+ cells in CD103+CD11b− population in CbfβF/F; CD11c-Cre mice (Fig S2A), we concluded that the development of CD103+CD11b− cDC2s was severely impaired in the absence of Cbfβ.

We also examined cDCs in another barrier tissue, the lung. In lung cDC population, our analyses of CD103 and CD11b expression detected two major cDC subsets, CD103+CD11b− and CD103+CD11b+, with a very minor population of CD103+CD11b− cells, which is consistent with a previous report (Greter et al., 2012) proposing that CD103+CD11b− subset is a major cDC2 subset in the lung. Interestingly, absolute cell numbers of CD103+CD11b− cDC2 subset were significantly reduced in the lung of CbfβF/F; CD11c-Cre mice (Fig S2B). In addition, CD101 expression was nearly undetected in both CD103+CD11b− and CD103+CD11b+ subsets upon loss of Cbfβ, indicating that Cbfβ deficiency also resulted in the impaired differentiation of CD101+expressing CD11b+ cDC2 subset in the lung.

All three mammal Runx genes, Runx1, Runx2, and Runx3, are expressed in all cDC subsets in the small intestine (Fig S3A). Using an anti-Runx3 antibody that detects Runx3 protein in flow cytometry analyses, we found that CD103+CD11b− and CD103+CD11b+ gut DC subsets expressed Runx3 at higher level than that in CD103+CD11b− cDCs (Fig S3B). To examine which Runx protein is responsible for the development of CD103+CD11b− cDC2s, we inactivated Runx1 and Runx3 gene by CD11c-Cre individually or in a combinational manner. Although ablation of Runx1 or Runx3 alone caused a slight decrease in CD103+CD11b− cDC2s, these cells were nearly absent in the small intestine and mesenteric lymph nodes upon loss of both Runx1 and Runx3 proteins (Fig 2). Decrease in CD103+CD11b− cDC2s was also observed in the large intestine by combination deletion of both Runx1 and Runx3 (Fig S3C), as was...
observed in CbfbF/F: CD11c-Cre mice. These observations indicate not only the redundant function between Runx1 and Runx3 proteins but also the essential requirement of both proteins to direct the differentiation of CD103+CD11b+ cDC2s.

**Impaired Rorγt+ T-cell differentiation by lack of Runx/Cbfβ in DCs**

One of the important functions of gut DCs is to prime CD4+ Th cells to differentiate into effector Th subsets such as FoxP3+ iTreg and Rorγt+ Th17 cells (Durai & Murphy, 2016). Because CD103+CD11b+cDC2s were shown to support differentiation of both iTreg and Th17 cells (Lewis et al, 2011; Persson et al, 2013; Bain et al, 2017), we examined the expression of FoxP3 and Rorγt in the lamina propria CD4+ T-cell population. In CbfbF/F: CD11c-Cre mice, the relative numbers of both FoxP3+Rorγt+ and FoxP3−Rorγt+ cells were significantly decreased in the small intestine (Fig 3A), whereas the reduction of such Rorγt+ T cells was not obvious in the large intestine (data not shown). Inactivation of both Runx1 and Runx3 genes in DCs also led to the decrease in small intestine Rorγt+ T cells (Fig S3D). These observations suggest an essential role of Runx/Cbfβ in DCs to support the differentiation of Rorγt+ T cells.

Functions of Runx/Cbfβ in T cells were shown to be important for the development of Treg (Kitoh et al, 2009; Rudra et al, 2009) and Th17 cells (Zhang et al, 2008). Previous report detected a leaky expression of the CD11c-Cre transgene in T lymphocytes (Bain et al, 2017). Similarly, we detected CD11c-Cre-mediated recombination of Cbfb gene to some extents in gut CD4+ T cells (Fig S1B). To examine cell intrinsic effects of Cbfb inactivation on the differentiation of gut Th17 and Treg cells, we inactivated Cbfb gene in T cells using CD4-Cre transgene (CbfbF/F:CD4-Cre mice). As previously reported (Kitoh et al, 2009; Rudra et al, 2009), FoxP3 expression level was slightly decreased upon inactivation of Cbfb in T cells by CD4-Cre transgene, whereas the relative numbers of FoxP3−Rorγt+ Th17 cells were not affected (Fig 3B). Instead, loss of Cbfβ in T cells resulted in an increase in FoxP3−Gata3+ Th2 cells in the small intestine (Fig 3B), presumably in part due to a failure of Il-4 silencing (Naoe et al, 2007) and loss of Runx3-mediated antagonism against Gata3 function (Yagi et al, 2010). We also examined the effect of combinational loss of Cbfβ in T cells with Cbfβ deficiency in DCs. In the gut of CbfbF/F: CD11c-Cre: CD4-Cre mice, reduction of Rorγt+ cells, in particular FoxP3+Rorγt+ cells, tended to be enhanced, indicating that the loss of Runx/Cbfβ in both T cells and DC synergistically impacts the differentiation of Rorγt+ T cells.

We previously showed that Rorγt expression in group 3 innate lymphoid cells (ILC3) and lymphoid tissue inducer (LTI) cells is lost by...
removal of a single intronic enhancer (E-11), which locates 11 kb downstream from the Rorgt transcriptional start site and is occupied by Runx/Cbfb complexes in thymocytes (Tenno et al., 2018). To further examine the Runx-mediated T-cell–intrinsic mechanism for Rorγt induction, we first addressed a role of the E-11 for Rorγt expression in T cells using the Rorcγf/gfp reporter allele in which the E-11 enhancer was removed from the Rorcγf allele. In the small ILP of Rorcγf/gfpΔE-11 mice, Rorγt T cells were developed. However, GFP expression from the Rorcγf/gfpΔE-11 reporter allele was not detected (Fig S4A). Similarly, Rorγt cells were absent from the small ILP of RorcΔE-11 mice lacking the E-11 enhancer in the Rorc locus. In addition, Rorγt-expressing cells in non–T-cell population, which correspond to ILC3 and LTi, were lost in RorcΔE-11 mice (Fig S4B), whereas the level of Rorγt in DP thymocytes was just reduced to the half of the one detected in control cells. In addition, naive CD4+ T cells from RorcΔE-11 mice failed to differentiate into IL-17–producing Th17 cells in Th17-inducing culture condition (Fig S4B). These results demonstrated that the E-11 enhancer is crucial for Rorγt expression during in vivo and in vitro differentiation to Th17, whereas other regulatory regions partially compensate Rorγt induction in DP thymocytes.

**Impaired signatures of CD103+CD11b+ cDC2 in the absence of Cbfb**

Our results showed that the loss of Runx/Cbfb during cDC development resulted in loss of CD103+CD11b+ cDC2s and a severe reduction in Rorγt-expressing T cells in the small intestine. A specific loss of CD103+CD11b+ cDC2s was also observed in other mutant mouse strains, such as the Notch2f/f: CD11c-Cre and Irf4f/f: CD11c-Cre, and Tgfbr2f/f: CD11c-Cre mice (Lewis et al., 2011; Persson et al., 2013; Satpathy et al., 2013; Schlitzer et al., 2013; Bain et al., 2017). However, the degree of Th17 cells reduction reported in those mice seems to be less than that in Cbfbf/f: CD11c-Cre mice. Indeed, a significant proportion of Rorγt+ T cells remained in Notch2f/f: CD11c-Cre mice (Fig 4A), although the relative numbers of Rorγt+ T cells tended to be reduced when compared with that of control mice. Thus, the relative numbers of Rorγt+ T cells were more than twofolds lower in Cbfbf/f: CD11c-Cre mice than in Notch2f/f: CD11c-Cre mice (Figs 3A and 4A), whereas the degree of CD103+CD11b+ cDC2s reduction was comparable between these two strains. Interestingly, the percentages of CD103+ cells in CD101+CD11b+ cDC2s of Notch2f/f: CD11c-Cre mice were lower than that of control mice but were higher than that of Cbfbf/f: CD11c-Cre (Fig 4B). These observations suggested that the impaired survival of CD103+CD11b+ cDC2s might occur by loss of Notch2, whereas developmental inhibition is likely to be the main mechanism for reduction of CD103+CD11b+ cDC2s in Cbfbf/f: CD11c-Cre mice, as was observed in the case of the loss of TGFβ receptor signaling (Bain et al., 2017).

It is conceivable that the severer loss of Rorγt+ T cells in Cbfbf/f: CD11c-Cre is not caused simply by the loss of CD103+CD11b+ cDC2s, and that the functions of other DCs supporting Rorγt+ T-cell differentiation might also be impaired by the loss of Cbfb. We, therefore, examined the gene expression profiles of Cbfb-deficient CD103+CD11b+ and CD103+CD11b+ DCs by RNA-seq. Principle component analyses (PCA) indicated that CD103+CD11b+ DCs became more different from control cells in PC2 values upon loss of Cbfb (Fig 5A). In CD103+CD11b+ DCs, more genes were up-regulated upon loss of Cbfb, such as IL-18, than down-regulated in their expression levels (Fig 5B and C). Interestingly, IL-18 was shown to limit colonic
Th17 cell differentiation (Harrison et al, 2015). On the contrary, in CD103+CD11b− DCs, more genes were down-regulated upon loss of Cbfβ, than up-regulated (Fig 5A). Recently, IL22ra2, one of the down-regulated genes in both CD103+CD11b− and CD103−CD11b+ DCs, was shown to be decreased by IL-18 and be involved in the control of intestinal tissue damage (Huber et al, 2012).

Spontaneous colitis development in CbfbF/F: CD11c-Cre mice

Our results indicate that the loss of Cbfβ during cDC development led to significant changes in cDC and Th-cell composition in the intestine. It was reported that Runx loss in hematopoietic cells (Brenner et al, 2004) and loss of CD103−CD11b+ cDC2s by lack of TGFβ receptor signaling (Bain et al, 2017) resulted in spontaneous colitis development. Interestingly, CbfbF/F: CD11c-Cre mice tended to develop colitis spontaneously around 6 mo and later, although significant histological changes were not observed at 7 wk old when differentiations of gut DCs and Th subsets were analyzed (Figs 6A and S5). Despite different cellular compositions in the intestine and colitis development in CbfbF/F: CD11c-Cre mice, the composition of the commensal gut microbiota assessed by feces of 5-mo-old CbfbF/F: CD11c-Cre mice were not significantly different from that in control mice (Fig S6), except for an increase in segmented filamentous bacteria (SFB) (Fig 6B), a known inducer of Th17 differentiation (Ivanov et al, 2009). These observations confirmed that the decreased Th17 differentiation that we observed is not caused by lack of SFB in microbiota.

Discussion

In this work, we provide concrete genetic evidence supporting a crucial function of Runx/Cbfβ complexes in gut cDC development, not only for the differentiation of CD103−CD11b+ cDC2s but also for their polarizing activity that primes both Rorγt+ Th17 and type 3 Rorγt FoxP3+ Treg cells. Several mutant mouse lines, in which CD103+CD11b+ cDC2s in the gut was diminished or lost, have been reported. However, roles of gut cDCs in priming gut Th-cell subset still remains elusive. For example, loss or reduction of CD103−CD11b+ cDC2s in intestinal lamina upon inactivation of Notch2 (Lewis et al, 2011; Satpathy et al, 2013) or Irf4 genes (Schlitzer et al, 2013) or by the expression of the Langerin-diphtheria toxin A transgene (Welty et al, 2013) was shown to result in a decrease in Th17 cells without significant reduction in Foxp3+ Treg cells. In contrast, ablation of TGFβR during DC development led to the reduction in CD103−CD11b+ cDC2s, which was accompanied with reductions of both Th17 and Foxp3+ Treg cells (Bain et al, 2017). It is noteworthy that gut Rorgt+ CD4+ T cells consists of at least two types of cells, Th17 and type 3 Treg cells. Although staining of both FoxP3 and Rorγt proteins is necessary to detect these two subsets, previous studies did not always examine these two subsets, making it difficult to fairly compare the phenotype of gut effector Th cells among several mouse strains. However, reduction of Rorgt+ T cells, including Th17 and type 3 Treg, seems to be more severe in CbfbF/F: CD11c-Cre mice than in Tgfbr1F/F: CD11c-Cre mice, suggesting that Runx/Cbfβ complexes may have unique roles other than...
transmitting TGFβ signals in regulating development or functions of gut cDCs. In line with this assumption, we found that CD101 expression in the gut and lung cDCs was nearly abolished in CbfbF/F: CD11c-Cre mice (Fig S2), whereas it was detected on those cells after loss of TGFβR (Bain et al, 2017). This difference suggests that Runx/Cbfb complexes might regulate DC development during earlier stages rather than later stages, which requires TGFβ signaling. CD101 expression on T lymphocytes and myeloid cells was reported to be important for preventing intestinal inflammation (Schey et al, 2016). Although relevance of CD101 expression in regulating cDCs remains unclear, it is possible that induction of CD101 expression during cDC differentiation is important for their polarizing functions of effector Th-cell differentiation.

It remains unclear whether a single DCs subset plays a predominant role in inducing both Th17 and type 3 Treg cells or distinct subpopulations of these CD103+CD11b+ cDC subsets prime each T-cell subset independently. Compared with other mutant strains lacking CD103+CD11b+ cDC subsets, the reduction in Rorγt-expressing T cells is prominent upon the loss of Cbfb. Such clear difference may suggest a presence of another DC population that compensate the
loss of CD103\(^{−}\)CD11b\(^{+}\) cDC2 subset for inducing Ror\(\gamma\) T-cell differentiation. Previously, CD103\(^{−}\)CD11b\(^{+}\) DCs was shown to have capacity to prime Th17 differentiation (Liang et al., 2016). Residual CD103\(^{−}\)CD11b\(^{+}\) cDCs in Cbf\(\beta\)\(^{−}\)CD11c-Cre mice showed impaired gene signatures including up-regulation of the IL-18 gene. Given a reported antagonistic function of IL-18 against Th17 differentiation (Harrison et al., 2015), ectopic IL-18 secretion from Cbf\(\beta\)-deficient CD103\(^{−}\)CD11b\(^{+}\) cDCs might alter the microenvironment toward inhibiting Th17 differentiation and intestinal inflammation. Consistent with a previous report showing a role of IL-18 in decreasing IL22ra2 level (Huber et al., 2012), IL22ra2 was down-regulated in both CD103\(^{−}\)CD11b\(^{+}\) and CD103\(^{−}\)CD11b\(^{+}\) cDC subsets. Thus, it is possible that dysfunction of remaining cDCs by loss of Runx/Cbf\(\beta\) have direct or indirect roles in regulating Ror\(\gamma\) T-cell differentiation. Either case, Runx/Cbf\(\beta\) complexes are indispensable for either differentiation or function of such cDC subset(s).

In addition to DC function, we showed that the single intronic enhancer, E-11, in the Rorc locus is crucial to integrate signals from cDCs to induce Ror\(\gamma\)T expression in T cells. The E-11 enhancer contains two conserved Runx-binding sequences and is occupied by Runx/Cbf\(\beta\) in thymocytes (Tenno et al., 2018). Ror\(\gamma\) T cells are still differentiated upon loss of Cbf\(\beta\) in T cells (Fig 3B), suggesting that other transcription factors compensate Runx/Cbf\(\beta\) function for activating the E-11 enhancer. However, once DC-derived priming signals are diminished by loss of Cbf\(\beta\) in cDCs, the effect of Cbf\(\beta\) deficiency in T cells would be beyond such compensatory pathway. On the contrary, loss of IL3C and LTI cells by attenuation of Runx/Cbf\(\beta\) activity (Tachibana et al., 2011; Ebihara et al., 2015; Tenno et al., 2018) indicates that activation of the E-11 enhancer in ILCs is more dependent on Runx/Cbf\(\beta\) complexes.

Collectively, our results demonstrate that a single transcription factor, Runx/Cbf\(\beta\), plays an essential role in shaping type 3 immune responses by regulating the developmental program of distinct immune cells, cDCs and T cells. It is unclear whether DCs and T cells coincidentally use the same transcription factor to enhance type 3 responses, or, alternatively, whether there is uncharacterized crosstalk between cDCs and T cells that modulates the type 3 developmental program in each cell type. Evolutional pressures to efficiently mount type 3 immune responses may have led to convergence of each program, leading to the use of a common transcription factor, Runx/Cbf\(\beta\). Along with an antagonism against type 2 responses in part through suppressing IL-4 production (Djuretic et al., 2007; Naoe et al., 2007), we show here that Runx/Cbf\(\beta\) complexes serve as central players for balancing type 2 and type 3 immune responses.

**Materials and Methods**

**Mice**

Cbf\(\beta\)\(^{floox}\) (Naoe et al., 2007), Runx\(\beta\)\(^{floox}\) (Taniuchi et al., 2002), Runx\(\beta\)\(^{floox}\) (Naoe et al., 2007), Rorc\(^{eo−E−}\)\(^{−}\) and Rorc\(^{E−}\)\(^{−}\) mice (Tenno et al., 2018) were previously described. Rorc\(^{eo−}\) reporter mice (Eberl et al., 2004), Cd11c-Cre (Caton et al., 2007) mice, and Cd4-Cre (Lee et al., 2001) mice were obtained from Dr DR Littman, Dr Boris Reis, and Dr CB Wilson, respectively. Notch2\(^{floox}\) mice were purchased from Jackson laboratory (010525). Cbf\(\beta\)\(^{floox}\) strains were maintained in the animal facility at the RIKEN Center for Integrative Medical Sciences. Runx\(\beta\)\(^{floox}\), Runx\(\beta\)\(^{floox}\), and Notch2\(^{floox}\) strains were maintained in the animal facility at the SigN. All animal experiments were performed in accordance with the protocol (28-017) approved by Institutional Animal Care and Use Committees of RIKEN Yokohama Branch and those approved by Institutional Animal Care and Use Committees of the Biological Resource Center (Agency for Science, Technology and Research, Singapore) and with the guidelines of animal care in RIKEN, Center for Integrative Medical Sciences, and the guidelines of the Agri-Food and Veterinary Authority (AVA) and the National Advisory Committee for Laboratory Animal Research of Singapore.

**Cell preparation**

Single-cell suspensions from mouse intestine were prepared using a previously described method (Bogunovic et al., 2009) with some...
In vitro Th17 differentiation

Naïve CD4⁺ T cells were purified from spleens by depleting CD4⁻ negative cells, followed by enrichment of CD62L⁺ cells using Naive CD4⁺ T cell Isolation Kit (Miltenyi Biotec) according to the manufacturer's protocol. 1.0 × 10⁵ cells were primed with plate-bound 2 μg/ml anti-CD3 (145-2C11; BD Pharmingen) and 2 μg/ml anti-CD28 (37.51; BD Pharmingen) in 100 μl DMEM supplemented with 10% FCS in round-bottomed 96-well plates at 37°C for 3 d in the presence of 10 μg/ml anti-IL-4 and anti-IFN-γ neutralizing antibodies (Biolegend), 50 ng/ml Mil-6 (R&D) and 1 ng/ml mouse Transforming Growth Factor-β (R&D). At day 3, 100 μl of fresh medium with the above reagents were added. At day 5, the cells were stimulated with PMA/ionomycin and fixed and permeabilized with the Fixation/Permeabilization Solution kit (BD Biosciences) and were stained for IL-17 (TC11-18H10.1, BioLegend) according to the manufacturer’s protocol.

RNA-seq analysis

Expression level of Runx genes in DCs subset in wild-type mice were analyzed using GSE100393 data. To compare gene expression profiles of wild-type and Cbfb⁻ deficient DCs subset, RNA-seq libraries were prepared by applying a RamDA-seq method for single cell RNA-seq (Hayashi et al, 2018) with some modifications. Briefly, 218-1000 CD103⁺CD11b⁺ and CD103⁺CD11b⁻ cells were sorted by FACSAria directly into tubes containing buffer T (QiAGEN). RNA was isolated using Agencourt RNAclean XP beads (Beckman Coulter). Then the RNA was eluted into 10 μl of RNase-free water and used for double strand (ds)-cDNA synthesis by RamDA-seq protocol adjusted for bulk input, which used 10 times larger reaction scale than for a single cell. After quantitation of ds-cDNA using Qubit dsDNA HS Assay Kit (Thermo Fisher Scientific), 3 ng of ds-cDNA was applied for Tn5 tagmentation in a 75-μl reaction with in-house Tn5 enzyme as previously described (Hennig et al, 2018). Then the DNA was amplified by 16–19 cycles of PCR using indexed primers and KAPA HiFi DNA Polymerase (Kapa Biosystems). The libraries were purified by Agencourt AMPure XP beads (+1.2 vol; Beckman Coulter), quantified by quantitative PCR and pooled, and were sequenced on an Illumina HiSeq 2500 instrument in a high-output mode. Low-quality part of short reads and adapter sequences were trimmed by trim_galore (v0.5.0, http://www.bioinformatics.babraham.ac.uk/projects/trim_galore/) and mapped to mm10 whole genome using STAR v2.7 with a GTF file from EMBL-EBI (GRCm38.92). Unmapped reads and reads mapped with low-quality scores (MAPQ < 5, 12–20%) and duplicated reads (14–39%) were filtered out using SAMTools view -q 5 (Li et al, 2009) and Picard Tools (Picard MarkDuplicates, http://broadinstitute.github.io/picard). Reads on each gene were counted using htsq-cound (Anders et al, 2015) with -s option and a GTF file from EMBL-EBI (GRCm38.92). To avoid non-informative genes, we removed genes on which counts were less than five in any samples from downstream analysis. Differentially expressed genes (2,560 genes, FDR < 0.01) were extracted with the “edgeR” package in R programming language (Robinson et al, 2010) to perform PCA using the “prcomp” function in R. To examine the most affected genes by Cbfb deficiency in CD103⁺CD11b⁻ DCs and CD103⁺CD11b⁺ DCs, genes were further selected by fold change (fold change > 5, control versus Cbfb⁻ deficient cells) and were plotted as heat map after z-score normalizations. For Gene Ontology enrichment analysis, 258 genes up-regulated in Cbfb⁻ deficient CD103⁺CD11b⁻ DCs (fold change > 1.3 and FDR < 0.05, control versus Cbfb⁻ deficient cells) were investigated using the “clusterProfiler” package in R (v3.8.1) (Yu et al, 2012).

Histology

Colonial tissues were processed as Swiss rolls (Bialkowska et al, 2016), fixed in Mildform 10N (Wako) overnight, embedded in paraffin, and sectioned at 5-μm thickness. Histological analysis was performed after staining with hematoxylin and eosin, and the severity of colitis was scored based on the criteria (Table S1) that was previously described (Asseman et al, 1999) in a blinded fashion.

Gut microbiotas analyses

Feces were collected from 20 to 30-wk-old mice. Bacterial DNA for 16S rRNA gene sequencing was extracted as previously described (Kato et al, 2018). The V4 variable region (515F to 806R) of the respective samples was sequenced using Illumina MiSeq by following the method previously described (Kozich et al, 2013). Taxonomic assignments and the estimation of relative abundances from sequencing data were performed using the analysis pipeline of the QIIME software package (Caporaso et al, 2010). An operational taxonomic unit (OTU) was defined at 97% similarity. OTU taxonomy was assigned based on comparison with the Greengenes database,
using UCLUST (Quast et al, 2013). Beta diversity was calculated using the UniFrac distance metric and visualized by principal coordinate analysis (PCoA) using weighted or unweighted UniFrac distances based on the OTU distribution across samples.

Genomic PCR analyses

To examine the genotype of ILP CD4+ T cells and MHC-II+ CD11c+ DCs of Cbfb+/Δ/, CD11c-Cre mice, DNA was prepared from sorted 5000 cells and analyzed by PCR. Primers used were CbfbEx5G2 5'-CCTCCTATTCTCAA-CAGGAAT-3', CbfbEx5G3 5'-GGTTAGGAGTCATTGTGATCAC-3' and CbfbEx5G6 5'-CATTGGATTTGCGTTACTGG-3'.

Data analyses

Statistical analysis was performed by Student's unpaired t test with GraphPad Prism6 (GraphPad software) unless otherwise stated.

Data Availability

The sequencing data from this publication have been deposited to the Genome Expression Omnibus database under accession number GSE130380.

Supplementary Information

Supplementary Information is available at https://doi.org/10.26508/lsa.201900441.

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Author Contributions

M Tenno: formal analysis and investigation.
F Ginhoux: supervision, funding acquisition, and writing—original draft, review, and editing.
I Taniuchi: supervision, funding acquisition, and writing—original draft, review, and editing.

Conflict of Interest Statement

The authors declare that they have no conflict of interest.

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