Absence of Batf3 reveals a new dimension of cell state heterogeneity within conventional dendritic cells

Highlights
- Single-cell proteogenomics identifies additional layers of DC heterogeneity
- cDC diversity is driven by proliferation, developmental stage, and maturation
- Lack of Batf3 increases cDCs with lineage-mixed features
- Sox4+ cDCs represent a cell state of lineage-mixed features
Article

Absence of Batf3 reveals a new dimension of cell state heterogeneity within conventional dendritic cells

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SUMMARY

Conventional dendritic cells (cDCs) are traditionally subdivided into cDC1 and cDC2 lineages. Batf3 is a cDC1-required transcription factor, and we observed that Batf3−/− mice harbor a population of cDC1-like cells co-expressing cDC2-associated surface molecules. Using single-cell RNA sequencing with integrated cell surface protein expression (CITE-seq), we found that Batf3−/− mitotic immature cDC1-like cells showed reduced expression of cDC1 features and increased levels of cDC2 features. In wild type, we also observed a proportion of mature cDC1 cells expressing surface features characteristic to cDC2 and found that overall cDC cell state heterogeneity was mainly driven by developmental stage, proliferation, and maturity. We detected population diversity within Sirpa+ cDC2 cells, including a Cd33+ cell state expressing high levels of Sox4 and lineage-mixed features characteristic to cDC1, cDC2, pDCs, and monocytes. In conclusion, these data suggest that multiple cDC cell states can co-express lineage-overlapping features, revealing a level of previously unappreciated cDC plasticity.

INTRODUCTION

Conventional dendritic cells (cDCs) play a central role in bridging innate and adaptive immune responses due to their specialization in antigen presentation and activation of T cells. The current terminology distinguishes cDC1 and cDC2 lineages based on differential ontogeny, expression of surface features, requirement for transcription factors, and function. cDC1 cells are highly specialized in cross-presentation and activation of cytotoxic T cells, whereas cDC2 cells are primarily driving helper T responses (Wculek et al., 2020).

The development of cDCs from precursor CD34+ cells is driven by specific transcription factors shared between mouse and human (Schultze et al., 2019), and cDC subtype divergence is largely driven by interferon-regulatory factors 4 and 8 (IRF4/IRF8) (Murphy et al., 2016; Seillet and Belz, 2013). cDC1 in mice express CD8α and/or CD103, and CD141 in humans. Across species, cDC1 are classified as IRF8+ or Batf3-dependent cDCs, and cDC1 development and maintenance depends on transcription factors IRF8, ID2, and Batf3. While IRF8 and ID2 are indispensable for lineage commitment, Batf3 is required for sustained IRF8 autoactivation and subsequently for the maintenance of cDC1 (Seillet et al., 2013; Grajales-Reyes et al., 2015). Function-conveying features of cDC1 are XCR1 and CLEC9A. cDC2 in mice are CD11b+ CD172a+, and CD4+ when residing in lymphoid tissues, whereas in humans, cDC2 express CD1c. Across species, cDC2 are classified as IRF4+ or Batf3-independent, and in human and mice, transcription factors driving cDC2 commitment include IRF4, NOTCH2, Rbp-J, and KLF4. In addition, our understanding of cDC2 heterogeneity is steadily increasing.

Recent advances in high-dimensional single-cell technologies have challenged the canonical cDC classification. Several groups utilizing single-cell omics technologies described unprecedented heterogeneity within cDCs in both humans and mice. The current consensus is that cDC1 are relatively homogeneous, but cDC2 are heterogeneous in phenotype and function (Brown et al., 2019; Villani et al., 2017). Recently, cDC2 were shown to comprise functionally distinct subpopulations expressing either TBET or RORGT,
transcription factors commonly associated with T cells and innate lymphoid cells (Brown et al., 2019). Additionally, transitional dendritic cells (DCs) and DC3 have been described in both mouse and human, which share monocyte features and are efficient antigen presenters (Leylek et al., 2019). Single-cell RNA sequencing (scRNA-seq) analysis of DCs has enabled the identification of subpopulation-specific gene signatures and advanced our understanding of DC heterogeneity. Despite this, accurate cDC2 classification remains unresolved due to the reduced ability to detect expression of cDC2-characteristic transcripts such as Itgam (CD11b), Cd44, and Ifi4 (Stoeckius et al., 2017).

We previously reported that the acquisition of a cDC1 phenotype and function depends on Batf3 and that in the absence of Batf3, a residual population of CD8+ cDCs co-expresses surface molecules characteristic to both cDC1 and cDC2 lineages (Chandra et al., 2017). Consequently, we sought here to explore cell state plasticity of cDCs in steady state and in the absence of Batf3 using multiple single-cell technologies including conventional and imaging flow cytometry, and scRNA-seq with integrated cell surface protein expression. This enabled us to identify cDC cell states co-expressing mixed cDC phenotypic and transcriptional signatures. We identified MHCII+ CD11c+ cells expressing both CD11b and CD8a in steady state, which were increased in Batf3−/− mice. We demonstrate that mature classical cDC1 cells are nearly absent in Batf3−/− mice, and that a population of mitotic cDC1-like cells remained immature and likely non-functional due to diminished expression of the classical cDC1 markers Tir3, Xcr1, and Clec9a. This population further expressed increasing levels of cDC2 genes, which supports the notion by others of a possible phenotypic plasticity between different DC subtypes (Grajales-Reyes et al., 2015; Lau et al., 2018). Our data further revealed a population of Sirpa+ Sox4+ Cd33+ cDCs expressing lineage-mixed features of cDC1, cDC2, pDCs, pre-DCs, and monocytes in both wild-type (WT) and Batf3−/− mice. cDCs with lineage-mixed phenotypes have recently caught attention in multiple other studies in both human and mice (Brown et al., 2019; Leylek et al., 2019; Bosteels et al., 2020; Bourdely et al., 2020; Cytlak et al., 2020). The Sirpa+ Sox4+ Cd33+ cDCs lineage-mixed cell state we describe here lacks expression of toll-like receptor (TLR) genes and genes indicating cDC1 functions, whereas it expresses the highest levels of Lag3 among cDCs. Our data further demonstrate differential expression of co-inhibitory molecules across different cDC cell states, which may assist cDC cell state-specific target prioritization for checkpoint inhibition.

RESULTS

Identification of cDCs that express both cDC1- and cDC2-characteristic surface features

Batf3 is required for continued autoactivation of IRF8 in cDC precursors, allowing commitment of these cells to the cDC1 lineage (Grajales-Reyes et al., 2015). However, this commitment can also be achieved in the absence of Batf3 by infection-associated induction of IL-12, which promotes expression of Batf, enabling cDC1 development in Batf3−/− mice (Tussiwand et al., 2012). It is therefore not surprising that cDC1 development in Batf3−/− animals can vary with the health of the studied animals (Mott et al., 2015). Using a multiparameter flow cytometry approach, we previously established that Batf3−/− mice harbor a residual population of CD8+ cDC cells which co-express increasing levels of cDC2-characteristic surface molecules CD11b, CD172a, and CD4, but lack IRF8 (Chandra et al., 2017). However, technical artifacts in conventional flow cytometry might explain this finding. Therefore we used imaging flow cytometry to confirm the existence of cDCs that express both cDC1- and cDC2-characteristic surface features. T- and B cell-depleted splenocytes were used to initially identify CD11c+ MHCII+ cDCs (Figure S1), which were subsequently analyzed for surface expression of CD11b and CD8. Classical cDC1 and cDC2 were identified as single positive for CD8 or CD11b, respectively (Figures 1A and S1). We further identified lineage-intermediate (lin-int) cDCs co-expressing CD8 and CD11b on the cell surface in both WT and Batf3−/− mice. We compared the expression level of CD8 and CD11b between lineage-intermediate DCs, cDC1 and cDC2 and found that WT and Batf3−/− lineage-intermediate cDCs expressed significantly more CD8 when compared with CD8− cDC2, but less than CD8+ cDC1 (Figure 1B). Similarly, lineage-intermediate DCs expressed significantly more CD11b when compared with CD11b− cDC1, and less when compared with CD11b+ cDC2 (Figure 1B). These CD8+ CD11b+ cDCs comprised a higher abundance in Batf3−/− animals than in wild-type animals (Figure 1C).

Cell state plasticity of splenic cDCs is driven by ontogeny, cell cycle, and maturation

Of note, our previous data indicated that CD8+ CD11b+ cDCs demonstrated a continuum of CD8 and CD11b expression (Chandra et al., 2017) suggesting that these cells do not represent a committed cell lineage, but more likely demonstrate cell state plasticity. To uncover the transcriptional basis of this
phenomenon, we carried out microfluidic droplet-based scRNA-seq of purified splenic cDCs in combination with barcoded cDC-specific antibodies, referred to as cellular indexing of transcripts and epitopes by sequencing (CITE-seq) (Stoeckius et al., 2017). After T and B cell depletion, cDCs were sorted to purity as live singlet CD11c+ MHCII+ cells (Figure S2). We sequenced RNA from 13,943 WT and 13,750 Batf3−/− cDCs and obtained ~30,000 reads per cell, from a median number of 2,352 genes per cell. After pre-processing and quality control, the data from 12,224 WT and 12,119 Batf3−/− cells underwent log normalization and uniform manifold approximation and projection (UMAP) dimensionality reduction.

Figure 1. Absence of Batf3 leads to enrichment of lineage-intermediate cDCs
(A and B) MHCII+ CD11c+ splenic cDCs of C57BL/6 WT and Batf3−/− mice were assessed for expression of CD8 and CD11b by imaging flow cytometry. (A) Pre-gated to MHCII+ CD11c+ cells, cDC1 cells were identified as CD8+CD11b−, cDC2 cells as CD8−CD11b+, and lineage-intermediate cDCs as CD8+CD11b+ (Figure S1). BF, brightfield; merge, overlay of CD8 and CD11b. (B) Intensities of CD8 and CD11b was compared between C57BL/6 WT and Batf3−/− cDC lineages. Each data point represents an individual cell with mean and interquartile range indicated. (C) cDCs in splenocytes of C57BL/6 WT and Batf3−/− mice (n=5) were immunoprofiled using conventional flow cytometry. The number of CD8+CD11b+ cDCs per 100,000 B cells was compared. Statistical significance was determined using one-way ANOVA followed by Tukey’s multiple comparison test. *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001. Shown is one of two independent experiments. See also Figure S1.
Unbiased graph clustering using the Seurat pipeline (Butler et al., 2018) at a default resolution of 0.5 identified 20 clusters (C) of the sorted CD11c+ MHCII+ cells (Figure 2A). To determine the differences between WT and Batf3−/− cells, composition analysis by genotype was undertaken, and showed that C4, C12, C17 and C18 were almost exclusively represented by WT, whereas no Batf3−/−-specific cluster was found (Figure 2B). We analyzed differentially expressed genes (DEGs) between the WT and Batf3−/− genotypes for each cluster. The majority of clusters comprised similar numbers of cells from both genotypes and, with the exception of C4, C6, and C12, revealed only Batf3 as a DEG (Table S1). Given the overall similarity of the WT and the Batf3−/− clusters, we proceeded to annotate cluster identities from the combined dataset.

Using SingleR (Aran et al., 2019) and the integrated Immunological Genome Project (ImmGen) data (Heng et al., 2008; Miller et al., 2012) to compare the single-cell transcriptional features with immune cell expression profiles, C13 were predicted to be NK/NKT/ILC/T cells, C15 to be monocytes, C19 to be stem cells, and the remaining 17 clusters to be DCs (Figure 2C). Of note, C2 and C11 were predicted to be DCs with high confidence, but were also moderately enriched with a monocyte signature. With the exception of the C13 cluster, no gene expression of Klrb1a, Cd19, or Cd3e was detected, demonstrating the absence of NK, B and T cells within the DC clusters (Figure S3A). All DC clusters expressed MHC class II genes and Itgax (CD11c) (Figure S3B). Using the DC lineage-specific cell signatures published by the Immgen group (Miller et al., 2012) (Table S2), we tested each cluster for enrichment of signatures of core cDCs, CD8+ cDCs (cDC1), CD8− cDCs, and pDCs using the AUCell package (Aibar et al., 2017). Consistent with the SingleR analysis, C13, C15, and C19 were not enriched for a core cDC signature, whereas all other clusters aligned (Figure 2D). Cells of C14 displayed the highest pDC area under curve (AUC) score. C4, C6, C12, C17, and C18 had the highest cDC1 AUC scores. Of note, the Immgen CD8− cDC signature only partially aligned with the remaining DC clusters and was enriched in the monocyte cluster C15, indicating that this signature is not cDC2 specific. However, this was expected with respect to our current knowledge of heterogeneity within CD8− cDCs.

Expression analysis of key canonical DC and monocyte genes further supported the identity of pDCs, cDC1, cDC2, and monocytes (Figure 2E). Expression of Sirpa, traditionally associated with a cDC2 phenotype, was the most reliable identifier of a non-cDC1 cDC2 phenotype (Figure 2E). Based on these analyses, we broadly subgrouped DC clusters into cDC1, Sirpa+ DCs and pDCs (Figure 2F).

The Sirpa+ DC clusters C2, C3, and C11 additionally displayed features characteristic of monocytes including Cd14 and Ccr2 (Figure 2E). Recent studies have identified at least two subtypes of Sirpa+ DCs. Brown et al. (2019) recently characterized cDC2a and cDC2b populations by the differential expression of T-bet and RORγt, respectively, with cDC2b displaying monocyte features. We used the previously published cDC2a- and cDC2b-specific gene signatures (Brown et al., 2019) (Table S2) to identify the presence of these cells within Sirpa+ expressing clusters and compare these signatures to monocytes (C15). Among Sirpa+ cDCs, C7 displayed the highest cDC2a AUC score, whereas C2 and C11 displayed the highest cDC2b AUC scores (Figure 2G). Indeed, the monocyte cluster C15 expressed the highest cDC2b AUC score overall (Figure 2G), but previous studies have shown that despite expression of monocyte features, cDC2b cells are not monocyte-related (Brown et al., 2019). The remaining Sirpa+ expressing clusters displayed moderate cDC2a AUC scores.

Furthermore, differential gene expression (DGE) analysis (Table S3) between clusters (one versus remaining) revealed Cd7 as a transcript feature dominantly expressed by cDC2a clusters (Figure 2H). We propose that Cd7 in combination with Sirpa, MHCII, and Itgax transcripts could therefore identify cDC2a. Cd7 expression by cDC2a cells has been observed by others in DC single-cell sequencing experiments (Brown et al., 2019; Bosteels et al., 2020). We further note that whereas no Cd7 expression was observed in cDC1 clusters, the level of expression varied within cDC2a clusters, and was highest expressed in C0, suggesting that Cd7 expression is regulated.

In addition to the cDC2a clusters (C0, C5, C7, C9, C8, and C10) and cDC2b clusters (C2 and C11), we identified two main Sirpa+ cDC clusters (C1 and C3) that expressed unique DEGs (Table S3). C1 was characterized by expression of genes associated with TNF signaling as well as the highest expression of Relb and Tcf7, whereas lacking expression of Itgax, Cd7, or any cDC2b features such as Cd14 and Ccr2 (Figure 2B). C1 also expressed Ccr7, Cd40, Fas, and the immune-suppressive genes Cd274 (PD-L1) (Figure 2J) and Il4i1. Based on expression of Fabp5, Spint2, and Il4i1, this cluster aligned with a migratory cDC2 phenotype.
described by others (Bosteeils et al., 2020). However, we note that C1 cells were distinct from other Ccr7-
expressing cDC2a (C8) and cDC2b (C11) cells in our dataset (see DEGs between C1, C8, and C11 in Table
S4). We annotated C1 as Il4i1+cDC2 as Il4i1 was a highly significant DEG (Figure 2I). The other distinct cDC
cluster C3 highly expressed Sox4, Cd33, and Tcf4 (Figures 2I, Table S3). Together with expression of Sirpa,
Cd8a, Cd14, Ccr2, and low levels of Siglech, this cluster displayed similarities to a murine Siglech+c DC
phenotype described by others, sharing pDC and cDC2 characteristics, which were termed transitional
DCs (Leylek et al., 2019; Brown et al., 2019), and similar populations have also been described in different
contexts (Bar-On et al., 2010; Schlitzer et al., 2011; Rodrigues et al., 2018). Based on the most significant
DEG, we annotated this cluster as Sox4+cDC (Figure 2I).

To test whether proliferation contributed to cluster heterogeneity, we predicted the cell cycle phase
of each individual cell using S and G2M signature genes defined by Tirosh and colleagues (Tirosh et al.,
2016) (Table S5) and identified C5, C6, C7, C10, and C19 as mitotic (Figure 2K).

DGE analysis between clusters revealed a prominence of mitochondrial gene expression in C9 and C17
(Suppl. File 1, Data S3, Figure S3C), and these clusters simultaneously expressed lower levels of MHCII
genes and Itgax (Figure S3B). Cluster C18 was enriched in ribosomal subunit genes (Table S3, Figure S2C).
After removing low-quality cells in the initial quality control (see transparent methods) and before PCA and
clustering, we further analyzed the data for expression of genes associated with apoptosis, and no cluster
expressed particularly increased levels of apoptosis genes (Figure S3D), however, C9 and C17 had mark-
edly lower unique molecular identifier (UMI) counts (Figure S3E).

We constructed a knowledge-informed list of canonical genes characteristic to cDC1, cDC2, and pDCs
(Tomasello et al., 2018). This list, consistent with the cDC1 AUC score, revealed high expression of
cDC1-associated genes in C4, C6, C12, C17, and C18 (Figure S4). Analyzing DEG-enriched pathways
(Table 1) and maturation features (Figure 2J), we identified C1, C8, C11, and C12 as clusters with
increased maturity based on expression of Ccr7, co-stimulatory molecules (Figure 2J), or genes asso-
ciated with TNF signaling. Analysis of TLR genes further enabled cDC lineage annotation (Figure 2J).
Using the information from each of these analyses, we annotated clusters according to cell lineage
and cell state (Table 1).

To enable identification of key genes differing between core WT DC clusters, we isolated WT cells and
removed clusters characterized by confounding cellular processes such as proliferation (C5, C6, C7, C10)
and high expression of mitochondrial (C9, C17) and ribosomal genes (C18), which would hinder identi-
fication of key steady state genes between DC clusters. We then built metaclusters for cDC1 (immature
C4 + mature C12), cDC2a (immature C0 + mature C8), and cDC2b (immature C2 + mature C11) (Figure
3A). We performed DGE analysis between these core cDC1 and cDC2 clusters as well as Il4i1+
cDC2 (C1), Sox4+cDCs (C3), pDCs (C14), and monocytes (C15) (Table S6). cDC1, Il4i1+cDC2, pDC,
and monocytes displayed unique gene signatures, whereas DEGs of cDC2a, cDC2b, and Sox4+cDC
were less clearly defined (Figure 3B). Many cDC2a DEGs were also expressed in Sox4+ DC, including
Mdh2, Cd7, Siglech, and Runx3, suggesting a partial overlap. cDC2b DEGs were largely expressed by
| Cluster | Cell lineage | Cell state       | GO Cellular Component enrichment                                                                 |
|---------|--------------|------------------|--------------------------------------------------------------------------------------------------|
| 0       | Sirpa+ cDC2a | Immature, phagocytic | Secretory granule lumen (GO:0034774), cytoplasmic vesicle lumen (GO:0060205), tertiary granule (GO:0070820) |
| 1       | Sirpa+ Il4i1+ cDC2 | Mature | CD40 receptor complex (GO:0035631), tertiary granule (GO:0070820), specific granule (GO:0042581), |
| 2       | Sirpa+ cDC2b  | Immature, phagocytic | Tertiary granule (GO:0070820), tertiary granule membrane (GO:0070821), azurophil granule (GO:0042582) |
| 3       | Sirpa+ Sox4+ cDC | Immature, phagocytic | Tertiary granule (GO:0070820), tertiary granule membrane (GO:0070821), specific granule (GO:0042581) |
| 4       | cDC1         | Immature, phagocytic | Lysosomal lumen (GO:0043202), lysosome (GO:0005764), phagocytic vesicle (GO:0045335)               |
| 5       | Sirpa+ cDC2a | Mitotic | Nuclear chromosome part (GO:0044454), chromosome, telomeric region (GO:0000781), nuclear chromosome, telomeric region (GO:0000784) |
| 6       | cDC1         | Mitotic, immature | Mitochondrial inner membrane (GO:0005743), nuclear chromosome part (GO:0044454), mitochondrial proton-transporting ATP synthase complex (GO:0005753) |
| 7       | Sirpa+ cDC2a | Mitotic | Nuclear chromosome part (GO:0044454), spindle (GO:0005819), chromosome, centromeric region (GO:0000775) |
| 8       | Sirpa+ cDC2a | Mature | P-body (GO:0000932), fibrillar center (GO:0001650), nucleolar part (GO:0044452)                  |
| 9       | Sirpa+ cDC2 (mixed) | Mitochondrial | Nuclear body (GO:0016604), nuclear speck (GO:0016607), chromatin (GO:0000785),                 |
| 10      | Sirpa+ cDC2a | Mitotic | Spindle (GO:0005819), chromosome, centromeric region (GO:0000775), condensed chromosome, centromeric region (GO:0000779) |
| 11      | Sirpa+ cDC2b | Mature | Tertiary granule (GO:0070820), ficolin-1-rich granule (GO:0101002), lysosomal lumen (GO:0043202) |
| 12      | cDC1         | Mature | CD40 receptor complex (GO:0035631), actin cytoskeleton (GO:0015629), cytoskeleton (GO:0005856) |
| 13      | NK/ILC/T     |                  | Cytosolic ribosome (GO:0022626), T cell receptor complex (GO:0042101), cytosolic part (GO:0044445) |
| 14      | PDCs         |                  | Lysosomal lumen (GO:0043202), lysosome (GO:0005764), specific granule membrane (GO:0035579) |
| 15      | Monocytes    |                  | Tertiary granule (GO:0070820), focal adhesion (GO:0005925), specific granule (GO:0042581) |

(Continued on next page)
monocytes. A significant number of Sox4+ cDC DEGs were also highly expressed in pDCs, such as Lag3, Siglech, Tcf4, Spib, and Cd37.

As myeloid cells have become an attractive target for immune checkpoint inhibition therapy to increase their priming capacity, we analyzed the expression of immune checkpoint genes across metaclusters (Figure 3C). This analysis identified that cDC1 cells are the main DCs expressing Btla and Havcr2 (encoding TIM-3), Il4i1+ cDC2 express high levels of Icosl, whereas monocytes express Vsir (encoding VISTA). Interestingly, both pDCs and Sox4+ cDCs expressed the highest levels of Lag3.

Simultaneous detection of transcript and epitope enables confident cDC lineage identification

Although a cDC1-characteristic gene transcript signature is clearly identified, confident classification of cDC2 is more challenging, as transcription of key cDC2 surface molecules, such as Itgam (CD11b) and Cd4, is limited (Stoeckius et al., 2017; Peterson et al., 2017). Although it is possible to partly overcome this challenge by recovering single-cell gene expression signals, for example, by using kernel density estimation algorithms implemented in Nebulosa (Alquicira-Hernandez and Powell, 2021), parallel detection of protein expression is required to compare transcriptomic and proteomic data. We integrated transcript and epitope multimodal analysis by adding barcoded antibodies specific to DC-characteristic surface molecules. Binding of the barcoded antibodies was confirmed using an oligo-dT probe conjugated to a fluorochrome dye and flow cytometry analysis (Figure 4A). Antibody-derived tag (adt) counts of the complete dataset, each genotype, and each cluster were converted to flow cytometry standard (FCS) files. Analysis of the median adt value for each cluster revealed that each Total-Seq antibody produced a substantial background read count (Figure 4B). This is in line with observations by others when using high antibody concentrations as recommended in suppliers’ protocols (Buus et al., 2021). Comparing the characteristic features of cDC1 (CD8A, CD24, XCR1) against those of cDC2 (CD11b, CD172A, CD4) in this way revealed that the expression of these markers was largely restricted to a single-cell type (Figure 4B). In line with previous analysis, the cDC1 clusters C4, C6, C12, C17, and C18 expressed increased levels of CD8A and CD24, whereas the cDC2 clusters C0, C1, C2, C5, and C7–C11 expressed increased levels of CD11b and CD172A (Figure 4B).

Visualization of the adt expression using a conventional FCS dotplot of CD8A versus CD11b or CD24 versus CD172A did not allow confident delineation of cDC1 and cDC2 clusters (Figure 4C); however, comparison between WT and Batf3−/− genotypes revealed that a population of CD24+ XCR1+ cells indicative of cDC1 cells was missing in Batf3−/− mice (Figure 4C).

A positive adt count for each antibody in each cluster suggested a level of non-specific binding. Informed by the percentage of positive staining in the flow cytometry validation data using the oligo-dT probe described in Figure 4A, we determined adt count thresholds above which we considered the adt signal specific, and all values below this threshold were set to 0. After thresholds were applied, the expression of Cd8a transcript and CD8A epitope correlated strongly (Pearson’s correlation, r = 0.75; Figure S3F).

Table 1. Continued

| Cluster | Cell lineage | Cell state | GO Cellular Component enrichment |
|---------|--------------|------------|----------------------------------|
| 16      | Mixed identity | Secretory granule lumen (GO:0034774), ficolin-1-rich granule lumen (GO:1904813), cytoplasmic vesicle lumen (GO:0060205) |
| 17      | cDC1          | Mitochondrial | Mitochondrial inner membrane (GO:0005743), mitochondrial respiratory chain complex I (GO:0005747), nuclear speck (GO:0016607) |
| 18      | cDC1          | Ribosomal   | Cytosolic ribosome (GO:0022626), cytosolic part (GO:0044445), cytosolic large ribosomal subunit (GO:0022625) |
| 19      | Stem cells    | Nuclear chromosome part (GO:0044454), spindle (GO:0005819), mitochondrion (GO:0005739) |
Figure 3. Diversity in Sirpa+ DCs is less distinct than diversity between other DC lineages

(A) Clusters defined by active cell cycle or metabolic activity were excluded, and metaclusters corresponding to cDC1, cDC2a, cDC2b, Il4i1+ cDC2m Sox4+ cDC, pDC, and monocytes were established.

(B) Dotplots showing the top 50 DEGs of each metacluster (see Table S6).

(C) Dotplots showing expression of immune checkpoint genes in each metacluster.
Figure 4. Total-Seq antibodies verify cDC1 and cDC2 identity

(A) Verification of Total-Seq antibody binding. Total-Seq antibodies were hybridized with a fluorochrome-conjugated dT probe and subsequently incubated with C57BL/6 splenocytes together with fluorochrome-conjugated antibodies against CD11c and MHCII. Cells were pre-gated on CD11c+ MHCII+ DCs and analyzed for binding of Total-Seq antibodies. Positive signal for Total-Seq antibodies were used to determine thresholds for bioinformatics analysis of adt signals. Total-Seq antibodies where specific staining could not be confirmed were excluded from further analysis of CITE-seq data.

(B) Adt sequencing data of each cluster was converted to FSC files using the flowCore package and analyzed using flow cytometric BD Kaluza analysis software. Shown are median adt tag counts for each cluster. Horizontal line represents background, based on cDC2 cluster C0 for cDC1 features CD8A, CD24, and XCR1, and based on cDC1 cluster C4 for cDC2 features CD11b, CD172A, and CD4.

(C) Adt data of cDC1 features CD8A, CD24, and XCR1 or cDC2 features CD11b and CD172A were compared between WT and Batf3−/− genotypes.

(D) Featureplots of RNA and corresponding thresholded adt counts.

(E) Dotplot of thresholded adt counts for each cluster.
C6: mitotic immature cDC1
C5: mitotic immature cDC2
Batf3-/-

A

| Cluster | Gene set | Biological Processes |
|---------|----------|----------------------|
| C6      | Neutrophil activation involved in immune response; Neutrophil-mediated immunity; Neutrophil degranulation | |
| C4      | Neutrophil activation involved in immune response; Neutrophil-mediated immunity; Neutrophil degranulation | |
| C12     | Neutrophil activation involved in immune response; Neutrophil-mediated immunity; Neutrophil degranulation | |
| C6      | RNAsplicing via transesterification reactions; RNAsplicing via spliceosome; mRNA processing | |
| C4      | RNAsplicing via transesterification reactions; RNAsplicing via spliceosome; mRNA processing | |
| C12     | RNAsplicing via transesterification reactions; RNAsplicing via spliceosome; mRNA processing | |
| C17     | Mitochondrial ATP synthesis coupled electron transport; Respiratory electron transport chain; Mitochondrial electron transport | |
| C18     | SRP-dependent cotranslational targeting to membrane; Proteintargeting to ER; Nuclear-transcribed mRNA catabolic processes | |

B

Average Expression

Percent Expressed

WT

Batf3-/-

C

Expression Level

Ltb Cybb S100a4 Ccr2 Tmem176a Tmem176b Sirpa

Irf8 Id2 Naad Cadm1 Xcr1 Clec9a Trl3 Trl11 Cd24a Cd8a

G

Expression Level

adt-CD24 adt-CD8A adt-XCR1 adt-CD11B adt-CD172A adt-CD4

Irf8 Cadm1 Xcr1 Clec9a Cor7

Trl3 Trl12 Cd8a Cd24a Cd40

H

Expression Level

adt-CD24 adt-CD8A adt-XCR1 adt-CD11B adt-CD172A adt-CD4

I

Expression Level

adm-CD24 adm-CD8A adm-XCR1 adm-CD11B adm-CD172A adm-CD4
Figure 5. Residual mitotic cDC1-like cells in Batf3−/−mice increase expression of cDC2 features
(A) GO analysis of cDC1 clusters C4, C6, C12, C17, and C18 (see Table S7).
(B) Distribution of cells assigned to WT and Batf3−/−genotypes.
(C) Trajectory analysis using RNA velocity.
(D) DEGs between WT and Batf3−/−of cDC1 cluster C6 (see Table S1).
(E) Top 40 DEGs between Batf3−/−mitotic cDC1 state C6 and Batf3−/−mitotic cDC2 state C5.
(F) Gene expression of canonical cDC2 and cDC1 features between mitotic cDC1 state C6 and mitotic cDC2 state C5 (related to Figure S5).
(G) Protein expression (adt) of canonical cDC2 and cDC1 features between mitotic cDC1 state C6 and mitotic cDC2 state C5. (H) DEGs between WT and Batf3−/−of cDC1 cluster C4 (see Table S1).
(I) Gene and protein expression of canonical cDC1 features between mitotic cDC1 (C6), immature cDC1 (C4), and mature cDC1 (C12).

We detected higher levels of adt-CD11B protein expression compared with the corresponding gene expression, indicating a substantial benefit to the simultaneous detection of CD11B as an important cDC2 feature (Figure 4D). In contrast, adt detection of CD24, XCR1, CD172A, and CD4 was weaker, when compared with the corresponding Cd24a, Xcr1, Sirpa, and Cd4 RNA detection. The combination of both RNA and adt protein data verified the identity of cDC1 and cDC2 clusters (Figures 4D and 4E).

Residual cDC1-like cells in Batf3−/−mice
Batf3 affects the development and maintenance of cDC1 cells, and we therefore explored the cDC1 landscape between WT and Batf3−/−mice. The cDC1 landscape was represented in three major clusters C4, C6, and C12 (Figures 2D and 2E) and two minor clusters C17 and C18 enriched in mitochondrial and ribosomal genes, respectively (Figure S3C). To gain more insight into the differences between cDC1 cell states, we analyzed DEGs between all five cDC1 clusters (Table S7). GO analysis of DEGs suggested that C4 was enriched in gene expression associated with neutrophil responses, whereas C6 was associated with RNA splicing, and C12 with cytokine signaling and NF-kB signaling (Figure 5A).

cDC1 cluster C6 was equally represented by 797 WT and 727 Batf3−/−cells, but the two genotypes presented as distinct C6 populations suggesting genotype-based differences (Figure 5B). Functional enrichment analysis suggested that Batf3−/−C6 cells were clustered together with WT C6 cells based on the expression of genes related to mitochondrial inner membranes and nuclear chromosome parts, likely connected to their active cell cycle state (Table 1, Figure 2K, Table S3). The mitotic state of C6 cells suggests that these cells might represent an early trajectory to cDC1 cells, but most Batf3−/−C6 cells would not reach the cDC1 states represented by C4 and C12. RNA velocity analysis, a technique used to predict future cell states from the abundance of spliced and unspliced transcripts in single cells (La Manno et al., 2018; Bergen et al., 2020), revealed a distinct path from WT cluster 6 to cluster 4 and finally to cluster 12 (Figure 5C).

Differential expression analysis of C6 between WT and Batf3−/−cells identified 101 DEGs (adj. p value < 0.05), of which 62 genes were significantly downregulated in Batf3−/−cells and 39 genes were significantly upregulated (Figure 5D, Table S1). Downregulated genes in Batf3−/−C6 included Naaa, Xcr1, Cadm1, Cd8a, Th3, and Thr11 as cDC1-defining genes, and Cxcl16, Cd81, Cd86, and Cd36 as genes related to maturation (Figure 5D); however, we did not observe specific GO or pathway-related enrichment. Upregulated genes in Batf3−/−C6 included Ltb, Cytb (NOX2), and S100a4 as cDC2 features, and Ccr2, Cd209a (DC-SIGN), and Tmem176a/b, indicative of a cDC2a phenotype (Figure 5D). Noting that Batf3−/−C6 cells demonstrated downregulation of cDC1 features and upregulation of cDC2 features when compared with WT C6 cells, we analyzed DEGs between Batf3−/−C6 and the corresponding mitotic cDC2 cell state C5. Compared with classical mitotic cDC2 C5 cells, Batf3−/−C6 cells displayed increased levels of cDC1-defining genes and decreased levels of cDC2-defining genes (Figure 5E). By comparing the expression of cDC1 and cDC2 characteristic transcriptomic features between mitotic cDC2 cluster C5 and mitotic cDC1 cluster C6, and between genotypes, we demonstrate an intermediate phenotype of Batf3−/−mitotic cDC1 C6 cells based on their expression of most cDC1 and cDC2 (Figure 5F), and comparable results were obtained at protein level (Figure 5G). Interestingly, Batf3−/−C6 cells downregulated many cDC1 features but maintained comparable expression levels of Cd24a and Id2 (Figure 5F). To analyze the distribution of cDC1 and cDC2 characteristic genes across Batf3−/−C6 cells, we paired (1) Batf3−/−C6 cells with WT C6 cells, revealing weak yet significant cDC2 gene expression compared with classical WT mitotic cDC1 cells or (2) Batf3−/−C6 cells with mitotic cDC2 C5 cells, which revealed weak yet significant cDC1 gene expression compared with classical mitotic cDC2 cells. The highest density of cDC1 and cDC2 genes was observed within the same Batf3−/−C6 cells (Figure S5). Our data suggest that mitotic early cDC1 cells might divert...
to the cDC2 lineage in the absence of Batf3. RNA velocity analysis predicted that mitotic Batf3−/− cDC1 (C6) cells either continued toward immature cDC1 cell state (C4) or followed a trajectory toward immature cDC2b (C2) (Figure 5C). It remains to be determined whether Batf3−/− mitotic cDC1 cells continue to increase expression of cDC2 features, resulting in a full convergence to cDC2, or abort a cDC2 trajectory altogether.

cDC1 cluster C4 identified 38 DEGs (adj. p value < 0.05) between genotypes, of which 14 were significantly downregulated in Batf3−/− cells, whereas 24 were significantly increased (Figure 5H, Table S1). Downregulated gene transcripts in Batf3−/− C4 included the cDC1-characteristic gene Naaa, whereas upregulated genes also included cDC2-characteristic genes Ltb, Cybb (NOX2), Cd209a (DC-SIGN), and Tmem176a/b (Figure 5H), similarly to those observed for C6. Of note, C4, predicted to be an immature cDC1 state when considering expression of activation features (Figure 2J), was represented by 2,202 WT cells but only 31 Batf3−/− cells, suggesting that Batf3 is required to reach this immature cDC1 cell state, and that the absence of Batf3 increases expression of cDC2 features.

Furthermore, cDC1 cluster C12 was represented by 394 WT cells but only 31 Batf3−/− cells, indicating that Batf3 was also required for this cDC1 cell state. We identified 12 DEGs (adj. p value < 0.05) between genotypes, of which only Batf3 was significantly downregulated in Batf3−/− cells, whereas expression of 11 genes was significantly increased (Table S1). These genes were, however, not associated with particular pathways and cell types. The expression of cDC1-characteristic genes Cadm1, Xcr1, Clec9a, Tlr3, and Tlr11 was virtually absent in C12 cells, and Irf8 expression was reduced (Figure 5I). C12 cells, however, expressed the highest levels of activation features Ccr7, Cd40, Cd83, Fas, and Cd274 (PDL1) (Figures 2J and 5I), indicating that C12 represents mature cDC1 cells. Protein features CD24, CD8A, and XCR1 were maintained on C12 cells and were comparable to immature C4 cells (Figure 5I). DEGs of C12 cells were further enriched in TNF signaling (Table S7). Maintenance of XCR1 protein with downregulation of Xcr1 gene expression on mature cDC1 has been reported by others (Ardouin et al., 2016; Crozat et al., 2011). DGE analysis between WT and Batf3−/− C12 cells suggested a high degree of similarity (Table S1), suggesting that the Batf3-dependent cDC1 trajectory follows an “abort/divert or proceed” principle.

**Characterization of Sox4+ cDCs**

Given our observation that a small proportion of cDCs express both the cDC1 feature CD8 and the cDC2 feature CD11B, as described in Figure 1, we aimed to identify cells with these features in the sequencing dataset. We analyzed the joint density of Cd8a and Itgam using the Nebulosa package and found that the Sox4+/cDCs cluster (C3) was the only cluster expressing both features (Figure 6A). This joint signal was only observed in Batf3−/− Sox4+/cDCs, and not in WT Sox4+/cDCs (Figure 6A). Rather, WT DCs with a joint Itgam/Cd8a signal were observed in C4 immature cDC1 cells, although at reduced density (Figure 6A). The adt protein expression data showed that a proportion of Sox4+/cDCs expressed CD8A, CD24, CD172A, CD11B, and CD4 (Figure 6B). These data suggest that the CD8+CD11b+ cDCs described in Figure 1 might represent a mixture of cells, including Sox4+ cDCs and cDC1. We further assessed this using the adt-converted FCS files of each cluster using standard flow cytometry software, where we compared expression of CD11B versus CD8A and CD172A versus CD24 between selected cDC2 clusters (C0 and C2), cDC1 clusters (C4 and C12), pDCs (C14), and Sox4+ cDCs (C3), confirming that a proportion of Sox4+ cDCs and cDC1 cells expressed CD24, CD8A, CD172A, and CD11B (Figure 6C).

DGE analysis of Sox4+/cDCs (C3) revealed Batf3 as the only significant DEG between WT and Batf3−/− genotypes (Table S1), whereas DGE analysis between Sox4+/cDC (C3) and all other clusters and metaclusters revealed that C3 expressed key genes of several DC lineages including Cd8a, Cd7, Siglech, Tcf4, and Spib (Tables S1, S3, and S6, top 50 in Figure 3B). C3 expressed levels of MHCII genes comparable with other cDC clusters (Figure S3B). We compared the expression of key canonical DC genes between Sox4+/cDCs and cDC1, cDC2a, cDC2b, Irf4+/cDC2, pDC, and monocytes (Figure 6D). Sox4+/cDCs expressed the highest level of Irf4 among these clusters. Of note, compared with cDC1 cells, Sox4+/cDCs cells did not express cDC1-characteristic transcription factors such as Irf8, Idd2, or functional genes Xcr1, Clec9a, Tlr3, or Tlr11, or only in a very small fraction of cells. Sox4+/cDCs cells expressed moderate levels of pDC canonical genes Siglech, Spib, Tcf4, Bcl11a, and Runx2, but also lacked expression of pDC-characteristic TLRs (Figures 2I and 2J). Instead, Sox4+/cDCs expressed high levels of Sox4, Runx1, and Flt3, which are also highly expressed by pre-DCs (Figure 6D). However, splenic pre-DCs are considered to be MHCII negative (Schlitzer et al., 2015), and we hypothesize that these cells might represent a cell state of pre-DC to cDC transition as described previously (Leylek et al., 2019). This cluster contained 2,245 cells,
Figure 6. Characterization of a Sox4+ cDC cell state

(A) Joint density of Itgam and Cd8a, using the Nebulosa package, identifying Sox4+ cDCs (C3) as cluster with co-expression in Batf3−/−/−.

(B) Protein expression (adt signals) between metaclusters defined in Figure 3A.

(C) Adt data of selected clusters were converted to FCS files and analyzed for co-expression of CD24 and CD172A or CD8A and CD11B using flow cytometric BD Kaluza software.

(D) Expression of canonical DC genes between metaclusters.

(E) Subclustering of Sox4+ cDCs (C3) using a resolution of 0.3.

(F) GO pathway analysis of C3 subcluster-specific DEGs (see Table S8).

(G) Distribution of canonical genes for DCs, pDCs, cDC1, cDC2, tDCs, and pre-DCs using the Nebulosa package.

(H) GSEA using gene signatures from Brown et al. (2019); Leylek et al. (2019); Villani et al. (2017) and the AUCell package (see Table S2).
representing 9% of all DCs, and was enriched in Batf3−/− DCs (12.7%) compared with WT DCs (5.8%). Recent single-cell analyses described similar populations of Siglech+ DCs (Brown et al., 2019; Leylek et al., 2019) and suggested a possible link to a phenotype of circulating human AXL+ DCs (See et al., 2017; Villani et al., 2017; Alcantara-Hernandez et al., 2017).

The distribution of gene and protein expression within the Sox4+ cDCs cluster indicated additional cell state heterogeneity. We therefore subclustered C3 into three cell states (Figure 6E). DGE analysis revealed a significant number of DEGs between C3.0 and C3.1, but only two DEGs in C3.2, suggesting that C3.2 represents a mixture of C3.0 and C3.1 cells (Table S8). Pathway analysis determined that DEGs of C3.0 were enriched in pathways of MAPK signaling, IL-17 signaling, NF-κB signaling, and Th1/Th2 differentiation, whereas DEGs of C3.1 were enriched in pathways of phagocytosis and antigen presentation (Figure 6F), suggesting that C3.0 is more mature than C3.1. We determined the expression of DC lineage characteristic features across C3 cell states. The three C3 cell states expressed comparable levels of Itgax (CD11c) and MHCII genes (Figure 6G). Expression of pDC-characteristic features varied across cell states, with C3.1 expressing high levels of Tcf4 and Sipb, whereas Bcl11a was similarly expressed across all C3 cell states (Figure 6G). The highest levels of cDC1 features Cd24a and Cd8a were expressed in C3.1, showing a low to high expression gradient from C3.0 to C3.1 (Figure 6G). Irf8 and Cadm1 were expressed only in a very small proportion of C3.1 cells. In contrast, the highest levels of cDC2 features were expressed in C3.0, with a low to high expression gradient from C3.1 to C3.0 (Figure 6G). We further examined features of previously described “transitional” Siglech+ DCs (tDC) (Leylek et al., 2019), including Cd14, Cxcr3, Irf4, and Runx2, which partly overlap with cDC2b cells and had the highest expression in C3.1 (Figure 6G). Last, we examined features characteristic to pre-DCs. We observed that, whereas Flt3 was similarly expressed across C3 cell states, Sox4 had the highest expression in C3.0, but also lacked Ly6c1 expression (Figure 6G). Gene set enrichment analysis (GSEA) determined that the Sox4+ cDC cluster was aligned with Siglech+ cDCs (Brown et al., 2019) and tDCs (Leylek et al., 2019), as well as with a transcription factor signature of AXL+ human cDCs (Villani et al., 2017) (Figure 6H, Table S2).

Our data revealed that splenic CD11c+ MHCII+ DCs contain at least two cell states with lineage mixed features, which share expression of Bcl11a, Sipb, and Flt3, but co-express either Tcf4, Cd8a, Cd24a, Cd14, Irf4, Cx3cr1, Runx2 and Siglech, or Itgam, Sirpa, Cd4, Esam, and Sox4 (Figure 6G). However, these cell states are connected via cells expressing moderate levels of these features, indicating cell state plasticity.

To verify whether Sox4+ cDCs contribute to the lineage-mixed phenotype observed in Figure 1, we analyzed splenic cDCs of WT and Batf3−/− mice with an extended flow cytometry panel. Expression of Cd33 was largely unique to pDCs and Sox4+ cDCs (Figures 2I and 3B). We thus compared expression of common cDC features across cDC1, cDC2, CD33+ cDCs, and pDCs (Figures 7A–7D). CD33+ cDCs were distinct from pDCs but overlapped with both cDC1 and cDC2 in both WT and Batf3−/− cDCs with regard to expression of CD172A, CD11B, CD24, and CD8A (Figures 7A and 7B). CD33+ cDCs contributed to a population of cells expressing CD172A, CD11B, CD24, and CD8A, but did not represent the whole population (Figures 7A and 7B). In line with this, the transcriptomic data suggested that less than 50% of Sox4+ cDCs expressed Cd33, and hence other identifying surface features will be required to investigate this population in detail. Flow cytometry analysis confirmed that CD33+ cDCs largely lacked expression of the pDC feature PDCA1 (Figure 7C). All CD33+ cDCs expressed increased levels of CD24 and CD8A compared with cDC2 cells and pDCs, and decreased levels compared with cDC1, but largely lacked expression of XCR1 (Figure 7C). Expression of CD11B, CD172A, and Ly6C on CD33+ cDCs was biphasic (Figure 7C), suggesting heterogeneity of two cell states as observed in Sox4+ cDCs of the transcriptomic data. The flow cytometric analysis further revealed that Batf3−/− cDC1 cells had a significantly increased expression of CD172A and CD11B, whereas expression of CD8A and XCR1 was decreased and CD24 was unchanged (Figure 7D). We further confirmed that a small proportion of WT cDC1 expressed CD11B (Figure 7B, C).

Combined, these data confirm that cDCs with lineage-intermediate phenotypes with regard to expression of CD11B, CD8A, and CD24 can represent either a fraction of cDC1 cells or a population of CD33+ cDCs, whose functionality remains to be determined.

**DISCUSSION**

We previously identified that in the absence of Batf3, a proportion of cDCs displays a lineage-intermediate phenotype defined by co-expression of CD11B, CD172A, and CD8A (Chandra et al., 2017). In the present
study, we observed cDCs with a lineage-intermediate phenotype, further differentiated by altered expression of classical cDC surface features. We implemented multi-modal single-cell transcriptome and epitope sequencing technology to characterize this cDC lineage-intermediate cell state in detail. We identified a significant proportion of cDCs with lineage-intermediate gene and protein expression signatures displaying characteristic features of cDC1, cDC2, and pDCs in both WT and Batf3−/−/−/− spleens. These cells lacked a transcriptional signature of maturity, but instead expressed high levels of Sox4, Flt3, and Cd33, features associated with a pre-DC phenotype. This population was enriched in the Batf3−/−/−/− cDC compartment compared with WT. Furthermore, Batf3−/−/−/− mice lacked classical cDC1 cells but harbored a population of mitotic immature cDC1-like cells expressing reduced levels of cDC1 genes but increased levels of cDC2 genes. Batf3−/−/−/− cDC1-like cells lacked expression of genes linked to cDC functionality including Xcr1, Clec9a, and cDC1-specific TLRs. Algorithms predicting developmental trajectories suggested that these Batf3−/−/−/− mitotic cDC1-like cells give rise to either cDC1 and cDC2 lineages.

Flow cytometric analysis of murine splenic DCs can delineate two main populations: cDC1 and cDC2 based on expression of XCR1/CADM1 and CD11B/CD172A, respectively (Leylek et al., 2019). With the advancement of high-dimensional single-cell technologies, recent work has redefined the comprehension of DC heterogeneity. Unsupervised clustering approaches based on principal-component analysis have enabled

Figure 7. Immunophenotyping of CD33+ DCs
Splenocytes of WT (n = 3) and Batf3−/−/−/−/− (n = 3) were incubated with a panel of 14 fluorochrome-conjugated antibodies. (A) CD33+ cDCs were gated as singlet live CD45+ LY6G−/− CD11C+ MHCII+. pDCs were gated as Lin−/− LY6G−/− PDCA1+ LY6C+. cDCs were gated as Lin−/− LY6G−/− CD11C+ MHCII+ LY6C−/−, and from these cDC1 and cDC2 were gated as XCR1+ or CD172A+, respectively. (B and C) Overlays of pDCs, cDC1, cDC2, and CD33+ cDCs. (D) Median fluorescence intensity (MFI) of DC features between WT and Batf3−/−/−/− cDC1, cDC2, pDCs, and CD33+ cDCs. Each data point represents data of individual animal with mean ± standard error of the mean (SEM). Statistical significance was determined using unpaired t test with Welch correction. **p < 0.01, ***p < 0.001, ****p < 0.0001.
the identification of cell state heterogeneities beyond pure phenotypic delineation of DC subtypes. However, because current algorithms do not distinguish between cell subsets and cell states when dividing data into clusters based on differential gene expression, the cluster annotation and interpretation remains a challenge, especially within a lineage where a continuum of different cell states rather than distinct subtypes is to be expected (Bassler et al., 2019), and this model is important to consider when interpreting clustered data. Our data support this paradigm as we observe that certain features are expressed only in a fraction of certain clusters. The presented data provide a basis for analysis of the continuum of DC states in more detail and reveal, for example, a certain cluster containing multiple cell states shared by cDC1, cDC2, and pDCs.

Recent high-dimensional analysis of DCs has uncovered additional heterogeneity particularly in the cDC2 compartment. At least two distinct cDC2 populations have been suggested in human and mice in recent years. In mice, these populations are distinguished by expression of Cd7/Dtx1/Ccr6 (cDC2a) and Cd209a/Clec10a/Clec12a (cDC2b) (Brown et al., 2019), and in human corresponding cells have been termed DC2 and DC3 (Cytlak et al., 2020; Bourdely et al., 2020; Bosteels et al., 2020; Dutertre et al., 2019). Additional minor populations of cDC2 cells have been described, such as Siglech+cDC2 (Brown et al., 2019) or transitional cDCs (Leylek et al., 2019). Our murine splenic DC dataset provides support of this cDC2 heterogeneity. We identified populations of immature and mature cDC2a and cDC2b populations, as well as a Il4i1+cDC2 population enriched in gene expression suggestive of an activated and inflammatory phenotype and distinguished through expression of Tcf7, with downregulated classical cDC2a and cDC2b features. It remains to be determined if and how this Il4i1+cDC2 population is related to the other cDC2 populations.

We further identified a population of Siglech+cDCs, which we termed Sox4+cDCs based on its most DEGs. These Sox4+cDCs expressed lineage-mixed features of cDC1, cDC2, pDCs, and pre-DCs and could be further divided in two substates. DC cell states combining lineage features of cDC1, cDC2, and pDCs have been described by others. Wylie et al. recently identified a CD8+XCR1– DC population, which expressed lineage-mixed gene signatures of pDCs, cDC1, and cDC2 (Wylie et al., 2018). They showed that these CD8+XCR1– DCs preferentially expressed TLR5 and TLR7, and a unique set of endocytic receptors, and their ability to induce T helper cell proliferation was comparable with cDC2. The Sox4+cDC cell state described here expresses low levels of Tlr5 gene, and no other TLR genes, suggesting that these cells are limited to respond to pathogen-associated stimuli. Furthermore, Bar-On et al. described a steady-state CD8+ CX3CR1+ non-canonical DC population, which is related to pDCs (Bar-On et al., 2010). In fact, the Sox4+cDC cell state we describe here highly expressed Cx3cr1 in one cell state. Although Cx3cr1 is expressed by many leukocytes including monocytes, macrophages, and DCs, it is significantly more highly expressed by pre-DCs (Choi et al., 2019), and the herein described Sox4+cDC cell state expressed elevated levels of Cx3cr1 and other pre-DC features such as Flt3, Cd33, and Csfr1, whereas any signs of maturity were absent. CCR2 was described as a required chemokine receptor to enable pre-DC migration (Nakano et al., 2017), and indeed the Sox4+cDC cell state expressed increased levels of Ccr2 compared with cDC1, cDC2a, and pDCs; however, CCR2 is also expressed by monocytes and the newly defined cDC2b population (Brown et al., 2019).

Schlitzer et al. described the transcriptional regulation of the MDP-to-CDP-to-pre-DC trajectory in detail and identified different pre-DC cell states that range from non-lineage primed to cDC1 or cDC2 lineage primed (Schlitzer et al., 2015). In spleen, Schlitzer et al. identified several pre-DC cell states distinguished by the expression of Ly6C, Ly6D, and SiglecH, and most of these cell states expressed moderate levels of Cd24, but importantly these pre-DCs were Cd11c+ MHCII–, whereas the Sox4+cDC cell state described here expressed comparable levels of MHCII to other cDC1 and cDC2 cell states, but lacked expression of Ly6C. We also found no evidence of enrichment of pre-DC gene signatures in Sox4+cDCs using genes listed by Schlitzer (Schlitzer et al., 2015) (data not shown). Other recent studies also identified a DC cell state sharing cDC and pDC gene expression (Brown et al., 2019; Leylek et al., 2019; Alcantara-Hernandez et al., 2017), and GSEA determined that Sox4+cDCs were similar to Siglech+cDCs and transitional DCs. An MHCII+ population expressing lineage-mixed features has also been identified in human blood and spleen (See et al., 2017). This population in humans expresses Cd33, Cx3cr1, Cd327, Cd123, Cd5, and Cd2. This population was shown to respond to CpG stimulation with secretion of TNFα and IL-12p40, and was able to induce CD4 T cell proliferation, whereas a pure pDC population was not. Furthermore, Villani et al. described a population of human DCs expressing Axl and Siglec6 (annotated as “AS DCs”), which...
expressed a continuum of cDC and pDC lineage-mixed gene signatures across a trajectory prediction and were able to induce T cell proliferation (Villani et al., 2017). We showed that the murine Sox4+ cDC cluster and the human “AS DCs” had a similar transcription factor expression profile, which was also shared with pDCs. It has been suggested that “AS DCs” are in fact aligned with the pre-DC population recently described by others (Amon et al., 2020), and Brown et al. showed that adoptive transfer of Siglech+ DCs leads to development of both cDC1 and cDC2 cells (Brown et al., 2019).

The simplified approach to classify cDC1 as CD8+ or CD103+ in mice is insufficient to predict cross-presenting function. Although not all CD8+ or CD103+ DCs possess cross-presentation ability (Kanazawa, 2007), the expression of XCR1 appears to be an exclusive cross-presentation feature (Bachem et al., 2012; Guillas et al., 2016). Batf3−/− mice are considered to lack cDC1 cells, but maintain a residual population of CD8+ cDCs. Our analysis determined that whereas Batf3−/− mice retained expression of CD8 and CD24 on cDC1-like cells, their expression of Xcr1/XCR1 and Clec9a was strongly diminished, although still elevated compared with classical cDC2 cells. These data suggest that the cDC1-like cells in Batf3−/− mice potentially retain suboptimal cross-presentation abilities, and this could explain some of the discrepancies described in the literature with respect to Batf3. We have previously reported that Batf3−/− mice were capable of rejecting neoantigen-expressing skin grafts, whereas a different mouse model lacking cDC1 cells (CD11cCre-Id2 flox/flox) was unable to do so (Chandra et al., 2017). Our interpretation of these data was that the residual population of CD8+ cDCs in Batf3−/− mice retained functionality to enable skin graft rejection. Intriguingly, in the same study, we showed that Batf3−/− mice were unable to mount a delayed hypersensitivity response and phagocytose dying cells, highlighting a threshold effect determining functions that are retained in Batf3−/− cDC1-like cells and others that are lost. Of note, the cDC1 compartment can be completely restored in Batf3−/− mice through bacterial infection or administration of IL-12, in which case Batf compensates for the lack of Batf3 (Tussiwand et al., 2012). This implies that the ability for Batf3−/− mice to mount a cDC1-dependent immune response is affected by the housing conditions of the animals, where Batf3−/− mice display more severe immunodeficiency when housed in a very clean facility (Mott et al., 2015). Similar to Batf3−/− mice, Etv6 deficiency also leads to a diminished cDC1 compartment with limited functionality. However, whereas Batf3−/− cDC1-like cells upregulated cDC2 features, Etv6−/− cDC1-like cells increased gene signatures of pDCs (Lau et al., 2018). Together these data support a hypothesis of phenotypic DC plasticity, and future studies are required to identify drivers of such plasticity.

With the extension to multimodal technologies including the incorporation of epitope-specific antibodies, holistic analysis of cell states can overcome some of the limitations associated with undetectable transcript expression and dropouts in unimodal scRNA-seq data. However, integration of epitope and transcript analysis highlights the need for caution when using barcoded antibodies, and verification of specific antibody staining and setting of thresholds is crucial. The need for antibody validation might pose a limitation when using extensive antibody panels in a range of hundreds as attempted by others. From an antibody panel of 10, we excluded 3 where specific binding was not clear. The current study also observed scenarios where transcript expression was better detected than epitopes (e.g., Sirpa/CD172A and Cdl4/CD4), and hence these antibodies did not add value to the analysis per se, but do act to support previous protein-based findings. This study provides the first CITE-seq dataset of murine splenic DCs of WT and Batf3−/− mice, which can serve to answer a variety of questions regarding steady-state DC heterogeneity and provides a road map to analyze DC cell state heterogeneity under inflammatory conditions in future studies.

**Limitations of the study**

Precise classification of dendritic cell populations has been proved to be a challenging task for the field. Although DCs can be broadly classified by cell surface protein expression and the expression of specific transcription factors, this approach is often insufficient for cell populations where the main distinguishing features are shared or are dynamically expressed. This is clear in the case of cells undergoing lineage conversion and/or differentiation. Furthermore, current clustering approaches cannot readily distinguish similar cell subtypes and cell states. This is visible in our data, where the identity of certain Sirpa+ cDCs as cell type or cell state remains elusive (e.g., Il4i1+ cDC and Sox4+ cDCs). To determine functional differences between these cDC cell states or cell subtypes would require extensive *in vitro* efforts. However, surface features exclusively expressed by these cells were not identifiable, making their isolation for further experiments not possible.
In this study, we combined high-resolution single-cell gene expression profiling with cell surface protein expression to track DC-specific features in the absence of a crucial DC transcription factor, Batf3. Despite the high resolution and throughput afforded by droplet-based scRNA-seq, the sensitivity of gene detection is limited to capturing only a portion of the transcriptome, in terms of numbers of genes detected and gene biotype. This is largely due to the short region that is sequenced using current 3' and 5' methods, and, for now, this limits the detection of population-specific transcript isoforms. Furthermore, current methods are designed to capture poly-A-tailed transcripts; developments in whole transcriptome profiling that capture a wider range of transcription products may reveal new and interesting markers.

One of the main limitations of CITE-seq is the potential for high levels of antibody background signal, which may be attributed to non-specific binding or high staining concentration. Although highly expressed proteins provide clear signals that enable robust cell type discrimination, classifying populations with low levels of expression remains a challenge. Our study was also limited to detecting proteins expressed at the cell surface; future techniques that also measure internal protein expression levels will allow investigation of protein families such as transcription factors. In addition, future studies in this direction would benefit from extensive antibody titration to achieve the best signal-to-noise ratio, because concentrations determined previously by other methods including flow cytometry may not be instructive.

**Resource availability**

**Lead contact**
Information and requests for resources may be directed to and will be fulfilled by the lead contact Janin Chandra (j.chandra@uq.edu.au).

**Materials availability**
This study did not generate new unique reagents.

**Data and code availability**
The Gene Expression Omnibus (GEO) accession number for the sequencing data reported in this article is GEO: GSE149544 (https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE149544).

**METHODS**
All methods can be found in the accompanying transparent methods supplemental file.

**SUPPLEMENTAL INFORMATION**
Supplemental information can be found online at https://doi.org/10.1016/j.isci.2021.102402.

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**AUTHOR CONTRIBUTIONS**
Conceptualization: S.W.L. and J.C.; data curation: S.L., I.R., S.B.A., A.N.C., and J.C.; formal analysis: S.W.L., I.R., S.K., S.M., S.B.A., A.N.C., M.Y., J.G., and J.C.; funding acquisition: I.H.F.; investigation: S.L., I.R., S.K., S.M., S.B.A., A.N.C., M.Y., J.G., and J.C.; methodology: S.L., I.R., S.K., S.M., S.B.A., A.N.C., M.Y., J.G., and J.C.; project administration: S.L. and J.C.; resources: I.H.F.; software: S.L., I.R., C.Z., and J.C.;
supervision: S.L., G.B., I.H.F., and J.C.; visualization: S.L., I.R., S.K., M.Y., J.G., and J.C.; writing – original draft: S.L., I.R., J.C.; writing – review & editing: S.L., I.R., S.K., M.Y., J.G., G.B., I.H.F., and J.C.

**DECLARATION OF INTERESTS**

The authors declare no competing interests. S.L. is currently an employee of Boehringer-Ingelheim.

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Absence of Batf3 reveals a new dimension of cell state heterogeneity within conventional dendritic cells

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Figure S1. Identification of cDCs co-expressing CD8 and CD11b, related to Figure 1. Splenocytes of C57BL/6 (WT) and Batf3-/- mice were T and B cell depleted and labelled with Hoechst stain and antibodies specific to MHCII, CD11c, CD8 and CD11b before acquisition by imaging flow cytometry. (A) CD11c+ MHCII+ DCs were pre-gated from Hoechst negative, in-focus singlets. The DCs subsets were further defined as CD11b- CD8+ cDC1s, CD11b+ CD8- cDC2s, and CD11b+ CD8+ lineage-intermediate (lin-int) cDCs. (B) Examples of CD11b, CD8, MHCII, and CD11c expression of lin-int cDCs. BF; brightfield.
Figure S2. Sorting purities of cDCs, related to Figure 2. Alive (live-dead stain aqua negative) single MHCII+CD11c+ cells were sorted from 6 C57BL/6 (WT) and 6 Batf3−/− T/B cell depleted splenocytes. Purity of the sorted cells was analysed and depicted >97%.
Figure S3. Additional single cell RNA sequencing information, related to Figure 2 and Figure 4. (A) Featureplots of Klrb1a, Cd19 and Cd3e to determine contamination by NK, B or T cells. (B) Ridgeplots of MHCII genes and Itgax (CD11c) across clusters. (C) Expression of mitochondrial and ribosomal subunit gene transcripts. (D) Heatmap of expression of apoptotic transcripts. (E) UMI counts across clusters. (F) Pearson correlations between RNA and antibody-derived tags (adt).
Figure S4. Expression of canonical DC genes across clusters, related to Figure 2. Heatmaps of canonical DC genes for WT (blue) and Batf3−/− (orange) clusters.
Figure S5. Residual mitotic cDC1-like cells in Batf3-/- mice increase expression of cDC2 features, related to Figure 5. Gene expression of cDC1 and cDC2 features was compared between WT and Batf3-/- of mitotic cDC1 (C6) cells (upper panels), or between Batf3-/- of mitotic cDC1 (C6) and Batf3-/- of mitotic cDC2 (C5) cells (lower panels).
## TRANSPARENT METHODS

### Key resources table

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Antibodies**      |        |            |
| Anti-mouse CD11c (PE-Dazzle) | Biolegend | Cat#117348; RRID: AB_2563655; Clone: N418 |
| Anti-mouse I-A/I-E (APC-Cy7) | Biolegend | Cat#107628; RRID: AB_2069377; Clone: M5/114.15.2 |
| Anti-mouse/human CD11b (Alexa Flour 700) | Biolegend | Cat#101222; RRID: AB_493705; Clone: M1/70 |
| Anti-mouse CD8a (PE) | eBioscience | Cat#12-0081-82; RRID: AB_465530; Clone: 53-6.7 |
| Anti-mouse/human CD11b (Brilliant violet 510) | Biolegend | Cat#101245; RRID: AB_2561390; Clone: M1/70 |
| Anti-mouse TCRb (Biotin) | Biolegend | Cat#109204; RRID: AB_313427; Clone: H57-597 |
| Anti-mouse CD19 (Biotin) | Biolegend | Cat#115504; RRID: |
| Antibody Description                        | Manufacturer | Cat#                                 | RRID                | Clone   |
|--------------------------------------------|--------------|--------------------------------------|---------------------|---------|
| Anti-mouse CD45 (BUV615)                    | BD Biosciences | Cat#751170; RRID: AB_2875194; Clone: 6D5 |
| Anti-mouse CD3 (BV750)                      | Biolegend    | Cat#100225; RRID: AB_10900444; Clone: 30-F11 |
| Anti-mouse CD19 (BV570)                     | Biolegend    | Cat#115535; RRID: AB_10933260; Clone: 17A2 |
| Anti-mouse Ly6G (BUV737)                    | BD Biosciences | Cat#741813; RRID: AB_2871151; Clone: B4 |
| Anti-mouse PDCA1 (APC)                      | Biolegend    | Cat#127015; RRID: AB_1967101; Clone: 927 |
| Anti-mouse CD11b (APC-R700)                 | BD Biosciences | Cat#564985; RRID: AB_2739033; Clone: M1/70 |
| Anti-mouse Ly6C (FITC)                      | Biolegend    | Cat#128005; RRID: AB_1186134; Clone: HK1.4 |
| Anti-mouse I-A/I-E (BV711)                  | Biolegend    | Cat#107643; RRID: AB_2565976; |
| Anti-mouse CD11c (BV421) | Biolegend | Clone: M5/114.15.2 |
|-------------------------|-----------|---------------------|
| Cat# 117329; RRID: AB_10897814; Clone: N418 |
| Anti-mouse CD24 (BV605) | Biolegend | Cat# 101827; RRID: AB_2563464; Clone: M1/69 |
| Anti-mouse XCR1 (BV650) | Biolegend | Cat# 148220; RRID: AB_2566410; Clone: ZET |
| Anti-mouse CD33 (PE) | Thermo Fisher Scientific | Cat# 12-0331-80; RRID: AB_2637178; Clone: 9A11 |
| Anti-mouse CD172A (PerCP eFluor710) | Thermo Fisher Scientific | Cat# 46-1721-82; RRID: AB_10804639; Clone: P84 |
| Anti-mouse CD4 (BUV496) | BD Biosciences | Cat# 612952; RRID: AB_2813886; Clone: GK1.5 |
| Anti-mouse CD11c (Totalseq-A0106, GTTATGGACGCTTTGC) | Biolegend | Cat#: 117355; RRID: AB_2750352; Clone: N418 |
| Anti-mouse/ human CD11b (Totalseq-A0014, TGAAGGCTCATTGT) | Biolegend | Cat#: 101265; RRID: |
| Antibody Description | Manufacturer | Cat#/RRID/Clone |
|----------------------|--------------|----------------|
| Anti-mouse CD172a (Totalseq-A0422, GATTCCCTTGTAGCA) | Biolegend | AB_2734152; Clone: M1/70 |
| Anti-mouse CD370 (Totalseq-A0556, AACTCAGTTGTGCCG) | Biolegend | Custom-built; Part #: 96461; Clone: 7H11 |
| Anti-mouse CD4 (Totalseq-A001, AACAAGACCCCTTGAG) | Biolegend | AB_2800670; Clone: P84 |
| Anti-mouse CD8a (Totalseq-A002, TACCGTAATAGCGT) | Biolegend | AB_2749956; Clone: RM4-5 |
| Anti-mouse F4/80 (Totalseq-A0114, TTAACCTCAGCCCGT) | Biolegend | AB_2749986; Clone: BM8 |
| Anti-mouse Siglec H (Totalseq-A0119, CCGCACCTACATTAG) | Biolegend | AB_2750537; Clone: 551 |
| Anti-mouse, rat XCR1 (Totalseq-A0568, TCCATTACCCACGTT) | Biolegend | AB_2783120; Clone: ZET |
| Anti-mouse CD24 (Totalseq-A0212, TATATCTTTTGCCGCA) | Biolegend | AB_2750380; Clone: M1/69 |
| Biological Samples     |               |           |
|-----------------------|---------------|-----------|
| N/A                   |               |           |

| Chemicals, Peptides, and Recombinant Proteins |               |           |
| Collagenase D          | Roche         | 11088858001 |
| DNase                  | Roche         | 11284932001 |

| Critical Commercial Assays |               |           |
| EasySep™ Mouse Streptavidin RapidSpheres™ Isolation Kit | EasySep | Cat#19860 |
| LIVE/DEAD™ Fixable Aqua Dead Cell Stain Kit | Invitrogen | Cat#L34965 |
| Viability stain FVS | BD Biosciences | Cat#566332 |

| Deposited Data |               |           |
| GSE149544      | NCBI GEO      |           |

| Experimental Models: Cell Lines |               |           |
| N/A                             |               |           |

| Experimental Models: Organisms/Strains |               |           |
| C57BL/6                         | Animal Resources Centre | Product code: A2KB |
| **Batf3-/-** | Jackson Laboratories | Stock#013755 |
|--------------|----------------------|-------------|

**Oligonucleotides**

| AF647-dT [5’-5Alex647N-T_{30}-3’] | IDT |
|-----------------------------------|-----|

**Recombinant DNA**

| N/A |
|-----|

**Software and Algorithms**

| IDEAS | Life Science Research |
|-------|------------------------|
| Kaluza | Beckman Coulter |

For Kaluza: [https://www.beckman.com.au/flow-cytometry/software/kaluza; RRID:SCR_016182](https://www.beckman.com.au/flow-cytometry/software/kaluza; RRID:SCR_016182);

| GraphPad Prism | Prism version 7 |
|----------------|-----------------|
| [https://www.graphpad.com/scientific-software/prism; RRID:SCR_002798](https://www.graphpad.com/scientific-software/prism; RRID:SCR_002798) |

| bcl2fastq | 2.2.0 |
|-----------|-------|

| Cellranger | 3.0.2 |
|------------|-------|

| Seurat     | 3.0.5 |
|------------|-------|

| SingleR    | 1.0 |
|------------|-----|

| Nebulosa   | 0.99.92 |
|------------|---------|
|     |     |
|-----|-----|
| AUCell | 1.12.0 |
| Velocyto | 0.17.15 |
| scvelo | 0.2.1 |
| Other |     |
| N/A |     |
Experimental models and subject details

Mice

C57BL/6 mice (WT) were obtained from the Animal Resources Centre (Perth, Western Australia, Australia). Batf3-/- mice were purchased from Jackson Laboratories (Bar Harbor, ME, USA) and bred in house. All mice were kept under specific pathogen-free conditions at the Biological Research Facilities of Translational Research Institute. All animal procedures and experiments were performed in compliance with the ethical guidelines of the National Health and Medical Research Council of Australia, with approval from the IMVS Animal Ethics Committee and the University of Queensland Animal Ethics Committee (UQDI/452/1)

Method details

Cell isolations

Spleens were removed from CO2-euthanised mice and injected with 0.5mg/ml collagenase D (Roche, 1108858001) and 0.1 mg/ml DNase (Roche, 11284932001) diluted in PBS 10% FCS for 30 min at 37° C. The tissue was mechanically disrupted by straining through a 70 mm cell strainer (BD Falcon) into 50 ml falcon tubes and washed with PBS to a total volume of 15 ml. Single cell suspensions were centrifuged at 350g for 7 min, followed by red blood cell lysis using ammonium chloride potassium (ACK) buffer for 2 min on ice and subsequent wash with PBS. Cell pellets were resuspended in 2.5 ml PBS buffer. Cells were counted using TC20 automated cell counter (Bio-Rad) by staining with trypan blue (Sigma-Aldrich) for dead cell discrimination.

T and B cell depletion

To enrich splenic dendritic cells, T and B cells were depleted from splenic single cell suspension using EasySep Mouse Streptavidin RapidSpheres (StemCell Technologies) in accordance to the manufacturer’s protocol. Biotin conjugated antibodies used for depletion were TCRb (Biolegend; clone H57-597; dilution 1:200) and CD19 (Biolegend; clone 6D5; dilution 1:200). In detail 1 x 10^8 cells were resuspended in 1 ml FACS buffer in round bottomed 5 mL tubes (BD Falcon) and incubated with 5 µl of each antibody for 10 minutes at room temperature. 75 µl of RapidSpheres magnetic beads were added followed by 2.5 minutes of incubation. The tube was placed in an EasySep magnet (STEMCELL Technologies) and incubated for 2.5 minutes before the enriched cells were carefully decanted into a new tube.

Flow cytometry analysis
Approximately 1 million cells from splenic single cell suspension was added to 5 mL round bottomed tubes (BD Falcon) for unstained and single stain controls. For samples with multiple stains approximately 4 million cells were added to each tube. 1 ml of PBS was added to all tubes before centrifuging at 350g for 5 min. To exclude dead cells, samples were treated with live/dead stain (Invitrogen, dilution 1:250 or BD Bioscience 1:500) diluted in PBS for 20 minutes on ice followed by PBS wash and centrifugation. Cells were incubated with FC block (BD Pharmingen; dilution 1:100) at 4°C for 10 minutes. Cells were incubated with anti-mouse monoclonal antibodies in FACS buffer (2% HI FBS, 1x PBS, 2 mM EDTA) for 20 minutes on ice. Cells were washed with FACS buffer and centrifuged, followed by resuspension in 400 µl FACS buffer. BD LSR Fortessa X20 and BD Symphony flow cytometers were used to acquire cells and Kaluza software (Beckman Coulter) was used for compensation and analysis.

Imaging flow cytometry

After T and B cell depletion the cells were incubated with Hoechst dye and FC block (BD Pharmingen; dilution 1:100) at 4°C for 10 minutes. mAbs against CD11c (Clone: N418; dilution 1:50), MHCII (Clone: M5/114.15.2; dilution 1:100), CD11b (Clone: M1/70; dilution 1:50), and CD8 (Clone: 53-6.7; dilution: 1:100) were used and the samples were incubated at 4°C for 30 minutes. The cells were acquired on an AMNIS ImageStreamX MkII (Luminex Corporation) and analysed using IDEAS software (Life Science Research).

Flow cytometric activated cell sorting

For CITE sequencing, 2 x 10^8 of pooled splenocytes from six C57BL/6 and six Batf3−/− mice respectively were resuspended in 1 mL FACS buffer and used for downstream analysis. Following T and B cell depletion (described above) cells were incubated with FC block (BD Pharmingen; dilution 1:100) simultaneously with live-dead Aqua stain (Invitrogen, dilution 1:250). Cells were subsequently stained with anti-mouse monoclonal antibodies against MHCII and CD11c in accordance with the staining protocol described above. Live MHCII^+CD11c^+ cells from each genotype were sorted into 1.5ml tubes (Eppendorf) containing 500μl 100% FCS using droplet-based sorters. Sorted cells were centrifuged at 350g for 5 minutes.

Incubation of barcoded antibodies

Sorted DC cell pellets were resuspended in 50 µl FACS buffer and incubated with 5 µl FC block for 10 minutes on ice, followed by PBS wash and centrifugation. We added 0.5 µg of each TotalSeq-A antibody (Biolegend) as an antibody pool to each of the samples. Cells were transferred to 1.5 mL tubes (Eppendorf) and incubated for 30 minutes on ice, followed by wash using 2% BSA/ 0.01%
Tween in PBS and centrifugation at 350g for 5 minutes at 4°C. DCs were resuspended in FACS buffer, cell viability and numbers were estimated by manual counting using trypan blue.

**Single cell RNA sequencing**

Sample preparation and 10x Chromium pipeline was performed in accordance with the manufacturer’s protocol. To target 12,500 cells per sample ~25,000 cells of each genotype were loaded onto the chip for Chromium Single Cell Gel Bead and Library Kit v3 (10x Genomics) to account for a ~50% cell loss during microfluidics processing. The 10x Genomics User Guide was followed for GEM generation and barcoding, cDNA amplification and library construction. Eleven cDNA amplification cycles were performed, and one quarter of the cDNA was used as input for library construction. Library construction for CITE-seq was conducted in accordance with Totalseq-A antibody manufacturer protocol (Biolegend, v.3). Final indexing PCR was performed in a C1000 Touch thermal cycler with a Deep Well Reaction Module (Bio-Rad). For gene expression and CITE-seq libraries 11 and 10 SI-PCR cycles were used, respectively. Using the High Sensitivity DNA kit on Agilent BioAnalyser 2100 the libraries were quantified (Agilent, 5067-4626). Equimolar ratios were used to pool gene expression libraries, and of the total sequencing pool concentration 10% were spiked in cell hashing libraries. Quantification of the final pool was conducted by qPCR using KAPA Library Quantification Kit in combination with the Life Technologies Viia 7 real time PCR instrument (KAPA Biosystems, KK4824). Illumina NextSeq-500 was used for sequencing the denatured libraries using a 150 cycle High-Output Kit. This included Read 1 (28 bp) supplying the cell’s barcode (16nt) and UMI (12nt), i7 (8bp) for sample index and Read 2 (111 bp) to capture the 3’ sequence of the transcripts. In total, two sequencing runs were performed to obtain an average read depth of ~30,000 reads per cell.

**Quantification and statistical analysis**

**Single cell RNA sequencing data analysis**

The Illumina sequencing output was processed using the 10x Genomics Cell Ranger 3’ gene expression and feature barcoding pipeline (v3.0.2). Illumina bcl2fastq (v2.2.0) was used to demultiplex the raw Illumina reads and generate FASTQ files for individual RNA and ADT libraries. Cellranger count was used to map the sequencing reads to the GRCh38 (mm10) mouse reference genome, and the aggregate function was used to combine the libraries into a single matrix and correct for differences in sequencing depth. The aggregated gene expression and antibody count data, comprising wildtype and knockout libraries were further processed and analysed using Seurat (v3.0.5) (Stuart et al., 2019). Filtering and quality control were performed to remove outlier cells and
transcripts. Transcripts were retained if they were expressed in at least 3 cells, and a preliminary filtering of cells was done to remove cells with fewer than 200 transcripts. Further filtering was done based on number of transcripts in each cell with a lower threshold of 500 transcripts and the maximum number of transcripts per cell set to 5000. Cells with mitochondrial or ribosomal reads accounting for more than 20% and 50% respectively, were excluded from downstream analysis. Cell-cell gene expression variation was normalised using log transformation and the antibody counts were normalised using centered log-ratio (CLR), each followed by scaling as implemented in the Seurat pipeline. To reduce the dimensionality, we performed principal component analysis (PCA) and the cut-off was set to where an ‘elbow’ was observed in the curve. The first 16 PCs were used to construct a k-nearest neighbour graph using the FindNeighbors function in Seurat. Similar cells were then grouped together using the Louvain algorithm in the FindClusters function using the default resolution of 0.5 (Stuart et al., 2019). Cells were visualised in a two dimensional space using non-linear dimensional reduction via uniform manifold approximation and projection (UMAP) implemented in the Seurat pipeline.

**Multi-modal cell type classification**

Cell identities were first predicted by SingleR analysis using the Immunological Genome Project (ImmGen) database as a reference (https://github.com/dviraran/SingleR). Single cells and clusters were annotated based on gene expression markers of immune cell subsets described in ImmGen. Gene set enrichment analysis (GSEA) was performed using DC lineage-specific gene signatures outlined and referenced in Suppl. File 1 and using the AUCell package. Populations were further identified using known markers characteristic to cDC1, cDC2 and pre-cDCs. Unbiased population identification was performed by differential gene expression analysis. Using the FindAllMarkers algorithm implemented in Seurat, we obtained positive gene markers, expressed in at least 25% of the cells at a minimum of 0.25-fold difference (log-scale) between the cluster and the remainder of the dataset. The top 10 differentially expressed genes were visualised in a heatmap. Clusters were further distinguished by differential expression between specific populations and expression of DC lineage-specific gene markers. Cell cycle phase prediction was performed in Seurat using a dataset of cell-cycle-specific gene markers (Kowalczyk et al., 2015). Cells were assigned to the G2/M and S phases based on expression of gene markers associated to these phases. Cells not expressing markers of either G2/M or S phases were assigned to G1.

**RNA velocity**

Spliced and unspliced transcripts were identified for each cell using velocyto in combination with the position sorted BAM files produced by cellranger for the wildtype and Batf3<sup>−/−</sup> samples. The
individual velocity results for each sample were merged and the cell barcodes were matched to only contain the cell barcodes present in the Seurat object used for the main expression analysis. Next, a loom file was created from the full Seurat object, and the merged velocity data was processed and analysed using scvelo.

Protein expression profiles using oligo-tagged antibodies (CITE-seq)

To account for non-specific binding of the TotalSeq-A antibodies, we defined antibody-specific thresholds using flow cytometry analysis of the TotalSeq-A antibodies hybridised to a fluorochrome-conjugated oligo-dT probe and incubated with C57BL/6 splenocytes together with fluorochrome conjugated antibodies against CD11c and MHCII (Fig. S4B). After pre-selection of DCs, the percentage of positive staining of TotalSeq-A antibodies was translated into an adt bin value, which was used as cut-off point for the adt data. All values below the antibody-specific cut off point were set to zero. The bin-corresponding adt threshold value determined in C57BL/6 was subsequently applied as cut-off in the Batf3-/ data.

Statistical analysis

Analysis of data was performed using either unpaired two-tailed Student’s t test or one-way ANOVA with Tukey’s multi-comparison post-hoc test with a 95% confidence interval (Prism, GraphPad Software). *p<0.05; **p<0.01; ***p<0.001; ****p<0.0001. Information regarding number of replicates, sample size, significance test, value and meaning of n for each experiment are included in each figure legend. CITE-seq experiment was performed once.

Data availability

The accession number for the sequencing data reported in this paper is GSE149544.
SUPPLEMENTAL REFERENCES

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