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## In situ maturation and tissue adaptation of type 2 innate lymphoid cell progenitors

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### Highlights:

- A single cell atlas of BM and lung ILCs of healthy, infected and parabiotic mice
- Identification of tissue-associated ILC progenitors in neonatal and adult lung
- Cells recruited from BM generate the entire spectrum of ILC2s in infected lungs
- Local cues imprint the phenotypes of ILC2s differentiating in the adult lung

eTOC blurb: To investigate how ILCs are locally maintained, Zeis et al. generated a single cell atlas of lung ILCs and tracked I18r1+ progenitor and effector ILC2s. Their work identifies tissue-resident and circulating ILCs, and highlights *in situ* differentiation and tissue adaptation as a mechanism of ILC renewal and phenotypic diversification.

## Summary

Innate lymphoid cells (ILCs) have ample roles in tissue homeostasis, inflammation and repair. ILCs are generated early during ontogeny and persist predominantly as tissue-resident cells. How ILCs are maintained and renewed remains unclear. Here, we generated a single cell atlas of lung ILC2s and found that  $\text{Il18r1}^+$  ILCs comprise circulating and tissue-resident ILC progenitors (ILCP) and effector-cells with heterogeneous expression of *Tcf7*, *Zbtb16* and *CD103*. We revealed a continuous differentiation trajectory and validated the potential of  $\text{Il18r1}^+$  ILCPs to generate ILC2s. Upon helminth infection, recruited and BM-derived cells generated the entire spectrum of ILC2s in parabiotic and shield chimeric mice, consistent with their potential role in the renewal of tissue ILC2s. Our work identifies local ILCPs and highlights their *in situ* differentiation and tissue adaptation as a mechanism of ILC maintenance and phenotypic diversification. Our results suggest that local tissue niches, rather than progenitor origin, or the developmental window during ontogeny, dominantly imprint ILC phenotypes in adult tissues.

## Introduction

Innate lymphoid cells (ILCs) have been identified in many different tissues where they contribute to immune surveillance, tissue homeostasis and repair, and barrier function. ILCs are found in the bone marrow, in secondary lymphoid organs and, most prominently, in non-lymphoid tissues including mucosal barriers. The size and subset composition of the local pool of ILCs can exhibit dramatic differences between organs. Despite considerable progress in ILC biology, ILC development and their adaptation to heterogeneous tissue environments remain poorly understood. ILCs are believed to seed tissues during embryonic development (Bando et al., 2014), where they expand and differentiate during the first weeks of life (Saluzzo et al., 2017; Schneider et al., 2019). In adult tissues, ILCs persist as tissue-resident cells. Originally established in parabiotic mice (Gasteiger et al., 2015; Huang et al., 2018; Moro et al., 2016; Peng et al., 2013), this concept is supported by analyses of human ILCs in transplanted livers and intestines (Cuff et al., 2016; Weiner et al., 2017). However, only a fraction of ILC2s labelled in the neonatal window persists in adult tissues (Schneider et al., 2019), reinforcing the question of how tissue-resident ILCs are maintained and replenished. Local pools of tissue-ILCs may receive input from distant cellular sources during inflammation, and local changes in ILC subsets and abundance correlate with a number of inflammatory diseases (Mazzurana et al., 2018). Consistently, murine ILC2s can traffic between organs during helminth infection (Gasteiger et al., 2015; Huang et al., 2018). It remains unclear whether recruited cells have a dedicated function, and whether they replenish the local pool of resident cells. Thus, ILCs detected in a given tissue may be generated from different cellular sources and at different times during ontogeny. How the local pools of ILCs are being maintained and renewed is currently unknown. In particular, we lack a detailed understanding of the defined stages of differentiation of tissue ILCs, the phenotype of “immature” ILCs, as well as the sites and tissue niches where maturation occurs.

To address these questions, we generated a comprehensive single-cell atlas of ILC populations in the BM, and neonatal and adult lung at steady-state, as well as during helminth infection in wild-type, parabiotic and shield chimeric mice. We identified immature stages of ILCs, inferred their differentiation trajectories, and validated their developmental potential *in vitro* and *in vivo*. We show that subsets of  $\text{Il18r1}^+\text{ST2}^-$  lung ILC2s express ILC progenitor genes, become proliferative, and gradually mature in the lung. Analysis of *Nippostrongylus brasiliensis* (Nb) infected parabiotic and shield chimeric mice suggests a contribution of these cells to the renewal of tissue ILC2s upon local inflammation. Furthermore, our study establishes that cells trafficking to the adult lung, particularly from the bone marrow, can give rise to the full spectrum of phenotypes of resident ILC2 subsets that have been generated in distinct waves during ontogeny, implying a dominant role of tissue imprinting on ILC phenotypes. Our work therefore identifies local progenitors as tissue ILCs and highlights tissue adaptation and *in situ* differentiation as a mechanism of ILC maintenance and phenotypic diversification. In addition, the single-cell RNA-seq datasets (<http://murine-ilc-atlas.ie-freiburg.mpg.de/>) represent a rich, interactive resource for further exploration of ILC biology.

## Results

### Heterogeneity of ILC2s in healthy lung tissue

To characterize the full phenotypic spectrum of adult lung ILC2s in an unbiased manner we FACS-purified lineage-negative ( $\text{Lin}^-$ )  $\text{CD45}^+\text{Il7ra}^+\text{Thy1}^{\text{low}}\text{NK1.1}^-$  “pan-ILCs” for scRNA-seq, excluding NK cells and ILC1, but not ILC3, which are relatively rare in the lung (Figure 1A, B and S1A). As committed ILC progenitors (ILCPs) and mature tissue ILCs express *Il7ra* (Constantinides et al., 2014; Klose et al., 2014), we expected that this sorting strategy samples the entire differentiation tree of ILC2s including putative early differentiation stages. Clustering cells using RaceID3 (Herman et al., 2018) yielded 12 clusters with >20 cells (Figure 1C). Except for clusters 1, 5 and 11, all cells highly expressed *Gata3* and *Il1rl1* (Figure 1D, E), which are markers of mature ILC2s in the lung. We detected different effector ILC2 sub-types such as cluster 10, up-regulating the neutrophil recruiting factor *Cxcl2*, *Il2*, *Klf2* and

*Cd69*, or cluster 12, up-regulating *Areg*, *Calca* and *Csf2* (Figure 1D, 1E and S1B). In addition, clusters 6 and 7 shared the up-regulation of *Cxcl2*, *Areg*, *Il13*, *Il5* and *Calca*, but were distinct with regard to expression of *Calca* and *Csf2* versus *Il2*, *Cxcl2*, and *Pdcd1*, respectively. Besides these effector-like ILC2s resembling subsets of ST2<sup>+</sup> natural ILC2s (nILC2s) (Nagashima et al., 2019; Wallrapp et al., 2017) we identified clusters 1, 5 and 11 characterized by low expression of *Gata3* and *Il1rl1*. Only a small fraction of cells in these clusters expressed low levels of ILC2 effector genes such as *Cxcl2*, *Areg* and *Il13* or *Il5*. We also detected cells up-regulating *Rorc* and *Tbx21* within cluster 5 and 11, respectively (Figure 1E, S1C), indicating mixed lineage contributions, as previously observed (Wallrapp et al., 2017). Few cells in clusters 1 and 5 expressed T cell markers *Cd3e* and *Trdc*, akin to reported subsets of murine and human ILCs (Björklund et al., 2016; Robinette et al., 2015), although a contamination of T cells with downregulated surface CD3ε cannot be ruled out (Figure S1A). Interestingly, cells in clusters 1, 5 and 11 up-regulated *Il18r1*, *Thy1*, *Cd7*, *Tcf7*. The Notch target *Tcf7* is expressed in ILCPs, and CD7 in circulating human ILC precursors (Constantinides et al., 2014; Harly et al., 2019; Ishizuka et al., 2016; Lim et al., 2017). Of note, a subset of cells in clusters 1, 5 and 11 expressed *Zbtb16*, a marker of committed ILCPs (Ishizuka et al., 2016; Klose et al., 2014; Lim et al., 2017) and few of these cells co-expressed combinations of *Rorc*, *Tbx21* and *Il17rb*, consistent with a mixed lineage-potential (Figure S1C). In addition, we observed up-regulation of *Igfr1* in clusters 1 and 5, a receptor suggested to be critical for recently identified ILC3-progenitors in neonatal lung (Oherle et al., 2020).

To investigate if these cells represent early differentiation stages of tissue ILCs we analysed the developmental relationship of lung ILC subsets with StemID2, which connects related clusters to construct a lineage tree (Figure 1F). This analysis predicted a differentiation trajectory connecting *Il18r1*+*Gata3*<sup>lo</sup>*Il1rl1*<sup>lo</sup> cells (clusters 1, 5, 11), to ILC2s with increasing levels of *Il1rl1* and intermediate expression of effector cytokines (clusters 9, 8, 4) and to mature effector clusters (10, 12, 6, 7 and 2) (Figure 1F). Elevated transcriptome entropy for clusters 1 and 5 suggested a more naïve state compared to the other clusters (Grün et al., 2016). After ordering cells along the trajectory starting at cluster 5 with the highest expression of known progenitor marker genes, and extending via clusters 11, 9, 8, and 4 towards cluster 10, pseudo-temporal gene expression profiles were grouped into distinct co-expression modules (Methods, Figure S1D, E). While the *Cxcl2*-expressing cluster 10 represents one possible endpoint of the trajectory, other mature clusters (2, 6, 7, and 12) correspond to alternative endpoints branching out from cluster 4 (Figure S1F).

Along the trajectory, we observed gradual up-regulation of *Gata3*, *Il1rl1*, *Klrg1* and *Bcl11b*, which has been shown to be up-regulated during ILC2 differentiation (Califano et al., 2015; Yu et al., 2015). *Il18r1* and *Cd7* were down-regulated along with ILC progenitor genes such as *Zbtb16*, *Tcf7*, *Maf* and *Ikzf2* (Figure 1G) (Harly et al., 2019; Ishizuka et al., 2016; Seehus et al., 2015). These inferred gene-expression dynamics suggested that mature *Gata3*<sup>hi</sup>*Il1rl1*<sup>hi</sup> ILC2s can differentiate from progenitor-like *Il18r1*<sup>+</sup>*Gata3*<sup>lo</sup>*Il1rl1*<sup>lo</sup> ILCs (clusters 1, 5, and 11) in the lung. FACS analysis confirmed that *Il18r1*<sup>+</sup>ST2<sup>-</sup>RORγt<sup>-</sup>Klrg1<sup>-</sup> cells, hereafter referred to as *Il18r1*<sup>+</sup>ST2<sup>-</sup> cells, represented about 2% of Lin<sup>-</sup>Il7ra<sup>+</sup> ILCs (Figure 1H, I), and were also detected in lungs of T cell deficient RAG<sup>-/-</sup> and TCRβ<sup>-/-</sup> mice (Figure S1G and data not shown). Consistent with the gene signatures predicted for cluster 11 (Figure 1D), a high fraction of these cells was positive for the proliferation marker Ki67, suggesting their capacity for local expansion or self-renewal (Figure 1J). Together, these observations indicated that *Il18r1*<sup>+</sup>ST2<sup>-</sup> cells comprise immature differentiation stages of ILC2s that are enriched in the lung, where they may undergo local maturation. Consistently, when we analyzed lung ILC2s by dividing them into three subsets that were predicted to map successively along the inferred maturation trajectory (i.e., *Il18r1*<sup>+</sup>ST2<sup>-</sup>; *Il18r1*<sup>+</sup>ST2<sup>+</sup> and *Il18r1*<sup>-</sup>ST2<sup>+</sup> cells), we found that cells downregulated Ki67, *Il18r1*, *Il17rb* and *Tcf7* along this trajectory, while gaining expression of CD25, ST2, *Gata3* and *Il5* protein, which all characterize mature ILC2s in the lung (Figure 1J- L).

To demonstrate that clustering and trajectory inference were not driven by proliferation associated programs, we removed proliferation associated genes prior to analysis (Methods) and obtained a similar trajectory connecting *Il18r1*<sup>+</sup>*Gata3*<sup>lo</sup>*Il1rl1*<sup>lo</sup> cells to effector-like ILC2s (Figure S1H-K). Together, these observations suggested that *Il18r1*<sup>+</sup>ST2<sup>-</sup> cells comprise early immature differentiation stages of ILC2s that are enriched in the lung, where they may locally mature.

Expression of *Il18r1*, *Tcf7* and *Cd7* as well as effector genes *Il5*, *Il13*, and *Il17a* has recently been reported for ILC2s in the skin (Bielecki et al., 2018; Kobayashi et al., 2019; Ricardo-Gonzalez et al., 2018), raising the possibility that  $Il18r1^{+}ST2^{-}$  cells comprise heterogeneous ILC subtypes. To better characterize  $Il18r1^{+}$  ILCs across tissues, we integrated our data with lung and skin single-cell transcriptomes from Ricardo-Gonzalez et al. and tested for the presence of discrete progenitor and effector states (Methods, Figure S2A). We found that *Il18r1* expressing lung ILCs largely clustered separately from mature skin ILC2s expressing ILC2 effector genes (Figure S2A, B). However, a fraction of skin ILCs clustered together with immature  $Il18r1^{+}$  lung ILCs and up-regulated *Igfl1r*, *Tcf7*, *Zbtb16* and *Pdcd1*, suggesting an immature progenitor phenotype (Figure S2A-C). It is important to note, that lung and skin-ILCs were sorted as  $Il5$ -mRNA-reporter<sup>+</sup> cells by Ricardo-Gonzalez et al., and therefore this analysis may underestimate the frequency of progenitor-like cells. Reminiscent of these distinct phenotypes in skin, we detected  $CD103^{+}Il18r1^{+}$  cells in the lung, a fraction of which also expressed cytokines *Il5*, *Il13* and *Il17a* upon *in vitro* re-stimulation (Figure S2D, E). These combined analyses reveal a previously unappreciated heterogeneity of  $Il18r1^{+}ST2^{-}$  ILC subtypes in skin and lung.

### **$Il18r1^{+}ST2^{-}$ ILCs in neonatal and adult lung share similarities with $Il18r1^{+}$ ILC progenitors in the bone marrow**

We next asked how the  $Il18r1^{+}ST2^{-}$  putative ILC2 progenitors from adult lung compare to known ILC progenitors. ILC2s are generated in different developmental windows during ontogeny, and cells emerging during neonatal life can persist in the adult lung (AL) (Schneider et al., 2019). To test whether corresponding progenitor populations are present in the neonatal lung (NL), we sequenced  $ST2^{+}$  and  $ST2^{-}$  ILCs from postnatal day 4 (P4) lung (Figure S3A), and performed clustering analysis (Figure 2A, 2B). Within the  $ST2^{-}$  compartment, cluster 9 exhibits a naïve ILC signature akin to AL clusters 1, 5, and 11, with up-regulation of *Il18r1*, *Tcf7*, *Zbtb16* and *Pdcd1*, as well as intermediate levels of *Gata3* and high expression of *Il17rb* (Figure 2C, 2D), suggesting predominant ILC2 bias. Cluster 4, 6, 7 and 12 share expression of *Il18r1* and *Tcf7*, and resemble ILC1s or NK cells with expression of *Tbx21*, *Nkg7* and *Klrd1* (Figure 2D). FACS analysis confirmed the heterogeneity of  $Il18r1^{+}ST2^{-}$  cells and the presence of  $Il18r1^{+}Tcf7^{+}$  cells that expressed *Zbtb16* (PLZF), c-Kit, PD1,  $\alpha4\beta7$ , and *Ki67* to variable degrees in both NL and AL (Figure 2E, F and S3B-D). These cells shared expression of many genes with  $Il18r1^{+}Icos^{+}$  BM ILCs that we further characterized by scRNA-Seq (Figure 2G, H) and cells that have recently been refined as  $Id2^{+}$  BM ILCPs (Wu et al. (2019)). By integrating our data with published BM ILC scRNA-seq data (Walker et al., 2019), we confirmed that BM  $Il18r1^{+}Icos^{+}$  cells contain ILCPs with ILC2 bias (Figure S3E-H). Accordingly, cells of  $Il18r1^{+}Icos^{+}$  BM cluster 1 up-regulated *Tcf7*, *Zbtb16*, *Pdcd1* and *Il17rb* (Figure 2H), reminiscent of NL cluster 9. While  $Il18r1^{+}$  NL clusters lacked expression of *Cd3* genes, BM clusters 4 and 5 as well as AL progenitor clusters 1 and 5 up-regulated these genes. In addition, we resolved clusters 2 and 7, up-regulating *Mki67* (Figure 2h). These data confirm the recently described phenotypic spectrum of BM ILC progenitors and reveal that corresponding populations with a similar heterogeneity exist in NL and AL (Figure 1, 2A-F). Circulating human ILC progenitors reminiscent of the populations we identified here were described recently (Lim et al., 2017). The  $IL7R^{+}c-KIT^{+}$  bulk population, which possibly include  $CRTH2^{-}$  cells biased toward ILC2 lineage differentiation (Nagasawa et al., 2019) displays the transcriptional profile of *IL18R1*, *TCF7*, *CD7* and *ZBTB16* (Lim et al., 2017). As expression of these genes was not tested previously at the protein level, we analyzed human ILCs in peripheral blood and lungs and identified a population of  $IL7R^{+}IL18R1^{+}TCF1^{+}$  (encoded by the *TCF7* gene) cells that expressed GATA3 while lacking markers of mature NK cells, ILC1, ILC2 and ILC3 (i.e., EOMES, T-bet,  $CRTH2$ , and  $NKp44$ , respectively) (Fig. 2I-K, S3I). Again, similar to cells in the murine lung, we observed graded expression of *IL18R1*, *TCF1*, *GATA3*, *c-KIT*, *ICOS* and *CD103*, reminiscent of the described spectrum of murine  $Il18r1^{+}$  ILCs in neonatal and adult lungs, and BM ILCPs (Walker et al., 2019; Xu et al., 2019).

Together, these observations raise the possibility that we have identified a murine counterpart to ILC progenitors present in human lungs, reinforcing the question about the developmental potential of these cells.

## II18r1<sup>+</sup> ILCs comprise immature cells that give rise to Gata3<sup>hi</sup> ILC2s in the lung

Consistent with the detection of human ILCPs in peripheral blood, we have previously observed rare donor-derived cells among BM ILCPs in parabiotic mice, suggesting that ILCPs, similar to HSCs, can physiologically traffic and systemically disperse at low levels in adult mice (Gasteiger et al., 2015; Wright et al., 2001). Of note, around 30-40% of II18r1<sup>+</sup>ST2<sup>-</sup> cells were labelled with intravenously injected anti-CD45 (ivCD45) (Figure 3A, B), and expressed genes associated with migration such as *Vim* and *Slpr1* (cluster 11, Figure 1E), and therefore could represent cells transiting through the pulmonary circulation. II18r1<sup>+</sup>ST2<sup>-</sup> cells were, however, extremely rare in the peripheral blood but readily detected in the ivCD45<sup>+</sup> fraction of the lung (Figure 3C), indicating that they may marginalize and enrich in lung vasculature. Interestingly, while II18r1<sup>+</sup> ILCs comprised a high fraction of ivCD45<sup>+</sup> and Ki67<sup>+</sup> cells, the majority of them was still ivCD45<sup>-</sup>, indicating their presence in the lung tissue at steady state (Figure 3D, E). Consistent with their residency in the lung and a minor exchange with circulating cells, all ivCD45<sup>-</sup>II18r1<sup>+</sup>ST2<sup>-</sup> subtypes were predominantly host-derived in parabiotic mice (Figure S4A). In addition, when comparing cells with an ILCP phenotype (II18r1<sup>+</sup>ST2<sup>-</sup>CD103<sup>-</sup>RORγt<sup>+</sup>) in lung versus BM, we found that about 20% fate-mapped positive for Il5 in lungs but not in the BM of Il5-Cre mice, further indicating that they are distinct from BM ILC progenitors "in transit" (Figure 3E). Instead, these data support the idea that ILCPs gained *Il5* expression locally and matured towards ILC2s within the lung tissue (Figure 3F).

To directly test their cell-intrinsic differentiation potential, we purified II18r1<sup>+</sup>Icos<sup>+</sup> lung ILCs by FACS for *in vitro* differentiation assays. In order to deplete cells that may be committed towards the ILC2 lineage, we additionally sorted ST2<sup>-</sup> and ST2<sup>-</sup>II17rb<sup>-</sup> double negative II18r1<sup>+</sup>Icos<sup>+</sup> ILCs from adult lungs, and cultured all three populations on OP9 in the presence of Il2, Il7, Il1β, and Il23 (Figure S4B), conditions which were recently used to expand human ILCPs (Lim et al., 2017). After ~21 days we performed scRNA-seq of lineage-negative output from these cultures along with the II18r1<sup>+</sup>Icos<sup>+</sup>ST2<sup>-</sup>II17rb<sup>-</sup> input population. First, we confirmed that the input population co-clustered with the naïve population from the uninfected lung ILC dataset and expressed only very low levels of *Il1r1l* and *Il17rb* (Figure S4C, D). Next, a combined analysis yielded 8 clusters, of which 5 corresponded to cells harvested from the culture (clusters 1, 2, 3, 4, 8) and 3 represent the input population (clusters 5, 6, 7, Figure 3G, H). Clusters 1 and 2 are marked by the expression of *Il5* and high levels of *Gata3*, *Il13* and *Calca* (Figure 3I), and were thus classified as ILC2s. The majority of cells differentiating from II18r1<sup>+</sup>Icos<sup>+</sup> and II18r1<sup>+</sup>Icos<sup>+</sup>ST2<sup>-</sup> input populations gave rise to these ILC2s (95 and 77%, respectively), while only ~15% of cells originating from II18r1<sup>+</sup>Icos<sup>+</sup>ST2<sup>-</sup>II17rb<sup>-</sup> cells, were ILC2s in clusters 1 and 2. Instead, these cells predominantly (61%) populated clusters 3 and 4, comprising a mixture of ILC2s and ILC3s with reduced levels of ILC2 specific gene expression. About 25% of cells originating from the II18r1<sup>+</sup>Icos<sup>+</sup>ST2<sup>-</sup>II17rb<sup>-</sup> population gave rise to NK/ILC1 cluster 8 characterized by expression of *Cxcr3*, *Klrl1* and *Gzmc*. Furthermore, whereas ILC2s largely reduced or lost *Il18r1* and *Tcf7* expression, cells in cluster 8 maintained high expression of these genes (Figure 3G-I). Of note, only a small fraction of cluster 8 originated from cultured II18r1<sup>+</sup>Icos<sup>+</sup>ST2<sup>-</sup> cells (Figure 3H), suggesting that depletion of II17rb expressing cells enriches the fraction of cells with ILC1/NK cell potential. We also assessed the potential of the II18r1<sup>+</sup>Icos<sup>+</sup> counterparts from the BM in the same conditions. Consistent with lung ILCs, the majority of II18r1<sup>+</sup>Icos<sup>+</sup>ST2<sup>-</sup> BM ILCs gave rise to ILC2s. In contrast, II18r1<sup>+</sup>Icos<sup>+</sup>ST2<sup>-</sup>II17rb<sup>-</sup> cells from the BM gave rise to only very few *Il13* expressing ILC2s (Figure S4E, F). Of interest, the presence of Il18 in these cultures led to the up-regulation of ILC2 effector genes (Figure S4H, I), as recently suggested (Ricardo-Gonzalez et al., 2018) but did not change the expansion of II18r1<sup>+</sup> ILC2s *in vitro* (Fig S4J).

These data suggest that *Il18r1*<sup>+</sup> lung and BM ILCs predominantly differentiated into *Gata3*<sup>hi</sup>*Il13*<sup>+</sup> ILC2. This bias towards ILC2s was evident under culture conditions that were recently shown to expand ILC progenitors (Lim et al., 2017) in the absence of alarmins Il25 and Il33, known to enforce ILC2 development. In order to test the expansion potential under such ILC2-activating conditions we cultured II18r1<sup>+</sup>ST2<sup>-</sup> and II18r1<sup>+</sup>ST2<sup>+</sup> ILC2s from lung in the presence of Il25 and Il33, and found that both populations generated similar numbers of *Gata3*<sup>hi</sup> ILCs as ST2<sup>+</sup> cells (Fig 3J). To test the differentiation potential *in vivo*, we activated the ILC2 compartment through infection with the parasitic helminth *Nippostrongylus brasiliensis* (Nb), which induces the local expansion of ILC2s. To this end, we sort-

purified I18r1<sup>+</sup>ST2<sup>-</sup> and I18r1<sup>+</sup>ST2<sup>+</sup> ILCs from lungs 2 days (d) post infection (p.i.) and co-transferred cells at a 1:1 ratio into sublethally irradiated RAGγc<sup>-/-</sup> hosts, as previously established (Ghaedi et al., 2020; Xu et al., 2019). Three weeks after transfer, we found that I18r1<sup>+</sup>ST2<sup>-</sup> cells partly gave rise to I18r1<sup>+</sup>ST2<sup>-</sup>, I18r1<sup>+</sup>ST2<sup>+</sup> and I18r1<sup>-</sup>ST2<sup>+</sup> ILC2s, consistent with the idea that these cells can differentiate into mature ILC2s (Figure 3K). Transferred I18r1<sup>-</sup>ST2<sup>+</sup> ILC2s, in contrast, only generated I18r1<sup>-</sup>ST2<sup>+</sup> progeny, suggesting that I18r1<sup>+</sup> cells represent early stages on the differentiation trajectory that are not readily generated from more mature cells in lymphopenic recipients.

Notably, I18r1<sup>+</sup> cells are a mixture of Ccr6<sup>+</sup>RORγt<sup>+</sup> ILC3s (Figure S4K) as well as Ccr6<sup>-</sup>RORγt<sup>-</sup> immature cells with NK/ILC1 and ILC2 potential, as suggested by the differentiation of these cells *in vitro* and *in vivo*, upon transfer (Figure S4 K-M). Therefore, only a fraction of the sorted I18r1<sup>+</sup> subsets will likely represent cells with ILC2 fate. To enrich for these cells, we sorted I18r1<sup>+</sup>ST2<sup>-</sup>IL5FM<sup>+</sup> cells for culture in the presence of IL25 and IL33, yielding many more Gata3<sup>hi</sup> ILC2s than culture of ST2<sup>+</sup> cells, highlighting the unique expansion potential of these immature lung ILCs (Figure 3L). Consistently, transfer of Ccr6<sup>-</sup> cells (i.e., RORγt-depleted and ILCP enriched cells) engrafted better and generated more ILC2s as compared to ST2<sup>+</sup> mature ILC2s *in vivo* (Figure S4M). To compare the clonal developmental potential of lung I18r1<sup>+</sup>ST2<sup>-</sup> subsets and BM ILCPs, we performed single-cell cultures in IL2, IL7, SCF in the presence of IL25 and IL33. Intriguingly, we found that both I18r1<sup>+</sup>ST2<sup>-</sup>CD103<sup>-</sup>Ccr6<sup>-</sup> lung and BM ILCPs gave rise to ILC2 or ILC1/NK cells or multiple lineages (Figure 3M). CD103<sup>+</sup> and ST2<sup>+</sup> cells, in contrast, preferentially generated ILC2s and ILC3s were undetectable under these type-2 polarizing conditions. Together, these data demonstrate that similar to the BM, the lung hosts a spectrum of immature cells with mixed lineage ILC potentials, including cells that are primed for ILC2 fate.

### Emergence of ILC2 subset heterogeneity during *N. brasiliensis* infection

Next, we investigated how the spectrum of phenotypic states of lung ILC2s and their differentiation trajectories change upon immune challenge. Infection with Nb causes local expansion of tissue-resident ILC2s during the acute phase of infection (Figure 4A), and around d15 postinfection (p.i.), in the tissue-repair phase, a fraction of lung ILC2s has emerged from cells recruited through the blood circulation (Gasteiger et al., 2015; Huang et al., 2018). During inflammation, ILCs exit from the small intestine (Huang et al., 2018), the lung (Ricardo-Gonzalez et al., 2020), and the BM (Stier et al., 2018) into the peripheral blood, and, hence, ILCs of different origins can traffic to the lung during infection to extend the local spectrum of phenotypes and functions. In order to dissect the heterogeneity of lung ILC2s upon immune challenge, “pan-ILCs” were sequenced at different time points p.i. along with “pan-ILCs” from the peripheral blood and ivCD45<sup>+</sup> ILCs from the lung vasculature (Figure 4B). After combining with the uninfected lung data (Figure 1) and removal of NK cell and macrophage contaminations, we inferred ILC2 sub-types using RaceID3 (Figure 4C). Our analysis highlights the dynamic emergence of ILC heterogeneity and specialization over time (Figure 4D, S5A). For example, cells from the uninfected condition contributed mainly to clusters 9, 11, 12 and 20 and *I18r1*<sup>+</sup> immature ILCs from the uninfected condition predominantly mapped to cluster 12 (Figure 4E, S5B, C). ILC2s derived from d4 p.i. expressed high levels of *Calca*, *Csf2* and *Cxcl2* and resided in cluster 4 and 10 together with a subset of ILC2s from the uninfected condition. However, ILC2s became more specialized during infection, exemplified by cluster 23 comprising almost exclusively d4 cells (Figure 4E, S5B), with marked up-regulation of *Csf2*, *Cxcl2*, *Cxcl1*, and *Il5*, which promote neutrophil and eosinophil recruitment, two myeloid subsets that expand in tissues early during helminth infection (Motran and Cervi, 2018). Similarly, ILC2s isolated at d15 p.i. comprised a variety of different effector subsets, e.g., cluster 14 with high expression of MHCII complex genes *Cd74*, *H2-Ab1* and *H2-Aa* (Oliphant et al., 2014; Symowski and Voehringer, 2019) and genes associated with T cell regulation such as *Il1r2*, *Ctla4* and *Furin*, or clusters 16 and 17, expressing high levels of *Il5*, *Il13*, *Cxcl2* and *Areg* (Figure 4E, S5B).

ILCs isolated on d7 and d10 p.i. contributed predominantly to clusters 2, 3, 5, and 15, which up-regulated *Vim* and, except for cluster 3, down-regulated expression of *Il1rl1*. Clusters 2 and 15 additionally up-regulated *Klrg1* (Figure 4E). We performed ivCD45 labeling to identify ILC2s that circulate through



the lung during infection (Huang et al., 2018; Ricardo-Gonzalez et al., 2020) and sequenced ivCD45<sup>+</sup> cells on d10 p.i. when their frequency peaked at ~10% of all lung ILC2s (Figure 4F). IvCD45<sup>+</sup> ILC2s clustered mainly with d7 and d10 ILCs (cl. 2 and 15) and few ivCD45<sup>+</sup> cells also clustered with mature effector ILC2s (e.g. cl. 6, 7 and 23). In contrast, blood ILC2s sorted on d15 p.i., when the wave of tissue-exiting ivCD45<sup>+</sup> cells has largely ceded, constitute a separate cluster (cl. 1) with high expression of *Vim*, *Klrg1*, *Id2*, *Klf2* and *S1pr1*, and have only minor contributions to clusters 2 and 3 (Figure 4D, E, S5B). Among them we detected a tiny population of cells expressing *Il18r1*, *Icos*, *Tcf7* and low levels of *Il1r1l*, consistent with the idea that Il18r1<sup>+</sup>ST2<sup>-</sup> cells may traffic to the lungs where they enrich in the lung vasculature and in the tissue (Figure 3A-C, and S5D, E).

We validated the dynamic regulation of selected markers, including PD1, MHCII, Il1r2, Ctl4, and CD25, on the protein level during the course of infection by flow cytometry, suggesting a close correlation of mRNA and protein levels for most genes (Figure 4G). Consistently, our analyses confirmed the regulated co-expression of Il1r2, PD1 and MHCII during late stages of infection (Figure S5F). Furthermore, we could validate the dynamic emergence of heterogeneous Il5-expressing ILC2s, co-expressing Il13 and/or PD1, resembling cluster 23 at d4 and clusters 16 and 17 at d10/d14 p.i. (Figures 4H).

### Differentiation trajectories of lung ILC2s during *N. brasiliensis* infection

We then investigated the differentiation dynamics of ILC2 subsets during Nb infection. Initially, we highlighted the position of cells from the early part of uninfected lung trajectory (uninfected lung clusters 1, 5, 11, 9, 8 in Figure 1) in the combined dataset (Figure 5A). Cells of the uninfected lung cluster 1 were largely localized to cluster 12, which was dominated by cells from the uninfected condition (Figure S6A). In contrast, the majority of cells from uninfected lung clusters 5, 11 and 9, representing early ILC2 differentiation stages, mixed with cells isolated during infection in clusters 2, 5 and 12 (Figure 5A), raising the question whether these cells may undergo dynamic differentiation and give rise to mature effector ILC2s during Nb infection. We applied StemID2 to infer potential trajectories starting from *Il18r1*<sup>+</sup> immature ILCs in either cluster 5 or 12 and ending in mature ILC2 effector clusters such as cluster 16 or 23 (Figure 5B). A large fraction of immature uninfected lung cells localized to clusters 2 and 8, suggesting that differentiating immature *Il18r1*<sup>+</sup> ILCs transition through these stages. From cluster 8 the trajectory branches out, potentially terminating in early (d4) or late (d15) effector clusters 23 or 16, respectively (Figure 5B). As the link between clusters 12 and 5 was populated well with intermediate single-cell transcriptomes (Figure S6B), we chose cluster 12 as a starting point, traversing cluster 5. Along both trajectories, we detected up-regulation of *Arg1*, *Klrg1*, *Gata3*, *Il1r1l*, and effector cytokines, as well as down-regulation of ILC progenitor genes (Figure 5C). Of note, we detected cells expressing *Il18r1* and *Tcf7* also within mature ILC2 compartments, albeit at reduced levels and frequencies compared to the progenitor cluster 12 (Figure 4E, S6C). The cluster composition along the trajectory suggested the presence of alternative starting points. Cluster 2 or 15, marked by a high contribution of potentially infiltrating ivCD45<sup>+</sup> cells (Figure 5B), may represent such alternative starting points for the *in situ* differentiation into mature nILC2 effector cells, consistent with the observation that both recruited and resident cells can contribute to lung ILC2s (Figure S6D) (Gasteiger et al., 2015; Huang et al., 2018).

These observations suggest that lung ILC2s can differentiate from an *Il18r1*<sup>+</sup> *Tcf7*<sup>+</sup> immature population during Nb infection. We found that Il18r1<sup>+</sup>ST2<sup>-</sup> cells were maintained during Nb infection while the proportion of ivCD45<sup>-</sup> cells among them gradually increased, consistent with the local expansion of these cells in the lung tissue (Figure 5D). At the same time, we observed a dramatic increase in highly proliferative Il18r1<sup>+</sup>ST2<sup>+</sup> cells over the course of infection (Figure 5E), which we characterized as differentiation intermediates along an expression gradient of *Tcf7* and *Gata3* and which were generated *in vivo* from transfers of Il18r1<sup>+</sup>ST2<sup>-</sup> immature cells (Figure 3K). Consistently, we observed the upregulation of ST2, Il5, Il13 and Ki67 within Il18r1<sup>+</sup> lung ILC2s over the course of infection (Figure 5F).

## Recruited cells contribute to the entire phenotypic spectrum of lung ILC2s during *N. brasiliensis* infection

A major open question is how cells recruited during inflammation contribute to the local pool of tissue ILCs, particularly, whether these cells retain specific phenotypes and functions, or adapt to their new tissue-environment. To address this question, we analyzed infiltrating versus resident ILC2s from lungs of Nb infected parabiotic mice. Here, two mice establish a common blood circulation and infiltrated donor-derived cells can be identified through allogeneic markers. Parabionts were infected with Nb, and similar numbers of donor and host ILC2s were sorted for scRNA-seq on d15 p.i., a time point when donor-derived, i.e., recruited, ILC2s represent about 10-15% of all lung ILC2s (Gasteiger et al., 2015; Huang et al., 2018). After RaceID3 analysis (Figure 6A) clusters were examined for their normalized host and donor contributions. Remarkably, all inferred clusters showed contribution of both host- and donor-derived cells (Figure 6B), with few differentially expressed genes between donor and host cells of the same cluster (Figure 6C). This observation demonstrates that cells recruited into the lung post parabiotic surgery contribute to all subpopulations that can be discriminated by unbiased scRNA-seq analysis at late stages of Nb infection.

Consistent with the idea of *in situ* differentiation of recruited cells from immature ILC2s, we found that some clusters were significantly enriched either for donor cells (clusters 3, 4, 5, 11 and 17) or host cells (cluster 1, 6, 7, 12, 13 and 15) (Figure 6B, S7A). While the former exhibited the signature of immature ILC2s with high levels of *Il18r1*, *Tcf7*, and *Cd7*, and low levels of *Gata3* and *Il1r1l* (clusters 4, 11 and 17), the latter showed expression of *Calca* and cells of clusters 12 and 13 additionally up-regulated *Cxcl2*, *Areg*, and *Il5* (Figure 6D, E and S7B). Thus, donor-derived cells are enriched in early differentiation stages. To further test this, we mapped the position of the donor- and host-enriched parabiosis clusters along the inferred trajectory of the time-course data (Figure 5b, Methods). The *Il18r1*<sup>+</sup> donor-dominated clusters 4, 11 and 17 of the parabiosis dataset had indeed the highest similarity to cells of time-course cluster 12, marking one of our inferred origins of ILC2 differentiation during Nb infection. Conversely, clusters enriched for host derived cells predominantly exhibited highest similarities with the terminal clusters of the inferred trajectories from the time-course data (Figure 6F and S7C).

These results demonstrate that donor-derived recruited cells populate all clusters while being enriched in immature *Il18r1*<sup>+</sup>ST2<sup>+</sup> ILCs, consistent with their *in situ* differentiation and their potential role in the renewal of tissue ILC2s. Importantly, our analyses reveal that ILCs recruited into the adult lung can generate the entire spectrum of tissue ILC2s based on their single cell transcriptomes.

## Bone marrow-derived ILCs give rise to the full phenotypic spectrum of ILC2s in the adult lung

We demonstrated that ILCs entering from the circulation give rise to all subtypes of tissue ILC2s during Nb infection (Figure 6B). However, the infiltrating cells can originate from different sources, including the BM (Constantinides et al., 2014; Klose et al., 2014; Stier et al., 2018), a circulating ILCP (Lim et al., 2017), or cells exiting from inflamed tissues (Huang et al., 2018; Ricardo-Gonzalez et al., 2020). These observations raise the possibility that cells from different origins give rise to limited specialized subsets of ILCs, and only together compose the full spectrum of tissue ILCs. Therefore, we investigated, whether cells originating from a defined source contribute to specialized subsets, or whether they can reconstitute the full phenotypic spectrum of lung ILC2s. To this end we adapted a shield chimera model (Gentek et al., 2018), where partial bone marrow chimerism was established through transfer of bone marrow after local irradiation of the hind legs (Figure 7A). Most of the body, including the lung, was shielded from the irradiation to avoid perturbation of the host tissue and immune cells. This procedure led to partial chimerism of BM-dependent cells of the myeloid and lymphoid lineages (e.g., NK cells), while very few ILC2s were detectable in the lung (Figure 7B). In contrast, ILC2s readily emerged from engrafted BM when shield chimeras were generated during the neonatal time window (data not shown), consistent with the local expansion of lung ILC2s during that time of ontogeny (Huang et al., 2018; Saluzzo et al., 2017; Schneider et al., 2019).

To test the differentiation potential of BM-derived cells in adult mice, adult shield chimeras were infected with Nb, and donor and host ILCs were sequenced on d15 p.i. (Figure 7A). Donor derived cells comprised ~5-8% of lung ILCs (Figure 7C, D). Similar to our observations for the parabiotic mice, all

inferred clusters comprised both host and donor derived cells (Figure 7E). Again, we detected donor enriched clusters with immature ILC2 signature up-regulating *Il18r1*, *Tcf7*, and *Zbtb16* (clusters 16 and 17) (Figure 7E, F and S7D, E). Within-cluster comparison revealed reduced levels of *Ccr2*, *Ctla2a*, *Cxcl2* and *Areg*, and higher levels of *Lmo4*, *Icos* and *Il17rb* in donor versus host derived cells (Figure 7G). These small but consistent differences suggest that donor derived cells carry a signature of less mature and migrating cells, consistent with their potential recruitment into the lung during infection. Our analyses do, however, not directly test this recruitment during infection, and donor-derived ILC2s could also differentiate from BM-derived cells that have migrated to tissues before infection. Consistent with both of these possibilities, our data establish that the adult bone marrow is a source of ILC2s, which can traffic to the lung and can generate the full spectrum of ILC2 subsets and differentiation stages in the lung during infection.

## Discussion

How ILCs are maintained and regenerated in the adult organism is currently unclear. Here, we provide evidence that differentiation and maturation of ILC2s from immature progenitors occurs within the adult lung. We analyzed the full spectrum of lung ILC2s at single-cell resolution and identified a heterogeneous population of  $Il18r1^+$  tissue-associated ILCs, that express *Tcf7* and *Zbtb16*, and low levels of *Gata3*, *ST2* and ILC2 effector molecules. We derived a continuous differentiation trajectory and validated the potential of  $Il18r1^+Tcf7^+$  cells to differentiate into  $Gata3^{hi}$  ILC2s.

Our extensive single-cell transcriptome comparison of ILC populations in the BM, and neonatal and adult lung at steady-state, as well as over the course of helminth infection in wild-type, parabiotic and shield chimeric mice support the idea that core programs of ILC2 differentiation are recapitulated during ontogeny as well as inflammatory challenge.

Intriguingly, recently described systemic ILC progenitors in men exhibit a transcriptional profile reminiscent of  $Il18r1^+$  ILCPs in murine lung (Ghaedi et al., 2020; Lim et al., 2017). Here, we confirmed that these cells are present in human lung tissue and can be further characterized by IL18R1 and TCF1 expression. Our findings suggest that  $Il18r1^+$  lung ILCPs represent the murine counterpart of human tissue ILCPs and thereby open new avenues to investigate how the local differentiation of tissue ILCs may contribute to inflammatory diseases (Mazzurana et al., 2018).

Il18 is best known as a myeloid derived cytokine that can activate lymphocytes, including ILC2s (Ricardo-Gonzalez et al., 2018). Il18 may, however, have additional functions and *Il18r1* should not be considered an exclusive marker of effector cells as it is also expressed by, e.g., BM hematopoietic stem cells (HSCs), common lymphoid progenitors, BM and circulating ILCPs, (Lim et al., 2017; Seillet et al., 2016; Silberstein et al., 2016; Xu et al., 2019), and local production of Il18 regulates the turnover of HSCs in their niche (Silberstein et al., 2016). Future studies should therefore investigate the niches of  $Il18r1^+$  ILC progenitors, as well as the potential functions of Il18 in regulating progenitor-like ILCs during homeostasis versus inflammatory challenge.

$Il18r1^+$  ILCs were virtually absent from peripheral blood of uninfected mice, but enriched in the lung vasculature which harbors many leukocytes that marginate from the blood stream, enrich at the endothelium and extravasate upon local inflammation (Anderson et al., 2014; Barletta et al., 2012; Yipp et al., 2017). It is therefore conceivable, that the vasculature may similarly serve as a local reservoir or niche for subsets of  $Il18r1^+$  ILCs. A large fraction of  $Il18r1^+$  ILCs was nevertheless  $ivCD45^-$ , confirming their presence in the lung parenchyma. In addition, analysis of parabiotic mice suggested that these cells are resident or exchange only very slowly in the lung-tissue. Importantly, lung-resident and BM ILCPs share core similarities. However, BM ILCPs were negative for *Il5* and *Il5-fatemap* label, while corresponding populations in the lung had intermediate levels of both, consistent with the idea that these cells receive local signals driving their maturation within the tissue. Intriguingly, the lung has been proposed as a reservoir of HSCs (Lefrançois et al., 2017), and may provide niches for the local differentiation of progenitors. Consistent with local differentiation, pulmonary  $Il18r1^+$  cells were highly

proliferative in uninfected tissue (in striking contrast to BM ILCPs or mature tissue ILCs) and gave rise to  $\text{Il18r1}^+\text{ST2}^-$ ,  $\text{Il18r1}^+\text{ST2}^+$  and  $\text{Il18r1}^-\text{ST2}^+$  ILCs in recipient lungs upon transfer. Transplanted mature  $\text{ST2}^+$  cells, however, did not generate  $\text{Il18r1}^+\text{ST2}^-$  or  $\text{Il18r1}^+\text{ST2}^+$  populations, suggesting that they are further downstream of a differentiation cascade, or that they cannot access the niche of immature ILCs in the lung.

Our data raise the possibility that tissue ILCPs may locally self-renew, but also indicate that they can receive input from circulating cells. First, we observed a small but significant donor contribution in the  $\text{ivCD45}^+\text{Il18r1}^+\text{CD103}^-$  subset in parabiotic mice. Second, we found that  $\text{Il18r1}^+\text{Tcf7}^+$  immature cells are generated from donor BM in shield chimeras. Third, donor-populations in infected parabiotic and shield chimeric mice were enriched with  $\text{Il18r1}^+\text{Tcf7}^+$  immature cells. Therefore, tissue ILCPs may receive low-grade influx from systemic counterparts. Increased competitive fitness of these cells may favour their contribution during aging, or when the local tissue niches are reorganized, e.g., upon regeneration of the ILC pool after infection. Such a model may explain why the frequency of donor-derived ILC2s is low early on, but gradually increases several months after parabiotic surgery, or in the reconstitution phase p.i. (Gasteiger et al., 2015; Huang et al., 2018; Ricardo-Gonzalez et al., 2020; Schneider et al., 2019). Despite a burst of  $\text{ivCD45}^+$  migrating inflammatory cells in the acute phase of helminth infection, these studies consistently showed that the majority of ILC2s were derived from tissue-resident cells around 2 weeks p.i.. Here we provide evidence for the *in situ* differentiation of ILCs from resident progenitors, highlighting a mechanism for the local, tissue-autonomous maintenance of ILCs in the adult organism. Importantly, we show that tissue ILCPs feature a phenotypic spectrum and mixed-lineage potential similar to the BM ILCP compartment. This heterogeneity may enable local “ILC-poiesis” to flexibly respond and generate the desired output upon distinct immune challenges.

Patients with inflammatory diseases such as psoriasis, hepatic fibrosis, or inflammatory bowel disease have elevated numbers of disease-associated ILCs in the blood. A major open question is how cells recruited during inflammation may contribute to the pool of tissue-resident cells. The extent to which local immune cells that have been generated early during ontogeny can be replenished in the adult organism has also been controversially discussed for macrophages, representing prototypic tissue-resident cells (Ginhoux and Williams, 2016; Mass et al., 2016). Here, we show that adult BM cells recruited into the adult lung can generate the full spectrum of lung-resident ILC2 subsets upon helminth infection. Our analysis reveals that ILCs differentiating in the adult organism can locally adapt the phenotypes of ILCs that have been suggested to emerge in distinct waves during ontogeny (Popescu et al., 2019; Schneider et al., 2019). Therefore, local tissue niches, rather than progenitor origin, may dominantly imprint ILC phenotypes. Together, our work highlights the local differentiation of immature ILCs as a mechanism of maintenance, phenotypic diversification and local adaptation of tissue ILCs.

### Limitations of Study

One current limitation in the study of ILCPs is the lack of conditional, stage-specific fate-mapping models to delineate the actual contribution of BM or tissue ILCPs during homeostasis or immune challenges in the adult organism. Moreover, we lack examples of conditions in which ILCPs are essential to maintain or regenerate a functional ILC pool. We speculate that ILCPs may have such critical roles upon depletion of ILCs, as documented in mice and men during different types of infection and environmental challenges (Kløverpris et al., 2016; Ng et al., 2018). It will be interesting to test whether local depletion and regeneration of ILC2s occurs during lung diseases as well. We are aware that our work highlights the tissue adaptation and differentiation potential of BM-derived ILCs in adult lungs but does not deconvolute potential contributions of other circulating cells. Future work should address if inflammatory ILC2s are a transient effector population or if (and how) they actually contribute to lung ILC2s once acute infection has resolved. Here, we mapped the heterogeneity of lung ILC2s and highlight multiple subsets that are dynamically emerging during infection, including, e.g., effector subsets expressing different combinations of cytokines and chemokines ( $\text{Il5/Il2/Cxcl2/Csf2}$  versus  $\text{Il5/Il13/Areg}$ ), or molecules involved in regulatory functions ( $\text{Pd1}$ ,  $\text{Ctla4}$ ,  $\text{Il1r2}$ ,  $\text{MHCII}$ ) and highly proliferative  $\text{Il18r1}^+$  and/or  $\text{CD103}^+$  states. While functional validation of these populations was beyond the scope of this study, we feel that our interactive cell atlas (<http://murine-ilc-atlas.ie-freiburg.mpg.de/>) provides a rich resource for future studies testing subset-specific functions of ILC2s.

## Figure Legends

**Figure 1. Heterogeneity of ILC2s in healthy adult lung tissue and immature  $\text{Il18r1}^+$  ILCs.** (A) Workflow for lung ILC scRNA-seq library preparation. (B) Representative sorting strategy for  $\text{Lin}^- \text{NK1.1}^- \text{Il7ra}^+ \text{Thy1.2}^{\text{hi/im}}$  ‘pan’-ILCs from lung. (C) t-SNE map of single-cell lung ILC transcriptomes clustered with RaceID3. (D) t-SNE maps indicating log2 normalized expression of candidate genes. (E) Candidate gene expression for clusters with at least 20 cells. Color represents z-score mean expression across clusters and dot size represents fraction of cells in the cluster expressing the gene. (F) Lineage inference using StemID2. Node color depicts transcriptome entropy and link color indicates p-value of StemID2 links ( $P < 0.05$ , Methods). (G) Pseudo-temporal gene expression profiles (local regression) of representative ILC2 genes (top) and ILC progenitor genes (bottom) along the predicted trajectory. Color bars indicate cluster identity. (H,I), Representative FACS analyses of  $\text{Lin}^- \text{Il7ra}^+$  ILCs in adult lung at steady state. Gating (H) and percentage (I) of  $\text{Il18r1}^+ \text{ST2}^- \text{ROR}\gamma\text{t}^- \text{Klrg1}^-$  ILCs. (J-K) Fraction of  $\text{Ki67}^+$  (J) and  $\text{Il5}^+$  (K) cells within the indicated lung ILC subsets (ICS after incubation with monensin,  $\text{ROR}\gamma\text{t}^+$  ILCs are not excluded). (L) FACS analysis of  $\text{Lin}^- \text{Il7ra}^+$  lung ILCs subdivided into  $\text{Il18r1}^+ \text{ST2}^- \text{ROR}\gamma\text{t}^- \text{Klrg1}^-$ ,  $\text{Il18r1}^+ \text{ST2}^+$  and  $\text{Il18r1}^- \text{ST2}^+$  ILCs. Depicted is the mean fluorescence intensities (MFI) for  $\text{Tcf7}$ ,  $\text{Il18r1}$ ,  $\text{Il17rb}$ ,  $\text{Gata3}$ ,  $\text{ST2}$  and  $\text{CD25}$ . Data in (H-L) are representative of 2-3 independent experiments ( $n=4-12$  mice). Graphs in (I), (J) and (K) depict data as mean  $\pm$  SD, (one-way ANOVA Tukey’s multiple comparisons test,  $*P < 0.05$ ,  $**P < 0.01$ ;  $***P < 0.001$ ;  $****P < 0.0001$ ; ns, not significant).

**Figure 2.  $\text{Il18r1}^+ \text{ST2}^-$  ILCs in neonatal and adult lung share similarities with  $\text{Il18r1}^+$  BM ILCs.** (A-C), t-SNE maps of neonatal lung ILC single-cell transcriptomes highlighting RaceID3 clusters (A) or sorted  $\text{ST2}^-$  and  $\text{ST2}^+$  ILCs (B), and log2 normalized expression of representative genes marking putative ILC progenitors (C). (D) Expression of candidate genes within neonatal lung clusters with at least 10 cells. Color represents z-score mean expression across clusters and dot size represents fraction of cells in the cluster expressing the gene. (E) FACS analysis of neonatal (P4) and adult (6-8w)  $\text{Il18r1}^+ \text{ST2}^-$  lung ILCs, represented as t-SNE maps, indicating expression levels for  $\text{Gata3}$ ,  $\text{ROR}\gamma\text{t}$ ,  $\text{PLZF}$ ,  $\text{Tcf7}$ ,  $\text{C-kit}$ ,  $\text{CD103}$  and  $\text{Ki67}$ . (F) Histograms comparing ILC subsets ( $\text{PLZF}^+ \text{Gata3}^+$ ,  $\text{ROR}\gamma\text{t}^- \text{CD103}^-$ ,  $\text{ROR}\gamma\text{t}^+ \text{CD103}^+$ ,  $\text{Il18r1}^- \text{ST2}^+$  ILC2s and  $\text{ROR}\gamma\text{t}^+$  ILC3) in adult lung for  $\text{Gata3}$ ,  $\text{Icos}$ ,  $\alpha 4\beta 7$ ,  $\text{PD1}$ ,  $\text{CD25}$  and  $\text{Il17rb}$  expression levels. Data are representative of 2-3 independent experiments ( $n=6-12$  mice). (G) t-SNE map of  $\text{Il18r1}^+ \text{Icos}^+$  BM ILCs highlighting RaceID3 clusters. (H) Expression of candidate genes for  $\text{Il18r1}^+ \text{Icos}^+$  BM clusters with at least 10 cells. Color represents z-score mean expression across clusters and dot size represents fraction of cells in the cluster expressing the gene. (I-J) FACS analysis of human lung and PB ILCs pre-gated on  $\text{CD45}^+ \text{Lin}^- \text{CD127}^+ \text{EOMES}^- \text{T-bet}^- \text{CRTH2}^- \text{NKp44}^- \text{c-KIT}^+$  (I) or  $\text{c-KIT}^{\pm}$  ILCs (J). Figures depict representative staining (I) and t-SNE maps (J) indicating the expression levels of selected markers. Data are representative of 3 independent experiments ( $n=2-4$  samples each).

**Figure 3.  $\text{Il18r1}^+$  ILCs are immature cells that give rise to  $\text{Gata3}^{\text{hi}}$  ILC2s in the lung.** (A-B) FACS analysis of  $\text{Ki67}$  expression and intravenous  $\text{CD45}$  label (ivCD45) in  $\text{Il18r1}^+ \text{ST2}^- \text{ROR}\gamma\text{t}^- \text{Klrg1}^-$  and  $\text{Il18r1}^- \text{ST2}^+$  adult lung ILC2s. Representative gating (A) and percentage of ivCD45<sup>+</sup> cells within indicated ILC subsets (B). (C) Number of  $\text{Lin}^- \text{Il7ra}^+ \text{Il18r1}^+$  cells per ml blood in PBMCs ( $\text{Il18r1}^+ \text{ST2}^-$  ILCs) and lung (ivCD45<sup>+</sup> $\text{Il18r1}^+ \text{ST2}^- \text{ROR}\gamma\text{t}^- \text{Klrg1}^-$  ILCs). Graphs in (B) and (C) depict data as mean  $\pm$  SD (unpaired t-test,  $*P < 0.05$ ,  $**P < 0.01$ ;  $***P < 0.001$ ;  $****P < 0.0001$ ; ns, not significant). (D-E) Representative gating (D) and percentage of  $\text{Ki67}^+$  and ivCD45<sup>+</sup> cells (E) for indicated  $\text{Il18r1}^+ \text{ST2}^-$  ILC subsets in adult lung. Data are pooled from 3-4 independent experiments ( $n=4-16$  mice). (F) Percentage of  $\text{Il5}^{\text{Cre}}$  fate-mapped ( $\text{Il5FM}^+$ )  $\text{Il18r1}^+ \text{ST2}^- \text{ROR}\gamma\text{t}^- \text{CD103}^-$  ILCs in lung and BM.  $\text{ROR}\gamma\text{t}$  staining was used to exclude ILC3. Data are pooled from 2-3 independent experiments ( $n=4-9$  mice). Graphs in (E) and (F) depict data as mean  $\pm$  SD (one-way ANOVA Tukey’s multiple comparisons test (E) or unpaired t-test (b, c, f),  $***P < 0.001$ ;  $****P < 0.0001$ ). Data are pooled from 3-4 independent experiments with a total of  $n = 8-15$  mice). (G) t-SNE representation of single-cell transcriptomes of lung  $\text{Il18r1}^+ \text{Icos}^+$  ILCs before (“input”) and after culture (“output”) highlighting RaceID3 clusters. (H) Cluster composition of samples (left) and sample composition of clusters (right, for legend see (G)). (I)

Expression of candidate genes for input and output populations. Color represents log<sub>2</sub> mean expression in the respective cluster and dot size indicates fraction of cells expressing the gene in the cluster. (J) Total number of Gata3<sup>+</sup> ILC2s after culture of indicated lung ILCs for 15 days in the presence of Il2, Il7, SCF, Il25 and Il33 on OP9-DL1 for 15 days. (K) FACS analysis of Lin<sup>-</sup>Il7ra<sup>+</sup> lung ILCs on d21 post transfer into sublethally irradiated CD45.2<sup>+</sup>RAG<sup>-/-</sup>γc<sup>-/-</sup> mice. Congenically marked Il18r1<sup>+</sup>ST2<sup>-</sup> and Il18r1<sup>-</sup>ST2<sup>+</sup> ILCs sorted from lungs of d2 Nb infected mice were cotransferred (2,000 cells each). Data are representative of 2 individual experiments (n=3 mice). (L) Total number of Gata3<sup>+</sup> ILC2 after culture of indicated lung ILCs as in (J). Data in (J) and (L), are pooled from 2 independent experiments with n=3-4 repeat wells each. (M) Indicated lung and BM ILC subsets were sorted as single cells onto OP9-DL1 monolayers and cultured for 19 days. Clonal progeny was analyzed by FACS. Pie charts indicate distribution of culture output across positive single cell cultures.

**Figure 4. Emerging heterogeneity of lung ILC2s during *Nippostrongylus brasiliensis* infection.** (A) FACS analysis of the fraction of Lin<sup>-</sup>Gata3<sup>+</sup> ILC2s among live CD45<sup>+</sup> cells (left) and of Ki67<sup>+</sup> cells among Lin<sup>-</sup>Gata3<sup>+</sup> ILC2s (right) in Nb infected lungs. Data are pooled from 3 independent experiments with a total of n=12-16 mice per time point. (B) Sequenced samples at the respective time-points during Nb infection. (C) t-SNE map of combined Nb infection time-course and uninfected lung data (cf. Figure 1) highlighting RaceID3 clusters. (D) Overlay of t-SNE map from (C) with pie charts indicating normalized sample contribution to clusters. (E) Expression of candidate genes for ILC subsets for inferred time-course clusters with at least 20 cells (left) or the respective sample (right). Color represents z-score of mean expression across clusters/samples and dot size represents fraction of cells positive for the gene in the cluster/sample. (F) Time-course FACS analysis of the fraction of ivCD45<sup>+</sup> ILC2s in Nb infected lungs. FACS data are pooled from 2-3 independent experiments with n=6-12 mice per time point. (G) Time-course FACS analysis of Il7ra<sup>+</sup> ILCs in Nb infected lungs (black) compared with fractions of RNA expressing cells of time-course dataset (red). FACS data are pooled from 2-3 independent experiments with n=6-9 mice per time point. (H) Time-course FACS analysis of Nb infected lungs. Expression of indicated markers represented as t-SNE maps for Il5<sup>+</sup> ILCs. ICS after incubation with PMA, ionomycin and monensin. Data are pooled from 5 mice for each time point and merged from equal numbers of Lin<sup>-</sup>Il7ra<sup>+</sup> ILCs. Graphs in (A) and (F) depict data as mean ± SD (one-way ANOVA Tukey's multiple comparisons test, \*\*\**P* < 0.001; \*\*\*\**P* < 0.0001).

**Figure 5. Differentiation trajectories of lung ILC2s during *Nippostrongylus brasiliensis* infection.** (A) t-SNE maps showing the distribution of uninfected lung immature clusters (cf. Figure 1) within the combined data set (top) and pie charts of the distributions (bottom) (cf. Figure 4C). (B) StemID2 lineage inference and normalized sample contribution to clusters depicted as pie-charts. Link color indicates p-value of StemID2 links (*P* < 0.05, Methods). (C) Pseudo-temporal gene expression profiles (local regression) along depicted trajectories. Color bars indicate cluster and sample identity, respectively, of cells ordered along the depicted trajectory by StemID2. (D) Time course analysis for the percentage of Il18r1<sup>+</sup>ST2<sup>-</sup>RORγt<sup>+</sup>Klrg1<sup>-</sup> ILCs (left) and the comparison of ivCD45 labelled and unlabelled fractions (right). (E) Time course analysis for the percentage of Il18r1<sup>+</sup>ST2<sup>+</sup> (left) and Il18r1<sup>-</sup>ST2<sup>+</sup> (right) ILC2s. Data in (D) and (E) are representative of 2 independent experiments with n=4-6 mice per timepoint. Graphs depict data as mean ± SD (one-way ANOVA Tukey's multiple comparisons test, \**P* < 0.05, \*\**P* < 0.01; \*\*\**P* < 0.001; \*\*\*\**P* < 0.0001). (F) Time-course FACS analysis of Nb infected lungs. Expression of indicated markers represented as t-SNE maps for Il18r1<sup>+</sup> ILCs. ICS after incubation with PMA, ionomycin and monensin. Data are pooled from 5 mice for each time point and merged from equal numbers of Lin<sup>-</sup>Il7ra<sup>+</sup> ILCs.

**Figure 6. Recruited cells contribute to the entire phenotypic spectrum of lung ILC2s during *Nippostrongylus brasiliensis* infection.** (A) t-SNE map of cells derived from parabiotic mice at d15 p.i. highlighting RaceID3 clusters. (B) StemID2 lineage inference and normalized donor and host contribution to clusters depicted as pie-charts. Link color indicates p-value of StemID2 links (*P* < 0.05, Methods). (C) Differentially expressed genes (*P* < 0.05, Methods) between donor and host cells within the depicted clusters. (D) t-SNE maps indicating log<sub>2</sub> normalized expression of candidate genes in the

parabiosis data. (E) Expression of representative candidate genes for ILC subsets in parabiosis clusters with at least 20 cells. Color represents z-score of mean expression across clusters and dot size represents fraction of cells expressing the gene in the cluster. (F) t-SNE map of parabiosis data showing normalized weights of indicated parabiosis cluster medoids for cells within the Nb time-course data (cf. Figure 4) as computed by cluster mapping (Methods).

**Figure 7. Bone marrow-derived ILCs give rise to the full phenotypic spectrum of ILC2s in the adult lung.** (A) Nb infection of the shield chimera model. (B) Frequency of donor-derived cells from transferred bone marrow (BM). (C-D) t-SNE map of ILCs from shield chimeric mice isolated at d15 p.i. highlighting RaceID3 clusters (C) and donor or host origin for each cell (D). (E) StemID2 lineage inference and normalized donor and host contribution to clusters depicted as pie-charts. Link color indicates p-value of StemID2 links ( $P < 0.05$ , Methods). (F) Expression of candidate genes for ILC subsets in the shield chimera clusters with at least 10 cells. Color represents z-score of mean expression in a cluster and dot size represents fraction of cells expressing the gene in the cluster. (G) Differentially expressed genes ( $P < 0.05$ , Methods) between donor and host cells within the depicted clusters.

### Supplementary Figure Legends

**Figure S1. Characterization of lung ILCs. Related to Figure 1.** (A) Full gating strategy for lung ILCs, including ROR $\gamma$ t expression and intracellular CD3 $\epsilon$  staining. (B) Scatter plot showing normalized expression of *Klf2* (y-axis), *Cd69* (x-axis) and log<sub>2</sub> normalized expression of *Cxcl2* (dot color) across cells of the indicated clusters of ILCs from uninfected lung. (C) Scatter plot showing normalized expression of *Rorc* (y-axis), *Il17rb* (x-axis) and log<sub>2</sub> normalized expression of *Tbx21* (left panel) or *Zbtb16* (right panel) across cells of the indicated *Il18r1*<sup>+</sup> uninfected lung clusters. (D) Self-organizing map (SOM) were used to group z-score transformed pseudo-temporal expression profiles into modules of co-expressed genes based on similarity. Cells were ordered by StemID2 along the predicted pseudo-temporal differentiation trajectory (5>11>9>8>4>10). Each row in the heatmap corresponds to a pseudo-temporal expression profile, and modules of co-expressed genes are separated by dashed grey lines. (E) KEGG enrichment (upper panel) or Gene ontology (GO) enrichment (lower panel), for genes expressed in module 5 or 1, respectively, of the pseudo-temporal SOM analysis from (D). Genes within the respective modules were used as input and the enriched GO terms or KEGG pathways are depicted in decreasing order of gene counts within GO term or KEGG pathway, respectively. Dot size represents the gene count within the pathway; the fraction of genes of the respective module matching KEGG pathway or GO term genes of total genes of the respective module is depicted on the x-axis and the color indicates the p-value. (F) Pseudo-temporal gene expression profiles of ILC progenitor (right panel) or ILC2 genes (left panel) along the predicted trajectory of uninfected lung ILC2s terminating in *Areg*<sup>+</sup> cells (cluster trajectory, 5>11>9>8>4>12). (G) *Il18r1*<sup>+</sup>ST2<sup>+</sup>ROR $\gamma$ t<sup>+</sup>Klrg1<sup>-</sup> ILCs in RAG KO mice. (H) Cluster-specific gene expression for RaceID3 clusters after removal of the proliferation signature (see Methods). Clusters with at least 20 cells are depicted. Color represents the z-score of the mean expression across clusters and dot size represents the fraction of cells in the cluster expressing the gene. (I) Lineage inference for RaceID3 clusters from (H) using StemID2. Link color indicates p-value of StemID2 links ( $P < 0.05$ , Methods). Node color indicates transcriptome entropy. (J) Pseudo-temporal gene expression profiles of ILC progenitor or ILC2 genes along the predicted trajectory of (I) (after removal of proliferation signature from clustering) terminating in *Cxcl2*<sup>+</sup> cells (cluster trajectory, 1>7>9>2>5>8). (K) Cell-cycle score (Methods) of RaceID3 clusters before (upper panel) and after (lower panel) removal of cell-cycle genes from clustering. Color bars below the SOM (D) or expression profiles (F) and (J) indicate cluster identity of cells ordered along the depicted trajectory based on StemID2 derived projection coordinates.

**Figure S2. *Il18r1*<sup>+</sup> ILCs comprise progenitor-like and effector-like ILCs in skin and lung. Related to Figure 1.** (A) Cluster information of healthy lung transcriptomes from this study integrated with lung and skin transcriptomes from Ricardo-Gonzalez et al. Integration and clustering was performed using

Seurat (see Methods). UMAP depicting cluster identity (left panel) or sample identity (right panel). Bar chart indicates sample composition of each cluster (middle panel). (B) Expression of candidate genes in skin cells in the depicted clusters. Color represents the z-score of the mean expression across skin cells in the respective clusters and dot size indicates the fraction of skin cells expressing the gene among skin cells in the cluster. (C) UMAP indicating log<sub>2</sub> normalized expression of selected candidate genes. (D) Representative FACS staining for ILC subsets within II18r1<sup>+</sup>ST2<sup>-</sup> ILCs based on the expression of RORγt and CD103. (E) FACS analysis showing cytokine expression (Il5, Il13 and Il17A) within lung II18r1<sup>+</sup>ST2<sup>-</sup>RORγt<sup>+</sup>CD103<sup>+</sup> effector-like ILCs, II18r1<sup>+</sup>RORγt<sup>+</sup> ILC3, II18r1<sup>+</sup>RORγt<sup>-</sup>CD103<sup>-</sup> ILCs and II18r1<sup>-</sup>ST2<sup>+</sup> ILC2. Intracellular staining of Il5, Il13 and Il17A protein after PMA and Ionomycin re-stimulation in the presence of monensin and brefeldin A. Cells were incubated at 37°C for 3.5h. Data are representative of 2 independent experiments (n=3 - 4 mice). Shown are FACS stainings of cytokines versus ST2 (upper panel) and percentage of cytokine producing cells across II18r1<sup>+</sup> populations (lower panel) depicted as mean ± SD, (one-way ANOVA Tukey's multiple comparisons test, \**P* < 0.05, \*\**P* < 0.01; \*\*\**P* < 0.001; \*\*\*\**P* < 0.0001; ns, not significant).

**Figure S3. II18r1<sup>+</sup>ST2<sup>-</sup> ILCs identified in neonatal and adult lung share similarities with II18r1<sup>+</sup> ILC progenitors in BM. Related to Figure 2.** (A) Sorting strategy for the enrichment of neonatal lung ILC2s. (B-C) Representative FACS analysis of Il7ra<sup>+</sup> ILCs (B) and percentage of Gata3<sup>+</sup>PLZF<sup>+</sup> ILCs within II18r1<sup>+</sup>ST2<sup>-</sup> ILCs (C) in P4, P7 and P10/11 neonatal lungs. Data are representative of 2 independent experiments (n = 4 mice). Graphs depict data as mean ± SD, (one-way ANOVA Tukey's multiple comparisons test, \**P* < 0.05, \*\**P* < 0.01; \*\*\**P* < 0.001; \*\*\*\**P* < 0.0001; ns, not significant). (D) Representative FACS analysis of PLZF<sup>+</sup> ILCs in adult lung. (E) Sorting strategy for II18r1<sup>+</sup>Icos<sup>+</sup> BM ILCs. (F) UMAP depicting cluster information (left panel) and sample information (right panel, cluster partition from Figure 2G was used as sample information) of BM transcriptomes from this study integrated with ILCP and ILC2P transcriptomes from Walker et al. Integration was performed using Seurat and clustering was done using RaceID3 (see Methods). (G) Sample composition of the indicated clusters from (F). (H) Expression of candidate genes in the respective clusters of the integrated dataset (F). Color represents the z-score of the mean expression across clusters and dot size represents the fraction of cells in the cluster expressing the gene. (I), Full gating strategy of human Lung and PB ILCs.

**Figure S4. II18r1<sup>+</sup> ILCs are immature cells that give rise to Gata3<sup>hi</sup> ILC2s in the lung. Related to Figure 3.** (A) Fraction of host-derived cells among Lin<sup>+</sup>CD3<sup>+</sup> T cells, Lin<sup>+</sup>CD3<sup>-</sup>Thy1.2<sup>int</sup> NK cells and Lin<sup>+</sup>Il7ra<sup>+</sup> ILCs (left panel) and among ivCD45<sup>-</sup> (middle panel) and ivCD45<sup>+</sup> (right panel) of the indicated ILC subsets in parabiotic mice. Data are pooled from 4 pairs of parabiotic mice n=8. Graphs depict data as mean ± SD. (B) Strategy for sorting of Lin<sup>+</sup>Il7ra<sup>+</sup>Thy1.2<sup>+</sup>II18r1<sup>+</sup>Icos<sup>+</sup> populations (upper panel) and schematic representation for co-culture of II18r1<sup>+</sup>Icos<sup>+</sup> ILCs and stromal cells (lower panel, related to Figure 3G-I). (C) t-SNE map showing a combined analysis of uninfected lung ILCs (Figure 1C, F) and sorted II18r1<sup>+</sup>Icos<sup>+</sup>ST2<sup>-</sup>Il17rb<sup>-</sup> ILCs used as input for the *in vitro* cultures (related to Figure 3G-I). (D) Histograms indicating the normalized expression of *Il1r1* (upper panels) or *Il17rb* (lower panel) within cells of *Il18r1*<sup>+</sup> immature uninfected lung clusters (1,5,11), all other clusters except *Il18r1*<sup>+</sup> clusters (referred to mature ILC2s), or II18r1<sup>+</sup>Icos<sup>+</sup>ST2<sup>-</sup>Il17rb<sup>-</sup> input cells. (E-F) Expression of candidate ILC genes within input and cultured output samples for sorted II18r1<sup>+</sup>Icos<sup>+</sup> lung populations (E) and cultured output of II18r1<sup>+</sup>Icos<sup>+</sup> BM populations (F). Color represents the log<sub>2</sub> mean expression and dot size represents the fraction of cells in a cluster expressing the gene. (G) t-SNE maps indicating log<sub>2</sub> normalized expression of *Il1r1* and *Il23r* expression in input and cultured output cells. (H) UMAP representation depicting status of Il18 stimulation (left panel) and sample information (right panel) of cultured II18r1<sup>+</sup>Icos<sup>+</sup> lung ILC transcriptomes. (I) Expression of candidate ILC and ILC2 genes within lung samples cultured in the indicated cytokine conditions. Color represents the z-score of mean expression across samples and dot size represents the fraction of cells positive for the gene in the sample. (J) Numbers of Gata3<sup>+</sup> ILC2s after culture of II18r1<sup>+</sup>ST2<sup>-</sup> ILCs for 15 days in the presence of OP9-DL1 and the indicated cytokines. (K) FACS analysis for Ccr6 expression on RORγt<sup>+</sup> ILCs. (L) Numbers of



ROR $\gamma$ t<sup>+</sup> ILC3 and Nkp46<sup>+</sup> NK/ILC1 (left panel) or Gata3<sup>+</sup> ILC2s (right panel) after culture of Il18r1<sup>+</sup>Ccr6<sup>+</sup>, Il18r1<sup>+</sup>Ccr6<sup>-</sup> and Il18r1<sup>-</sup>ST2<sup>+</sup> ILC2 for 15 days in the presence of OP9-DL1 and the indicated cytokines. Data are representative of 2 independent experiments with n= 4 repeat wells each. (M) ILC2, ILC3, ILC1 and NK cells in lungs of RAG $\gamma$ c KO mice 6 weeks post co-transfer of either Il18r1<sup>+</sup>Ccr6<sup>+</sup> and Il18r1<sup>-</sup>ST2<sup>+</sup> ILCs, or Il18r1<sup>+</sup>Ccr6<sup>-</sup> and Il18r1<sup>-</sup>ST2<sup>+</sup> ILCs. 1,500 congenically marked cells were transferred for each indicated ILC subset. Data represent 2 individual mice each.

**Figure S5. Emerging heterogeneity of lung ILC2s during *Nippostrongylus brasiliensis* infection. Related to Figure 4.** (A) t-SNE map of combined Nb time-course and uninfected lung data highlighting the sample-of-origin for each cell. (B) Expression of candidate genes for ILC and ILC2 subsets within combined Nb infection time-course and uninfected lung clusters. Color represents log<sub>2</sub> mean expression of the gene in the respective cluster and dot size indicates the fraction of cells expressing the gene in the cluster. (C) t-SNE map indicating log<sub>2</sub> normalized expression of selected ILC progenitor genes. (D) t-SNE map indicating RaceID3 clusters of ILC2s sorted from peripheral blood at day 15 p.i.. (E) Expression of candidate genes for ILC and ILC2 subsets within peripheral blood derived ILC2 clusters. Color represents the z-score of the mean expression across clusters and dot size represents the fraction of cells in the cluster expressing the gene. (F) FACS analysis of d4 and d10 Nb infected lungs. Expression of indicated markers represented as t-SNE maps for ST2<sup>+</sup> ILC2. Data are pooled from 5 mice for each time point and merged from equal numbers of Lin<sup>-</sup>Il7ra<sup>+</sup> ILCs.

**Figure S6. Differentiation trajectories of lung ILC2s during Nb infection. Related to Figure 5.** (A) Pie charts showing cluster distribution of each sample. Clusters with <1% contribution are not labelled. (B) StemID2 projection coordinates of each cell from cluster 12 onto inter-cluster links between the medoid of cluster 12 and the medoids of all other clusters. Cells of cluster 12 are ordered by hierarchical clustering along the y-axis and links are ordered by hierarchical clustering along the x-axis. (C) Scatter plot showing normalized expression of *Tcf7* (y-axis), *Il18r1* (x-axis) and log<sub>2</sub> normalized expression of *Il1r1* (dot color) across cells of the depicted time-course clusters. (D) Pseudo-temporal gene expression profiles of representative migratory genes (lower panel) and ILC2 genes (upper panel) along the predicted ivCD45 trajectory (15>2>8>6>16). Colored lines indicate a local regression of the expression profiles. Color bars below the expression profiles indicate cluster and sample information of cells ordered along the depicted trajectory based on StemID2 derived projection-coordinates.

**Figure S7| Recruited cells contribute to the entire phenotypic spectrum of lung ILC2 during *Nippostrongylus brasiliensis* infection. Related to Figure 6 (S7A-C) and Figure 7 (S7D, E).** (A) Enrichment scores and respective p-values of donor (upper panel) and host (lower panel) enriched clusters. P-values were calculated by an one-sided Fisher-test and shown as negative log<sub>10</sub> p-values. \**P* < 0.05. (B) Expression of representative candidate genes for ILC and ILC2 subsets in the parabiosis clusters. Color represents the log<sub>2</sub> mean expression of a gene in a cluster and dot size represents the fraction of cells expressing the gene in the cluster. (C) t-SNE map of Nb time-course data indicating normalized weights of each cell from the Nb time-course for the parabiosis cluster medoids inferred by cluster mapping (Methods). (D) Enrichment scores and respective p-values of donor and host enriched shield chimera clusters. P-values are calculated by an one-sided Fisher-test and shown as negative log<sub>10</sub> p-values. \**P*<0.05. (E) Expression of representative candidate genes for ILC and ILC2 subsets in the shield chimera clusters. Color represents the log<sub>2</sub> mean expression of a gene in a cluster and dot size represents the fraction of cells expressing the gene in the cluster.

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### **Author Contributions**

Conceptualization, funding acquisition and project coordination, D.G. and G.G.; scRNA-seq experiments, computational analysis and web interface, P.Z.; FACS analysis and transfers, M.L.; in vitro culture experiments, M.L., P.Z., J.S.H.; parabiosis, X.F., S.E.; shield chimera, R.G.; human ILCs, D.C.H.; experimental support, K.K., N.P., C.F., R.D.L., A.M.K.; Il5 reporter mouse, R.M.L.; Nb culture, C.S.; experimental design, M.L., P.Z., D.G., G.G.; supervision, A.R., D.V., M.B., C.R., D.G., G.G.; paper writing and visualizations, P.Z., M.L., D.G., G.G.; review and editing, all authors.

### **Declaration of Interests**

The authors declare no conflict of interests.

## STAR METHODS

REAGENT or RESOURCE Antibodies	SOURCE	IDENTIFIER
<b>Antibodies</b>		
InVivoMAb anti-mouse CD16/CD32 (Clone 2.4G2)	Bioxcell	Cat # BE0307
Fixable Viability Dye eFluor® 780	eBioscience	Cat # 65-0865-18
Anti-mouse CD3 $\epsilon$ Biotin (Clone 145-2C11)	BioLegend	Cat# 100304
Anti-mouse CD5 Biotin (Clone 53-7.3)	BioLegend	Cat# 100604
Anti-mouse CD19 Biotin (Clone 6D5)	BioLegend	Cat# 115504
Anti-mouse Gr1 Biotin (Clone RB6-8C5)	BioLegend	Cat# 108404
Anti-mouse F4/80 Biotin (Clone BM8)	BioLegend	Cat# 123106
Anti-mouse CD11b Biotin (Clone M1/70)	BioLegend	Cat# 101204
Anti-mouse CD11c Biotin (Clone N418)	BioLegend	Cat# 117304
Anti-mouse NK1.1 Biotin (Clone PK136)	BioLegend	Cat# 108704
Anti-mouse TCR $\beta$ Biotin (Clone H57-597)	BioLegend	Cat# 109204
Anti-mouse TCR $\gamma\delta$ Biotin (Clone eBioGL3)	eBioscience	Cat # 13-5711-85
Anti-mouse Fc $\epsilon$ R1 Biotin (Clone MAR-1)	BioLegend	Cat# 134304
Anti-mouse TER-119 Biotin (Clone TER-119)	BioLegend	Cat# 116204
Anti-mouse CD45R/B220 Biotin (Clone RA3-6B2)	BioLegend	Cat# 103204
Anti-mouse Ly6G Biotin (Clone 1A8)	BioLegend	Cat# 127604
Streptavidin V500	BD Biosciences	Cat# 561419
Anti-mouse Thy1.2 FITC (Clone 30-H12)	BioLegend	Cat# 105306
Anti-mouse Thy1.2 AF786 (Clone 30-H12)	BioLegend	Cat# 105331
Anti-mouse CD127/IL7R $\alpha$ APC (Clone A7R34)	BioLegend	Cat# 135012
Anti-mouse CD127/IL7R $\alpha$ BV605 (Clone A7R34)	BioLegend	Cat# 135041
Anti-mouse CD25 PE-Cy7 (Clone PC61)	BioLegend	Cat# 102016
Anti-mouse ICOS PE-Cy7 (Clone C398.4A)	BioLegend	Cat# 313520
Anti-mouse/human KLRG-1 FITC (Clone 2F1/KLRG1)	BioLegend	Cat# 138410
Anti-mouse/human KLRG-1 Percp-Cy5.5 (Clone 2F1/KLRG1)	BioLegend	Cat# 138417
Anti-mouse PD1/CD279 PE-Cy7 (Clone 29F.1A12)	BioLegend	Cat# 135216
Anti-mouse PD1/CD279 FITC (Clone 29F.1A12)	BioLegend	Cat# 135213

Anti-mouse CCR6 PE (29-2L17)	BioLegend	Cat# 129804
Anti-mouse CD103 FITC (2E7)	BioLegend	Cat# 121420
Anti-mouse CD45.1 FITC (Clone A20)	BioLegend	Cat# 110706
Anti-mouse CD45.2 BV711 (Clone 104)	BioLegend	Cat# 109847
Anti-mouse CD45.2 Pacific Blue (Clone 104)	BioLegend	Cat# 109820
Anti-mouse CD45 AF-700 (Clone 30-F11)	BioLegend	Cat# 110724
Anti-mouse CD3 AF-700 (Clone 17A2)	BioLegend	Cat# 100215
Anti-mouse CD3 BV711 (Clone 17A2)	BioLegend	Cat# 100241
Anti-mouse IL-5 PE (Clone TRFK5)	BioLegend	Cat# 504304
Anti-mouse IL-17A AF700 (Clone TC11-18H10.1)	BioLegend	Cat# 506914
Anti-mouse IL13 AF488 (Clone TWAJ)	eBioscience	Cat # 53-7133-82
Anti-mouse GATA-3 eF660 (Clone TWAJ)	eBioscience	Cat # 50-9966-42
Anti-mouse GATA-3 PerCP-eF710 (Clone TWAJ)	eBioscience	Cat # 46-9966-42
Anti-mouse Eomes AF488 (Clone Dan1 1mag)	eBioscience	Cat # 53-4875-82
Anti-mouse IL25R/IL17RB eF660 (Clone MUNC33)	eBioscience	Cat # 50-7361-82
Anti-mouse IL18R $\alpha$ PE-Cy7 (Clone P3TUNYA)	eBioscience	Cat # 25-5183-82
Anti-mouse NKp46 / CD335 PercP-eF710 (29A1.4)	eBioscience	Cat # 46-3351-82
Anti-mouse/human TCF1/7 AF488 (Clone C63D9)	Cell Signalling	Cat # 6444S
Anti-mouse IL33R/ST2 BV421 (Clone U29-93)	BD Biosciences	Cat# 566309
Anti-mouse ROR $\gamma$ t PE (Clone Q31-378)	BD Biosciences	Cat# 562607
Anti-mouse ROR $\gamma$ t AF647 (Clone Q31-378)	BD Biosciences	Cat# 562682
Anti-mouse ROR $\gamma$ t BV650 (Clone Q31-378)	BD Biosciences	Cat# 564722
Anti-mouse PLZF PE (Clone Q31-378)	BD Biosciences	Cat# 564850
Anti-mouse Ki67 PerCP-Cy5.5 (Clone B56)	BD Biosciences	Cat# 561284
Anti-mouse CTLA-4 PE (Clone UC10-4B9)	BD Biosciences	Cat# 553720
Anti-mouse CD121b PE (Clone 4E2)	BD Biosciences	Cat# 554450
Anti- mouse ICOS (CD278) BUV395 (Clone 398.4A)	BD Biosciences	Cat# 565884
Anti-mouse CD69 BUV805 (Clone H1.2F3)	BD Biosciences	Cat# 741927
Anti-mouse KLRG-1 BUV661 (Clone 2F1)	BD Biosciences	Cat# 741586
Anti-mouse CD279 (PD-1) BUV737 (Clone RMP1-30)	BD Biosciences	Cat# 749306
Anti-mouse CD90.2 BU496 (Clone 30-H12)	BD Biosciences	Cat# 741047

Anti-mouse LPAM-1(a4b7) BV711 (Clone DATK32)	BD Biosciences	Cat# 740701
Anti-mouse CD103 BV786 (Clone M290)	BD Biosciences	Cat# 564322
Anti-mouse CD25 PE-Dazzle 594 (Clone PC61)	BD Biosciences	Cat# 562694
Anti-mouse CD3e PE-Cy5 (Clone 145-2C11)	BD Biosciences	Cat# 553065
Anti-mouse CD45.2 BUV737 (Clone 104)	BD Biosciences	Cat# 612778
Anti-mouse CD45 BUV496 (Clone 30-F11)	BD Biosciences	Cat# 749889
Anti-mouse IL33R/ST2 PE (Clone U29-93)	BD Biosciences	Cat# 566311
Anti-mouse CD90.2 BV605 (Clone 30-H12)	BioLegend	Cat# 140317
Anti-mouse CD45 BV510 (Clone 30-F11)	BioLegend	Cat# 103137
Anti-mouse CD45.2 FITC (Clone 104)	BioLegend	Cat# 109806
Anti-mouse CD45.1 Percp-Cy5.5 (Clone A20)	BioLegend	Cat# 110728
Anti-mouse CD4 PE-Texas Red (Clone RM4-5)	eBioscience	Cat # MCD0417
Anti-mouse CD8a PE-Texas Red (Clone 5H10)	eBioscience	Cat # MCD0817
Anti-mouse I-A/I-E AF-700 (Clone M5/114.15.2)	BioLegend	Cat# 107621
Anti-mouse TCR $\gamma\delta$ Biotin ( Clone GL3)	BioLegend	Cat# 118103
Anti-mouse Epcam Biotin (Clone G8.8)	BioLegend	Cat# 118203
Anti-mouse CD8 $\alpha$ Biotin (Clone 53.67)	BioLegend	Cat# 100704
Anti-mouse CD3 Biotin (Clone 17A2)	BioLegend	Cat# 100243
Anti-mouse CD4 Biotin (Clone GK1.5)	BioLegend	Cat# 100404
Anti-mouse CD45 FITC (Clone 30-F11)	BioLegend	Cat# 103107
Anti-mouse NK1.1 PE-Cy7 (Clone PK136)	BioLegend	Cat# 108713
Anti-mouse CD19 FITC (Clone MB19-1)	BioLegend	Cat# 101505
Anti-mouse CD3 $\epsilon$ FITC (Clone 145-2C11)	BD Bioscience	Cat# 563061
Anti-mouse CD4 FITC (Clone H1 29.19)	BD Bioscience	Cat# 553651
Anti-mouse CD8a FITC (Clone 53-6.7)	BD Bioscience	Cat# 553031
Anti-mouse Gr1 FITC (Clone R86-8C5)	BD Bioscience	Cat# 553126
Anti-mouse Siglec-H FITC (Clone 551)	BioLegend	Cat# 129603
Anti-mouse B220 FITC (Clone RA3-6B2)	BD Bioscience	Cat#553088
Anti-mouse c-Kit/CD117 BV510 (Clone ACK2)	BioLegend	Cat#135119
Anti-mouse Sca-1 BV650 (Clone D7)	BioLegend	Cat#108143

Anti-mouse Flt3/CD135 PerCP-eFluor 710 (Clone A2F10)	eBioscience	Cat#46-1351-80
Anti-mouse CD27 PE (Clone LG3A10)	BD Bioscience	Cat# 558754
Anti-mouse Thy1.2 AF-700 (Clone 30-H12)	BioLegend	Cat# 105320
Anti-mouse ICOS BV785 (Clone C398.4A)	BioLegend	Cat# 313533
Anti-mouse IL18R $\alpha$ AF647 (Clone BG/IL18RA)	BioLegend	Cat# 132903
Anti-mouse IL33R/ST2 (ST2) PE-Cy7 (Clone RMST2-2)	eBioscience	Cat#25-9335-82
Streptavidin BV510	Biolegend	Cat#405233
Anti-mouse IL25R/IL17RB PE (Clone 9B10)	BioLegend	Cat#146305
Anti-mouse CD127/IL7Ra BV421 (Clone A7R34)	BD Biosciences	Cat# 566377
Anti-mouse TCR $\gamma\delta$ Antibody APC (Clone GL3)	BioLegend	Cat#118115
Anti-mouse/human CD11b FITC (Clone M1/70)	eBioscience	Cat# 11-0112-82
Anti-mouse TER-119 FITC (Clone TER-119)	eBioscience	Cat# 11-5921-82
Anti-human CD14 APC-eF780 (Clone 61D3)	eBioscience	Cat# 47-0149-42
Anti-human CD19 APC-eF780 (Clone HIB19)	eBioscience	Cat# 47-0199-42
Anti-human CD123 APC-eF780 (Clone 6H6)	eBioscience	Cat# 47-1239-42
Anti-human CD3 APC-eF780 (Clone SK7)	eBioscience	Cat# 47-0038-42
Anti-human Fc $\epsilon$ R1a APC Vio770 (Clone CRA1)	Miltenyi	Cat# 130-105-500
Anti-human CD127 PE (Clone MB15-18C9)	Miltenyi	Cat# 130-113-410
Anti-human CD218(IL18Ra) PEVio770 (Clone H44)	Miltenyi	Cat# 130-101-723
Anti-human ICOS APC (Clone REA192)	Miltenyi	Cat# 130-101-723
Anti-human EOMES FITC (Clone WD1928)	eBioscience	Cat# 11-4877-42
Anti-human CRTH2 PerCP-Cy5.5 (Clone BM16)	BioLegend	Cat# 350116
Anti-mouse/human T-bet BV605 (Clone 4B10)	BioLegend	Cat# 644817
Anti-human CD117(c-KIT) BV650 (Clone 104D2)	BioLegend	Cat# 313222
Anti-human CD103 BV711 (Clone Ber-ACT8)	BioLegend	Cat# 350222
Anti-human CD336(NKP44) Biotin (Clone P44-8)	BioLegend	Cat# 325106
Streptavidin BV785	BioLegend	Cat# 405249

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### Experimental Models: Organisms/Strains

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Mouse: wild-type:	C57BL/6, Charles River Laboratories
Mouse: CD45.1/Ly5.1	B6.SJL-Ptprca Pepcb/BoyJ, Jax mice JAX stock #006584

Mouse: <i>Ubiquitin</i> <sup>tdT</sup> (`Ubow`)	(Ghigo et al., 2013)
Mouse: IL5cre x LSLtdTom	(Nussbaum et al, 2013)
Mouse: RAG1 <sup>-/-</sup>	B6.129S7-Rag1tm1Mom/J JAX stock #002216
Mouse: RAG2 <sup>-/-</sup> $\gamma$ c <sup>-/-</sup>	C57BL/6NTac;B10(Cg)-Rag2 <sup>tm1Fwa</sup> Il2rg <sup>tm1Wjl</sup> , Taconic, 4111-F

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## Software and Algorithms

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FlowJo v.10.5.0	FlowJo	<a href="https://www.flowjo.com">https://www.flowjo.com</a>
Prism v.6.0.	GraphPad	<a href="https://www.graphpad.com">https://www.graphpad.com</a>
R v. 3.5.1 and 3.6.1		<a href="https://www.r-project.org/">https://www.r-project.org/</a>

## CONTACT FOR REAGENT AND RESOURCE SHARING

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, D. Grün( [gruen@ie-freiburg.mpg.de](mailto:gruen@ie-freiburg.mpg.de) ) and G. Gasteiger ([georg.gasteiger@uni-wuerzburg.de](mailto:georg.gasteiger@uni-wuerzburg.de))

## EXPERIMENTAL MODEL AND SUBJECT DETAILS

### Mice

C57BL/6 mice were purchased from Charles River Laboratories. All mice (IL5<sup>Cre</sup> x LSL-tdTom mice, RAG1<sup>-/-</sup> mice, RAG2<sup>-/-</sup>  $\gamma$ c<sup>-/-</sup> mice, etc.) were bred and housed under specific pathogen free conditions at the animal facilities of the University of Würzburg (ZEMM and Institute for Immunology and Virology), Max Planck Institute of Immunobiology and Epigenetics, Memorial Sloan-Kettering Cancer Center and Centre Immunologie Luminy and were used in accordance with institutional guidelines and as approved by the local authorities. For all experiments except for shield chimeras, female mice at 6-8 weeks of age were used.

## METHOD DETAILS

### Parabiosis

Congenic CD45.1 and CD45.2 mice were surgically connected in parabiosis as previously described (Gasteiger et al., 2015). After corresponding lateral skin incisions were made from elbow to knee in each mouse, forelimbs and hindlimbs were tied together using nylon suture, and the skin incisions were closed using stainless steel wound clips. After surgery, mice were maintained on a diet supplemented with trimethoprim-sulfamethoxazole for prophylaxis of infection.

### Shield chimeras

Shield irradiation was performed as described before (Baratin et al., 2017; Scott et al., 2016). Briefly, mice were placed in a 6mm thick lead cylinder that only exposed their hind legs, irradiated (9Gy) and reconstituted i.v. with BM cells from *Ubiquitin*<sup>tdT</sup> (`Ubow`) (Ghigo et al., 2013) donors. Newborn (1-2 days old) mice were transplanted with BM depleted of monoblasts by magnetic depletion of CD11b<sup>+</sup> cells. Adult mice were anesthetized with a Ketamine/Xylazine mixture during irradiation and received total BM prepared using standard procedures. In both setups, BM cells were transplanted at a ratio of 1 donor per recipient.

### ivCD45 labeling and cell isolation

2 $\mu$ g anti-mouse CD45 AF-700 antibody were injected via retroorbital vein (200 $\mu$ l/mouse) after mice were anesthetized with intraperitoneal injection of ketamine/xylazine. Mice were sacrificed 5 mins after antibody injection. Peripheral blood was taken from the left ventricle and bone marrow cells were prepared using standard procedures. To isolate mononuclear cells from lung, the tissues were physically dissociated using scissors and incubated for 45 minutes in 2ml DMEM (10mM HEPES) digestion buffer (0.25mg/ml Liberase LT, Roche) before being passed through 70- $\mu$ m strainers. Red blood cells in blood and lung were lysed using ACK lysis buffer.

### **Helminth Infection**

Mice were injected subcutaneously (s.c.) with 350 L3 *Nippostrongylus brasiliensis* (*Nb*) larvae cultured as previously described (Symowski and Voehringer, 2019). Briefly, *Nb* were maintained by passage in rats. These were injected s.c. with 1000 L3 *Nb* and stool was collected on days 7-9 p.i.. Stool was mixed with water and activated charcoal in petri dishes and cultured at 25°C for up to 4 weeks in a chamber lined with wet paper towels to maintain humidity. L3 *Nb* were collected as needed using a Baermann apparatus after at least 7 days of culture. L3 larvae were extensively washed with PBS to eliminate culture contaminants before infection.

### **Murine cell isolation**

Lungs were harvested, chopped into small pieces and incubated in digestion buffer DMEM (Gibco 31966021); 0.25 mg/ml Liberase<sup>TM</sup> TL (Research Grade 05401020001, Roche); 10mM Hepes (Gipco 1563049) for 45 mins at 37°C on Eppendorf rotator at 120rpm. Digestion was stopped with 20ml RPMI 1640 medium Thermo Fisher Scientific; 31870074) added with 50 $\mu$ M 2-mercaptoethonal (AppliChem, A1108). Cells were filtered through 70 $\mu$ m cell strainer (Falcon 3522350) and washed and collected in FACS Buffer or 1 $\times$  MojoSort buffer (BioLegend; 480017). Bone marrow cells from ilia, femora, and tibiae were isolated by crushing the bones in FACS Buffer or 1 $\times$  MojoSort buffer. Bone marrow cells were filtered (CellTrics, 30  $\mu$ m; Sysmex) and washed and collected in FACS Buffer or 1x MojoSort buffer.

### **Human samples and cell isolation**

Peripheral blood (PB) leukocytes were isolated from buffy coats of healthy adults provided by the German Red Cross (DRK, Berlin). Lung samples were obtained at the Helios Klinikum, Berlin, from non-microscopically affected surgical excisions of lung tumor patients. All samples were processed within 24 hours. The institutional review boards of the Charité Universitätsmedizin Berlin approved the protocols and all probands gave their written informed consent according to the Declaration of Helsinki. All studies were approved by the Ethics Committee. Human mononuclear cell suspensions from PB and lung were isolated by Ficoll density gradient centrifugation. Lung tissues were thoroughly dissociated in sterile PBS/BSA/2mM EDTA and passed through a 70 $\mu$ m strainer prior to density gradient.

### **Antibody staining for sorting and flow cytometric analysis**

Cell suspensions were prepared as described above and incubated with Fc block (2.4G2, Biotcell) and Fixable Viability Dye, and were subsequently stained with fluorophore-conjugated antibodies against: CD3 $\epsilon$ , CD5, CD19, Gr, F4/80, CD11b, CD11c, NK1.1, TCR $\beta$ , TCR $\gamma\delta$ , Fc $\epsilon$ R1, TER-119, CD45R/B220, Ly6G, Thy1.2, CD127/IL7R $\alpha$ , CD25, IL25R/IL17RB, CD218a/IL18R $\alpha$ , IL33R/ST2, ICOS, KLRG-1, PD1/CD279,  $\alpha$ 4 $\beta$ 7, C-kit, CD45.1, CD45.2, CD45, CD3, IL-5, IL13, IL17A, NKp46, Eomes, ROR $\gamma$ t, Gata3, TCF1/7, PLZF, Ki67, IL1R2, I-A/I-E, CD103 and CTLA-4. Lineage (Lin) positive cells were stained with a cocktail of biotin-labeled antibodies against CD3 $\epsilon$ , NK1.1, CD5, Gr-1, CD19, TCR $\gamma\delta$ , TCR $\beta$ , CD11b, CD11c, F4/80, TER-119, and Fc $\epsilon$ R1, followed by staining with Streptavidin (V500, BD). For intracellular cytokine staining, cells were incubated in the presence of 1 $\mu$ g/ml monensin (BioLegend) or re-stimulated with 50ng/ml PMA and 500ng/ml Ionomycin in the presence of 1 $\mu$ g/ml monensin with/without 1 $\mu$ g/ml Brefeldin A for 3.5h in indicated experiment. Intracellular staining of CD3 $\epsilon$ , IL5, IL13, IL17A Ki67, IL1R2, CTLA-4, PLZF, Tcf7, Eomes, Gata3 and ROR $\gamma$ t was performed using Foxp3 / Transcription Factor Staining Buffer Set (eBioscience). To reserve tdTom from IL5<sup>Cre</sup> fate-mapped mice, cells are first fixed with 2% FA for 1h, and then washed and stained ROR $\gamma$ t with Foxp3/Transcription Factor staining buffer. Human cells, were incubated with antibodies against EOMES, CRTH2, TCF1/TCF7, T-bet, NKp44, CD117, CD103, ICOS, CD45, CD127, GATA3 and IL-



18R. In addition, Lin was stained using antibodies against CD14, CD19, CD123, CD3, FcεRIa. For flow cytometric analysis cells were analyzed on a BD Fortessa, BD Celesta or Cytex Aurora cytometer and analyzed with FlowJo software.

Pan-ILCs were sorted as live Lin<sup>-</sup> CD45<sup>+</sup> Il7ra<sup>+</sup> Thy1.2<sup>+/lo</sup> ILCs for scRNAseq. Additional markers were included for host and donor-derived cells (Parabiosis and Shield chimeras), intravascular cells (ivCD45-labeling) and neonatal lung (ST2).

For transfer and culture experiments, BM and lung ILCs were sorted as live Lin<sup>-</sup>CD45<sup>+</sup> Il18r1<sup>+</sup> Icos<sup>+</sup> Il7ra<sup>+</sup> Thy1.2<sup>+</sup> and further divided into subsets based on the expression of ST2 and/or Il17rb for culturing and input for sequencing. Moreover, CD45<sup>+</sup> Il7ra<sup>+</sup> Thy1.2<sup>+</sup> lung ILCs were sorted as Il18r1<sup>+</sup> ST2<sup>-</sup>, Il18r1<sup>+</sup> ST2-IL5FM<sup>+</sup>, Il18r1<sup>+</sup> Ccr6<sup>+</sup>, Il18r1<sup>+</sup> Ccr6<sup>-</sup> and Il18r1<sup>-</sup> ST2<sup>+</sup> subsets for in vitro cultures and transfer experiments as indicated in respective figures. Cells were enriched by negative selection and depletion of Lin<sup>+</sup> cells with anti-biotin Microbeads (Miltenyi) or (MojoSort Streptavidin Nanobeads, 480016, Biolegend) according to the manufacturer's recommendations before sorting.

Staining and enrichment of sorted Il18r1<sup>+</sup>Icos<sup>+</sup> BM ILCs for single-cell sequencing: BM cells were harvested as described above. Hematopoietic progenitors were enriched with the MojoSort mouse hematopoietic progenitor cell isolation kit (BioLegend; 480004) according to the manufacturer's protocol. After enrichment,  $5 \times 10^6$  to  $7 \times 10^6$  cells per 100  $\mu$ l of 1 $\times$  MojoSort buffer were stained on ice in the dark for 40 min with following fluorophore-conjugated antibodies against: c-Kit/CD117, Sca-1, Il18r1, Thy1.2, NK1.1, Icos, Flt3/CD135, CD127/Il7ra $\alpha$ , CD27 and the lineage markers: CD19, Siglec-H, CD3 $\epsilon$ , B220, Gr1, CD74, TER-119 and CD11b/Mac-1. Cells were sorted as live Lin<sup>-</sup>Il18r1<sup>+</sup>Icos<sup>+</sup> and index information was recorded for the other markers.

### **Adoptive cell transfer**

Lung ILCs were isolated and stained as above. Cells were derived from lungs of congenically marked B6 mice. 1500-2000 cells each were co-transferred into sub-lethally irradiated (3Gy) RAG<sup>-/-</sup> $\gamma$ c<sup>-/-</sup> mice. ILCs were analyzed 3 or 6w post transfer as indicated in the Figure legends.

### **Cell culture of sorted ILCs subsets**

Sorted ILCs were seeded onto mitomycin-C-treated OP9 or OP-DL1 feeder cells. Approximately 100 cells were plated in 1 well of a 96-well plate and grown at 5% CO<sub>2</sub> in RPMI 1640 medium (complete formulation; Thermo Fisher Scientific; A1049101) supplemented with 10% fetal calf serum (Corning BV; 35-016-CV), 1 $\times$  penicillin–streptomycin (Gibco; 10378016) and 25 ng/ml IL-2 (BioLegend; 575402), 25ng/ml IL-7 (BioLegend; 577802), complemented with different combinations of the following cytokines at 10ng/ml IL-1 $\beta$  (BioLegend; 575102), IL-23 (BioLegend; 589002), IL25 (R&D, 1399-IL-025/CF), IL33( Biolegend, 580506) and 25ng/ml SCF(Peprotech, 250-03) where indicated. One day after sorting and then every other day, 100  $\mu$ l of medium was replaced with fresh medium containing cytokines. For clonal analysis of progenitor cell potential, single cells were sorted directly onto OP9-DL1 monolayers and cultured for 19 days when FACS analysis was performed. Cultures were considered to be positive for a cell population if > 20 events appeared within the indicated FACS gates.

### **Preparation of OP9 feeder cell layer**

OP9 feeder cells were grown in 1 $\times$  MEM Alpha (Gibco; 12561-056) supplemented with 20% fetal calf serum (Corning BV; 35-016-CV), 1 $\times$  penicillin–streptomycin (Gibco; 10378016). OP9 cells were grown until 80–90% confluent and treated with 10  $\mu$ g/ml mitomycin C (Sigma-Aldrich; M0503-2MG) for 1.5 h. Cells were washed four times with PBS and detached with trypsin–EDTA (Gibco; 25200056) at 37 °C for 5 min. Cells were resuspended in fresh complete MEM Alpha medium, and cell concentration was adjusted to seed approximately 40,000–50,000 cells in 100  $\mu$ l of medium per well of a 96-well plate.

### **Single cell sorting, single-cell RNA amplification and library preparation of ILCs.**

Single cells were FACS sorted into 384-well plates on a BD AriaIII, BD Influx cell sorter or BD Aria Fusion using single-cell mode and index information was recorded (primary cells) or sorted on BD Aria Fusion using 4-way purity mode (output cultured cells). After sorting, plates were centrifuged for 5 min at 2200g at 4°C, snap-frozen in liquid nitrogen and stored at -80°C until processed. Single cell RNA sequencing was performed using the CEL-Seq2 method (Hashimshony et al., 2016) with modifications as described in (Herman et al., 2018). ILCs were sequenced on a HiSeq 2500 or HiSeq 3000 sequencing system (Illumina paired-end multiplexing run, high output mode) at a depth of ~72,000~600,000 reads per cell.

### Quantification of transcript abundance

Paired end reads were aligned to the transcriptome using bwa (version 0.6.2-r126) with default parameters (Li and Durbin, 2010). The transcriptome contained all gene models based on the mouse ENCODE VM9 release downloaded from the UCSC genome browser comprising 57,207 isoforms with 57,114 isoforms mapping to fully annotated chromosomes (1 to 19, X, Y, M). All isoforms of the same gene were merged to a single gene locus. Furthermore, gene loci overlapped by >75% were merged to larger gene groups. This procedure resulted in 34,111 gene groups. As described in (Wallrapp et al., 2017) the loci for *Nmur1* was extended at the 3' end to position 86,384,000 on the chromosome 1. The right mate of each read pair was mapped to the ensemble of all gene loci and to the set of 92 ERCC spike-ins in sense direction (Baker et al., 2005). Reads mapping to multiple loci were discarded. The left read contained the barcode information: The first six bases corresponded to the unique molecular identifier (UMI), followed by six bases representing the cell specific barcode. The remainder of the left read contained a polyT stretch. The left read was not used for quantification. For each cell barcode, the number of UMIs per transcript was counted and aggregated across all transcripts derived from the same gene locus. Based on binomial statistics, the number of observed UMIs was converted into transcript counts (Grün et al., 2014).

### Clustering

The datasets were analyzed using RaceID3 (v0.1.3 and v0.1.6). Rescaling to 3,000 transcripts per cells was used for data normalization (2,000 for the parabiosis dataset due to lower sample quality). Prior to filtering and normalization, ERCCs and mitochondrial genes were excluded and cells expressing >2% of *Kcnqlot1* transcripts, identified as a marker of low-quality cells (Grün et al., 2016) were removed from analysis. RaceID3 was run with the following parameters: mintotal= 3,000, minexpr=5, outminc=5, FSelect=TRUE, probthr=10<sup>-3</sup>, ccor = 0.65 (probthr=10<sup>-4</sup>, for the parabiosis and shield chiemera dataset, and ccor=0.4, mintotal= 2000, minexpr=3, outminc=3 for the parabiosis dataset). In addition, we used imputing implemented in RaceID3 with knn=10. For the BM dataset cln=6 was set, since cluster dispersion decreases linearly after 6 clusters. For cultured lung Il18r1+Icos+ ILCs outlier identification was omitted. To remove batch effect and technical variability we initialized CGenes. CGenes and highly correlating genes (Pearson's correlation > ccor) were removed from clustering, representing a minimal invasive batch effect removal approach. The following sets of genes were initialized as CGenes: ensemble of all *Gm*, *RP*, *Hsp*, *Igl*, *Igh* and *A4300 genes*, *Scgb1a1*, *Jchain*, *Igkc*, *Malat1*, *Xist*, *Mid1* and *Kcnqlot1*. To remove cell-cycle associated variability from the uninfected lung dataset we integrated cell cycle related genes (Buettner et al., 2015) and highly correlating genes (Pearson's correlation >= 0.4) into CGenes. As a large fraction of neonatal cells expressed proliferation associated genes with a dominant effect on cluster, *Mki67*, *Pcna*, *Tuba1a/Tuba1b*, *Top2a*, *Tubb5* were additionally included into CGenes. We used the internal RaceID batch correction method for the *in vitro* culture datasets. Briefly, this method identifies a pair of cells across the Il18r1+Icos+ and the Il18r1+Icos+ST2- batches with minimal k nearest neighborhoods distance and infers differentially expressed genes between cells of these two neighborhoods. Thus, *Gzmb*, *Calca* and *Ucp2* were additionally included in CGenes to remove batch effect associated variability for this dataset. For the parabiosis dataset, *Lars2*, *Mki67* and *Pcna* were included into CGenes, to remove batch effects or cell cycle associated variability. For contaminant cell removal, cells within clusters with signatures for non-ILCs, NK-cells or low quality cells (Grün et al., 2016) were removed from the dataset after initial clustering and RaceID3 was rerun. The single-cell RNA-seq data of uninfected adult lung comprised (after removal of a cluster of cells highly expressing natural killer (NK) cell markers such as *Nkg7*, *Klrd1* and *Klrc2*) 1,043 ILC transcriptomes with a median count of 4,470 transcripts and a median of 1,780 different genes per cell. Neonatal lung ILC data encompassed 366 cells with a median of 7,047 transcripts and 2,546 different genes per cell.

Il18r1+Icos+ BM cells consisted of 230 transcriptomes with a median transcript count of 6,372 transcripts and a median of 2,609 different genes per cell. Cultured lung and input cells (related to Figure 3G-I) comprised of 436 transcriptomes with a median of 5,035 transcripts and 1,908 different genes per cell. The Nb time-course dataset comprised 3200 ILC transcriptomes(excluding uninfected lung data) with median counts of 6,451 transcripts and 2,147 different genes after removal of NK cell and macrophage contaminants. The parabiosis dataset comprised 2,500 ILC transcriptomes, with a median count of 4,249 transcripts and 1,252 different genes after removal of NK cell and macrophage contaminants. In total, 1,422 cells were derived from host, and 1,078 cells from the enriched donor populations of cells. The shield chimera dataset comprised 2,281 ILC transcriptomes(995 host derived and 1,286 donor derived) with a median count of 5,729 transcripts and 2,047 different genes after removal of macrophage contaminations and low quality cells (Grün et al., 2016).

### **Cell-cycle Scoring**

To evaluate the impact of proliferation signature on clustering we adapted Seurat's cell cycle scoring. Genes passing filtering of the uninfected lung dataset were defined as universe. S phase and G2/M phase genes proposed by Tirosh and colleagues (Tirosh et al., 2016), which were elements of universe, were utilized. Genes of universe were binned into 24 bins based on the mean expression across cells. For every cell-cycle gene 100 genes were sampled from the respective bin and combined to a control gene set, containing only unique genes. Mean expression across all genes of the control gene set within each cell was subtracted from mean expression across combined S phase and G2/M phase gene sets within each cell. Cells were considered as cycling if this score was >0 and non-cycling if the score was <0. S phase and G2/M phase genes were combined as we detected correlation between genes of the two gene sets. To compare cell-cycle score with and without removal of cell-cycle variability in uninfected lung (see above), universe before removal of cell-cycle variability was used.

### **Differential expression analysis**

Differentially expressed genes between two subgroups of cells were identified similar to a previously published method (Anders and Huber, 2010). First, negative binomial distributions reflecting the gene expression variability within each subgroup were inferred based on the background model for the expected transcript count variability computed by RaceID3. Using these distributions, a p-value for the observed difference in transcript counts between the two subgroups was calculated. Multiple testing correction was performed using the Benjamini-Hochberg method and referred to as "adjusted p-value" (Herman et al., 2018).

### **Lineage inference, pseudo-temporal ordering and SOM**

For derivation of differentiation trajectories the StemID2 algorithm was used. (Grün et al., 2016; Herman et al.,2018). StemID2 was run with the following parameters: cthr=20 (for shield chimera cthr = 10), pdishuf=2000, pthr=0.01 and nmode=TRUE. Pseudo-temporal order was derived based on StemID2 projection coordinates of highest projection of cell to medoid links onto inter-cluster links of the subsequent cluster along the selected trajectories (uninfected lung: 5>11>9>8>4>10, 5>11>9>8>4>12 & time-course: 12>5>2>8>6>16, 12>5>2>8>4>23, 15>2>8>6>16). Self-organizing maps (SOMs) were used to infer models of pseudo-temporal expression profiles using the following parameters: nb=500, alpha=0.5, corthr=0.75, minsom=10.

### **KEGG/GO pathway enrichment**

GO and KEGG enrichment analysis was performed using the clusterProfiler R package (Yu et al., 2012). Genes of SOM inferred gene modules along the respective trajectory were used as input and the ensemble of all genes, excluding ERCCs and mitochondrial genes, were used as universe. In addition the following parameters were used: pAdjustMethod = "BH", pvalueCutoff = 0.01, qvalueCutoff = 0.05.

### **Fraction dot plot**

For expression color code, log<sub>2</sub> mean expression above for 4 or below -4 are replaced by 4 and -4, respectively. Z-scores above 1 and -1 are replaced by 1 and -1, respectively.

### **Normalisation of pie charts**

For pie charts, sample composition of RaceID3 clusters was normalized by dividing the number of cells within a cluster of the respective sample by the total number of cells of the respective sample in the dataset.

### **Enrichment scores**

Enrichment scores in RaceID3 clusters were obtained by dividing the relative frequency of donor or host cells within a cluster by the expected frequency, based on the frequency of donor and host cells in the total population. P-values of enrichments were calculated by a one-sided fisher-test using the *R fisher.test* function in R. For combined enrichment score and P-value calculation for shield chimera cluster 16 and 17, respective numbers of donor and host cells were aggregated

### **Dataset integration**

Integration of I118r1+Icos+ BM ILCs with ILCs and ILC2Ps from Walker et al.(Walker et al., 2019) Initially, RaceID3 was run on the filtered raw counts for cells from cluster 1 and 2 of Walker et al., representing ILCs and ILC2s, respectively. Cells with expression of *Cd302*, *Sox13*, *Cd19*, *Spi1* and *Gata1* were considered contaminants and removed prior to further analysis. For data set integration, the normalized counts of the remaining cells from cluster 1 and 2 of Walker et al. for genes shared with our I118r1+Icos+ BM dataset were used as feature set. For this purpose, Ensembl gene IDs from Walker et al., were matched with their respective gene symbol. If two or more Ensembl IDs were matching the same symbol, their counts were aggregated per cell, if the differences of their aggregated counts across cells was  $\geq 1$  or averaged per cell if their difference across cells was  $< 1$ . Dataset integration was performed using Seurat's standard workflow (Stuart et al. Cell 2019) with `nfeatures=3000` as argument for the "FindVariableFeatures" function. Variable features passing RaceID3 filtering (`minexpr=5`, `minnumber=1`, CGenes initialization as described above with `ccor=0.65`) were used for anchoring and integration. The resulting integrated object was subjected to RaceID3 distance matrix calculation and clustering.

Integration of uninfected lung ILCs with lung and skin ILCs from Ricardo-Gonzalez et al.(Ricardo-Gonzalez et al., 2018):

Before integration, cells of our uninfected lung dataset from Figure 1 (after removal of NK cells) and lung and skin ILCs from Ricardo-Gonzalez et al., with less than 2000 transcripts or less than 200 different genes were discarded. Genes which were unique to any of the two datasets were removed and Ensembl gene ID counts matching the same gene symbol were treated as described above. Uninfected lung cells were assigned to batch1, whereas lung and skin cells from Ricardo-Gonzalez et al. were assigned to batch2. Dataset integration was performed using the SCTransform function from the Seurat package with selection of 3000 integration features. Integration features were filtered by removal of CGenes and correlated genes as described before. In addition, cell-cycle genes from Buettner et al.(Buettner et al., 2015) and correlating genes (correlation  $\geq 0.4$  across uninfected lung cells) were removed from integration feature set, which was subsequently subjected to anchoring and integration. Clustering was performed using Seurat, using 30 principle components and the resolution factor 1.2 for clustering.

### **Cluster mapping: Inferring transcriptome similarities between two scRNA-seq datasets based on cluster annotation**

The unbiased identification of cells sharing a similar transcriptome in different tissues can be formulated as a classification problem: given a set of classifiers, i.e. cluster medoids in one dataset (e.g., parabiosis data with 22 clusters), we would like to identify non-negative weights reflecting similarities for each single cell in a given query (e.g., a single cell in the time-course data) under the constraint that the weights sum up to one. Hence, if a cell in the query set does not resemble a distinct cluster in the classifier set, it will have uniform weights for all clusters in the classifier set. On the other hand, if a cell resembles a specific cluster, the corresponding weight will be close to one, and all other weights will be close to zero. This optimization problem can be solved by quadratic programming, which was implemented using the `solve.QP` function from the `quadprog` R package.

**Accession Codes**

scRNA-seq data generated in this study are deposited in Gene Expression Omnibus (GEO) under accession code GSE141330.

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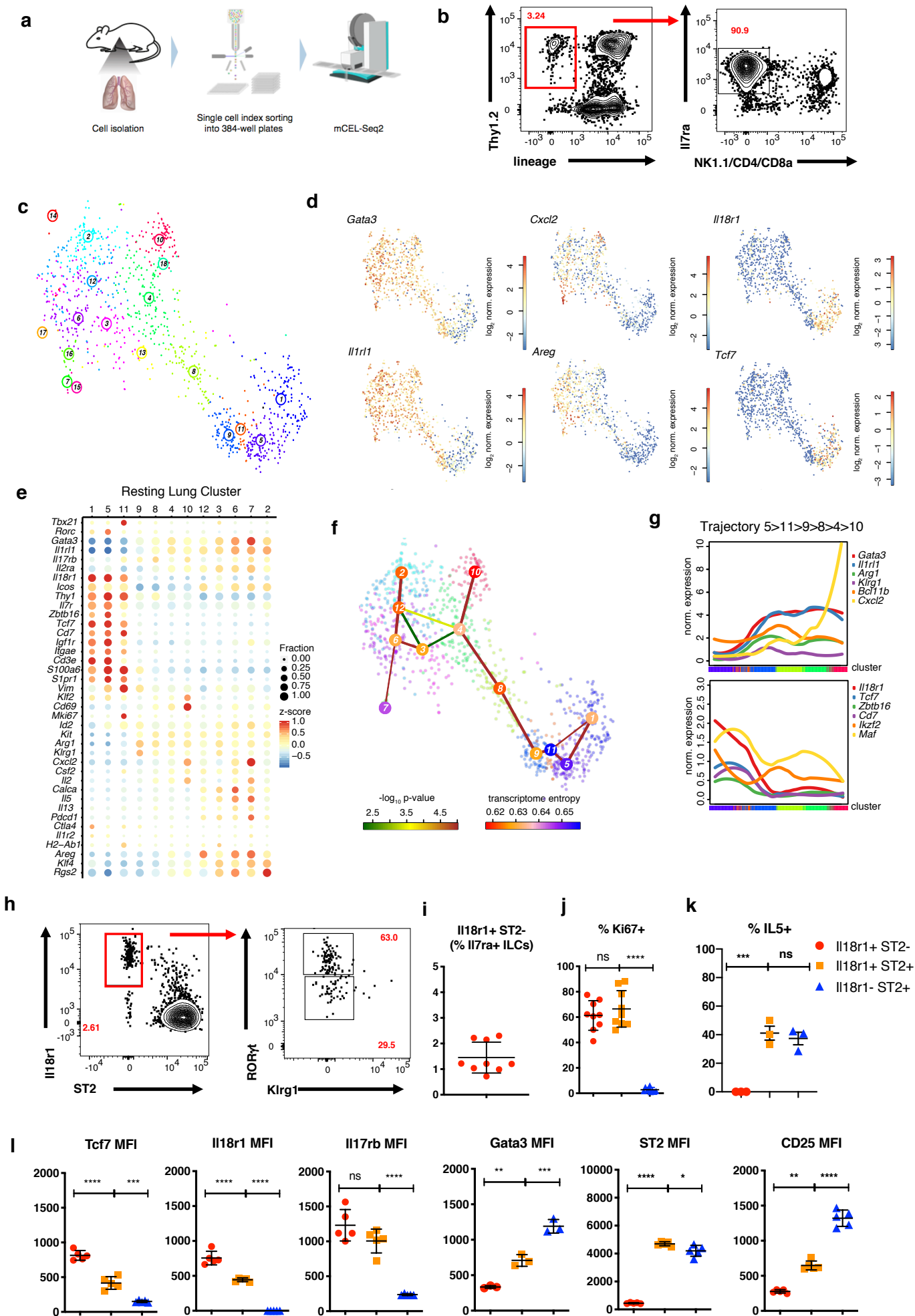
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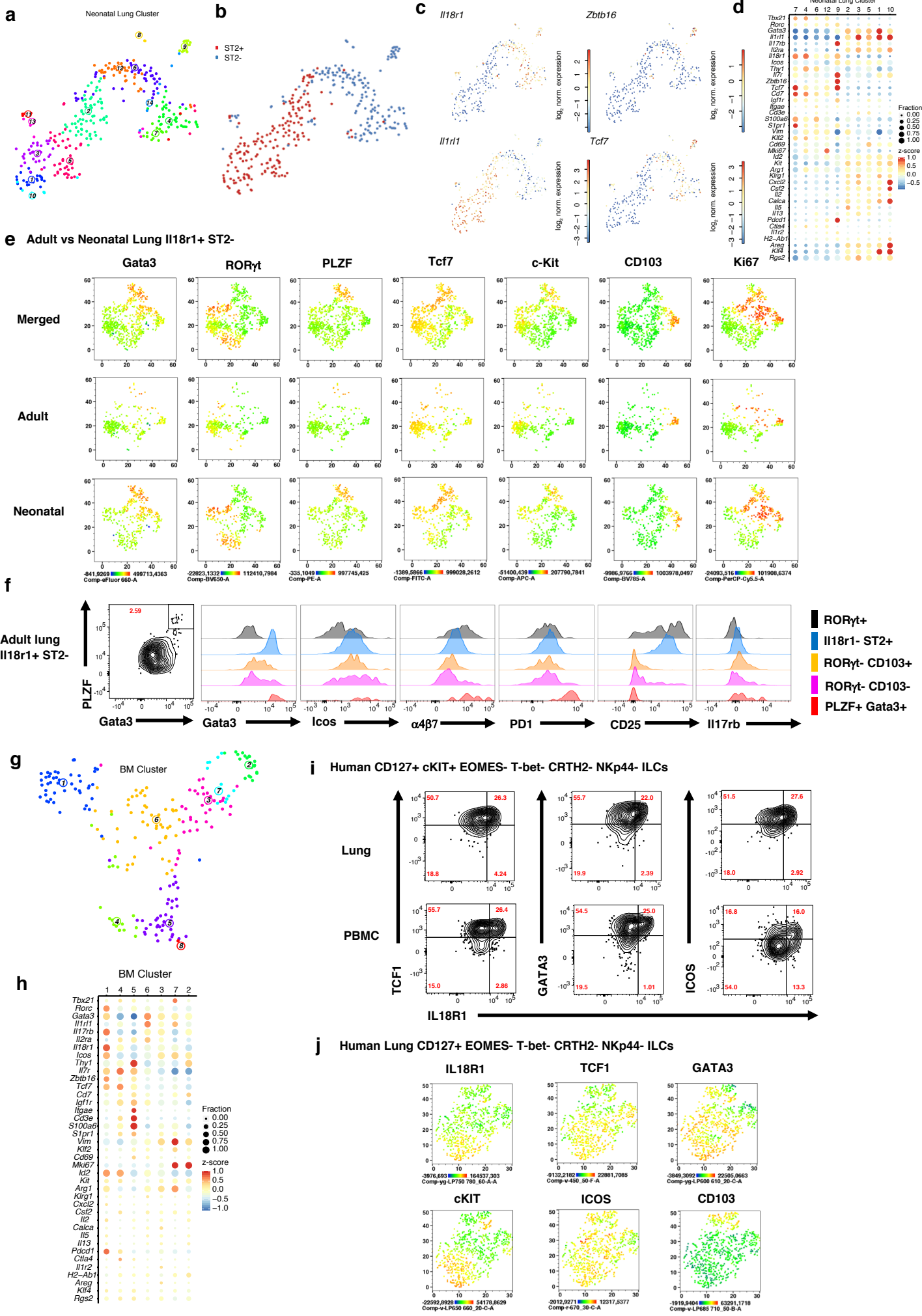
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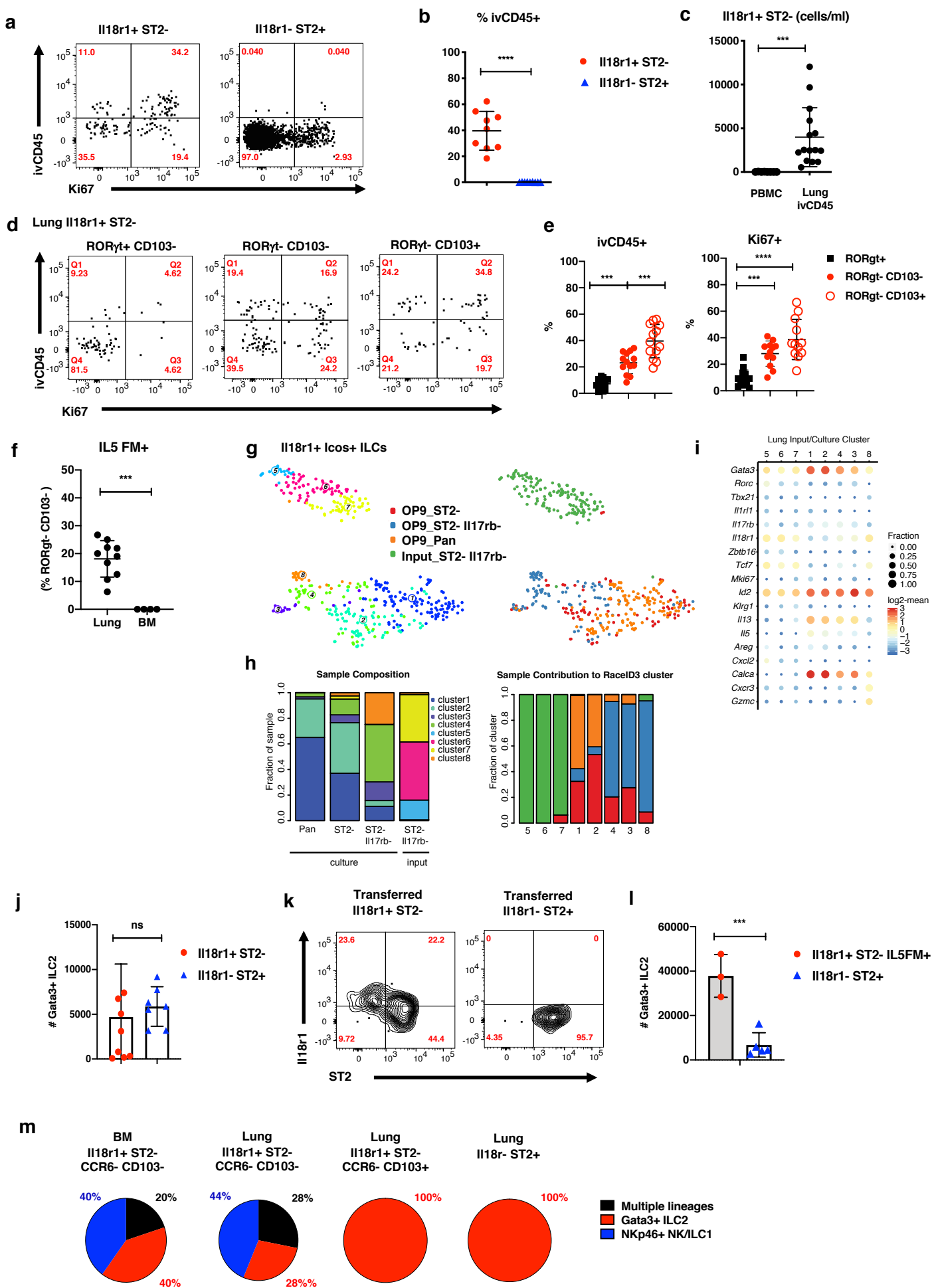


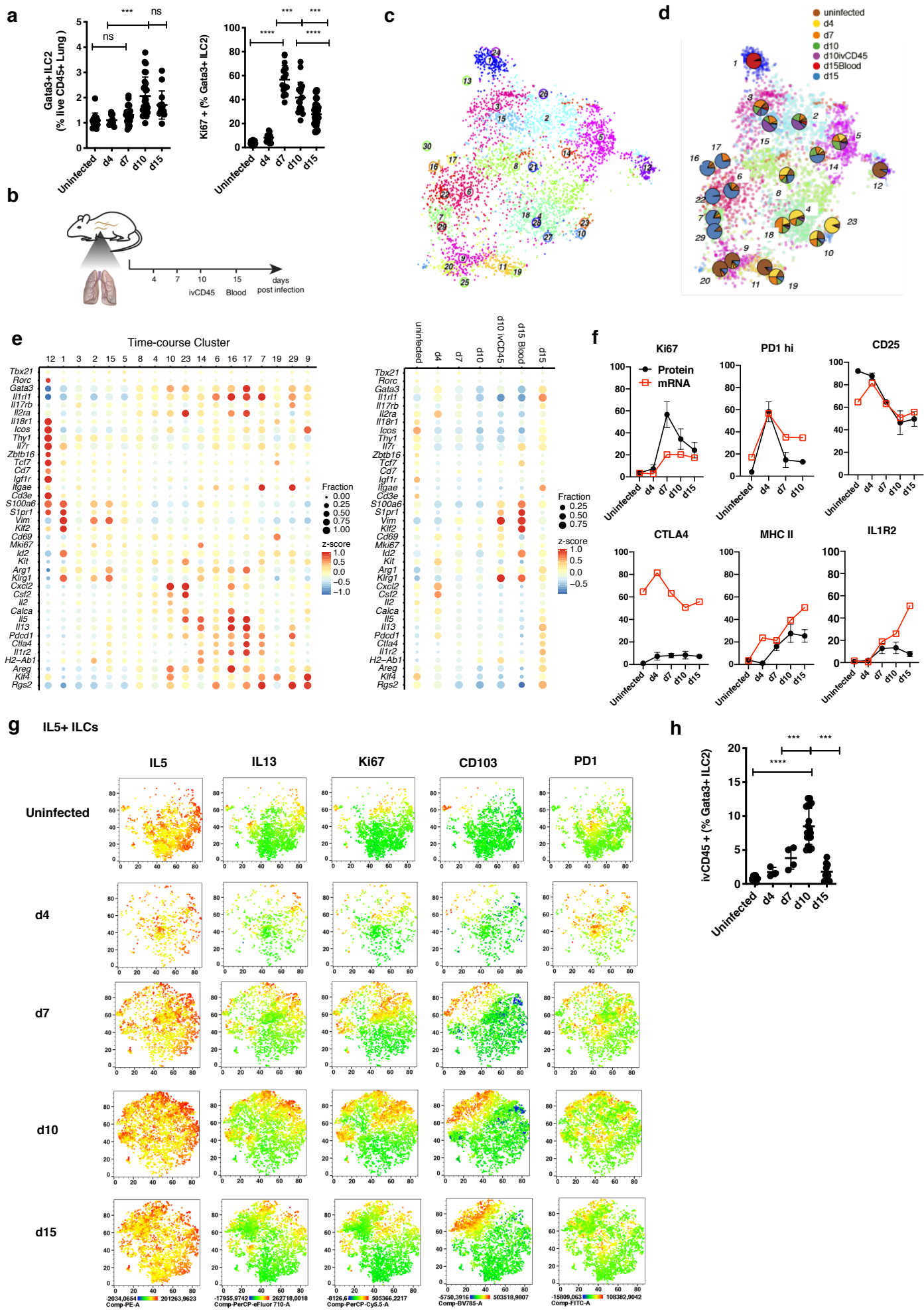
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**Figure 1**

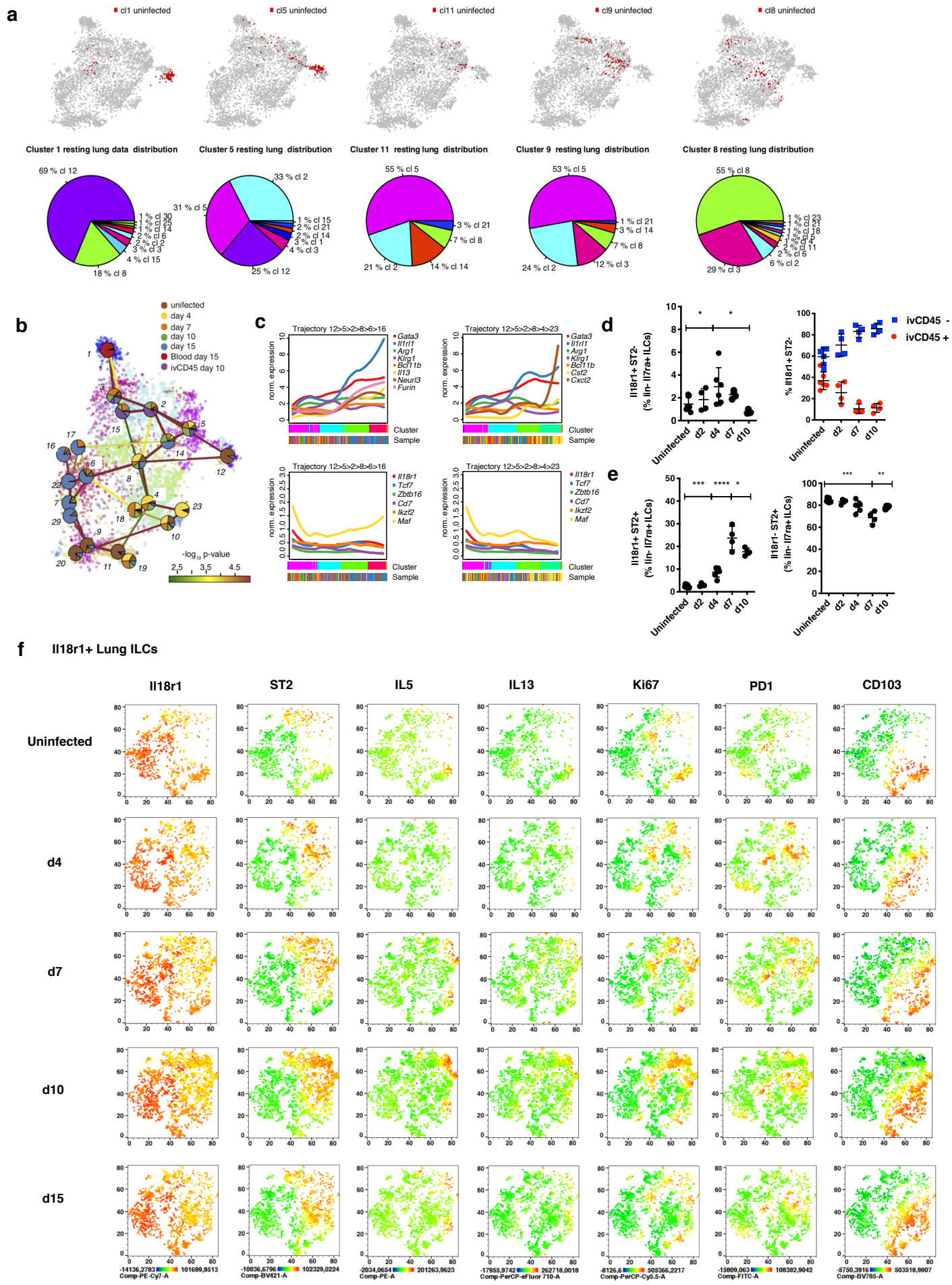
**Figure 2**

**Figure 3**

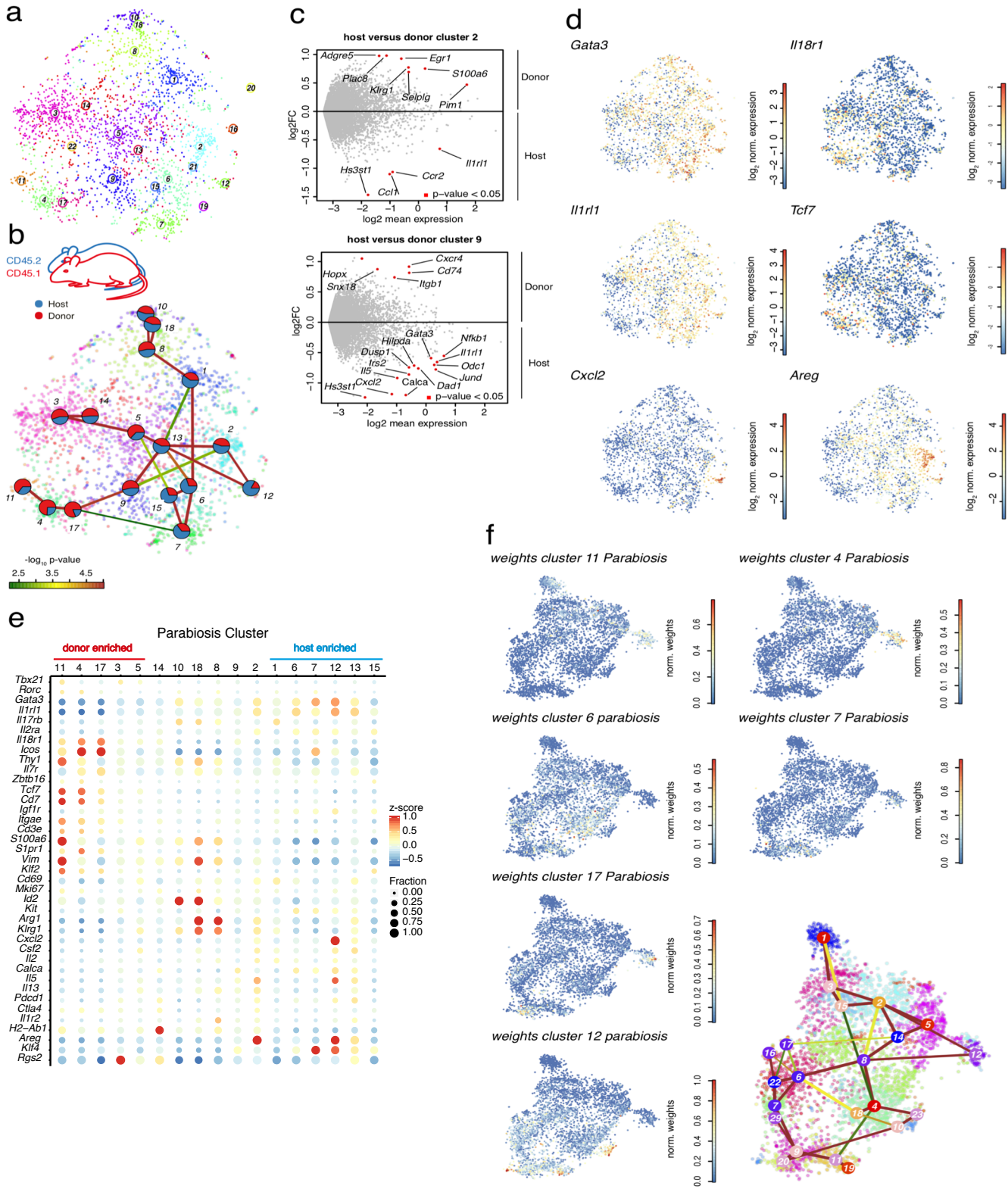


**Figure 4**

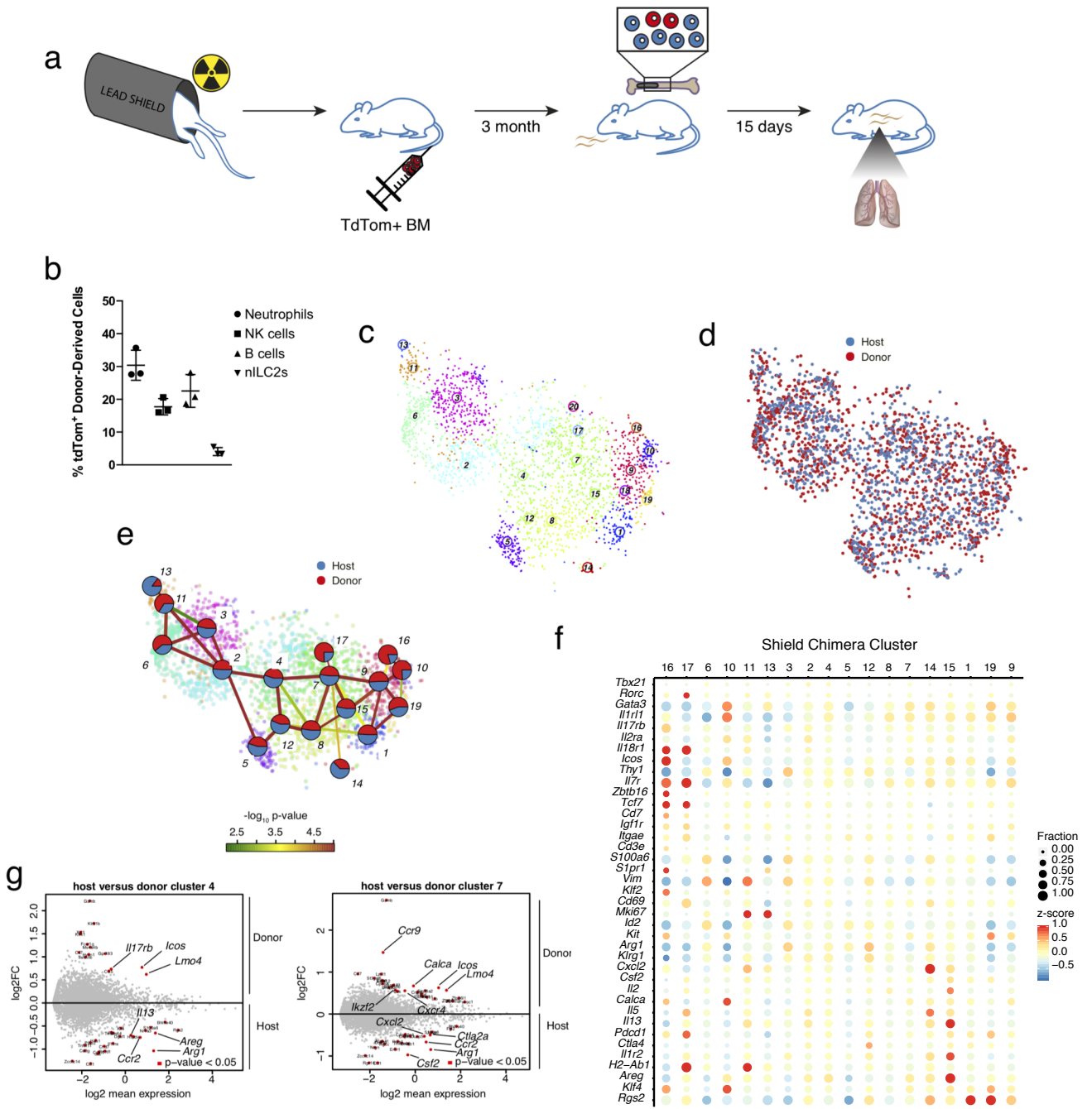
**Figure 5**



**Figure 6**

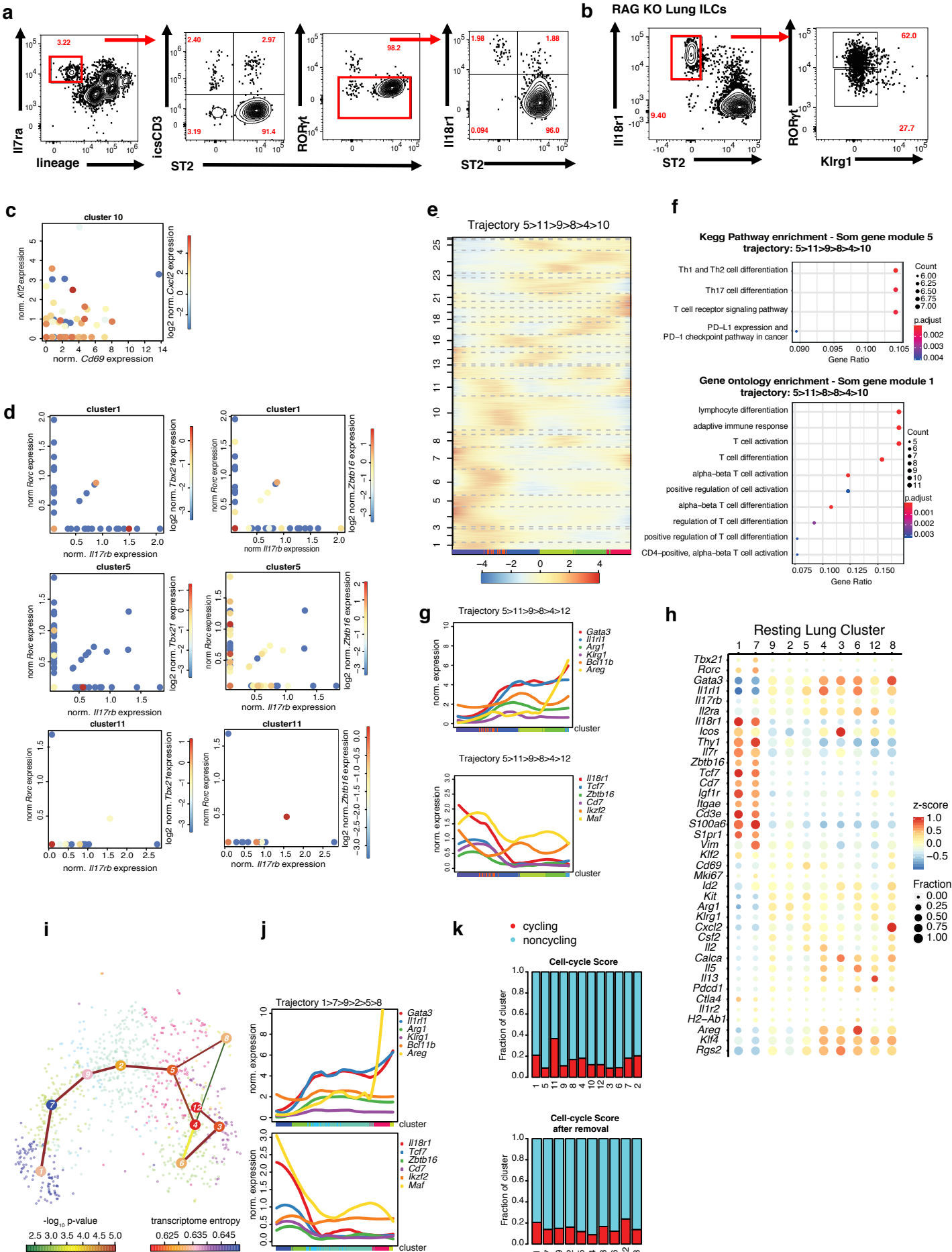


**Figure 7**

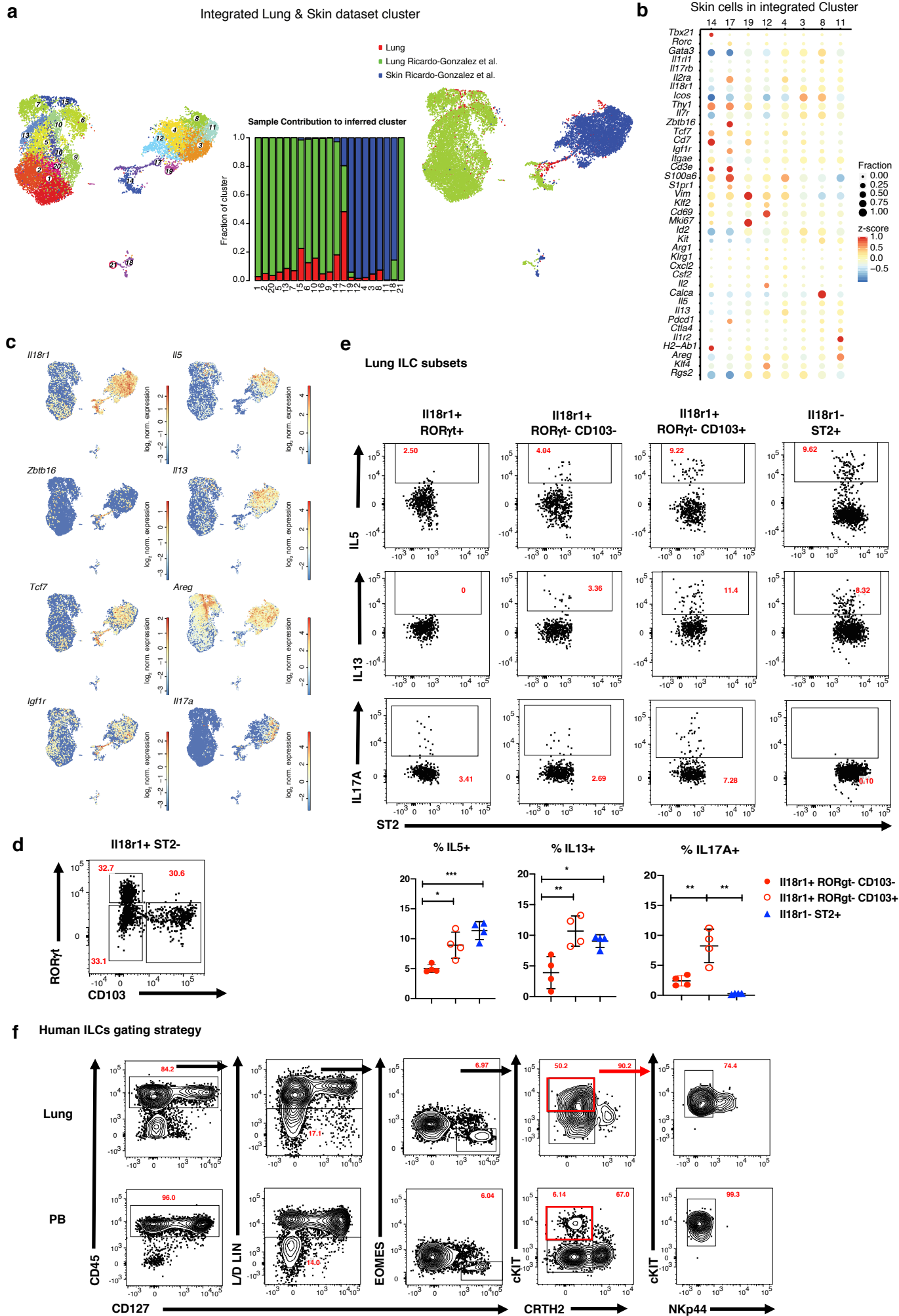




**Figure S1**

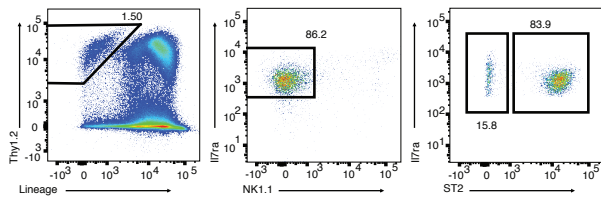


**Figure S2**

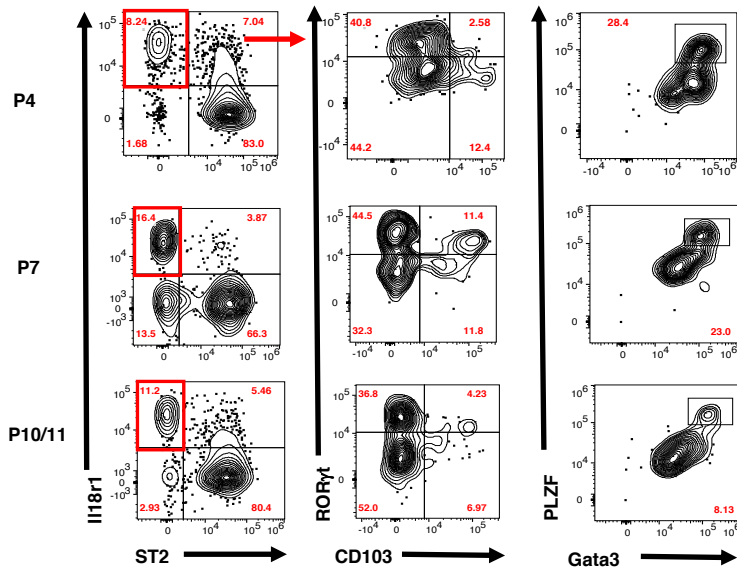


# Figure S3

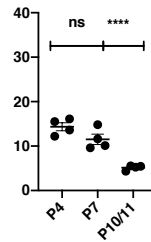
## a Neonatal lung sorting strategy



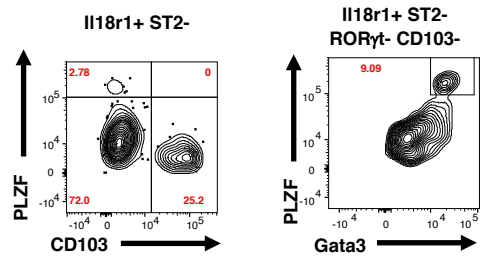
## b Neonatal lung Il7ra+ ILCs



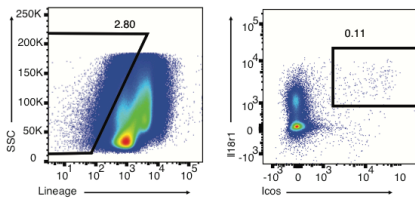
## c PLZF+ Gata3+ (% Il18r1+ ST2- ILCs)



## d Adult lung Il7ra+ ILCs



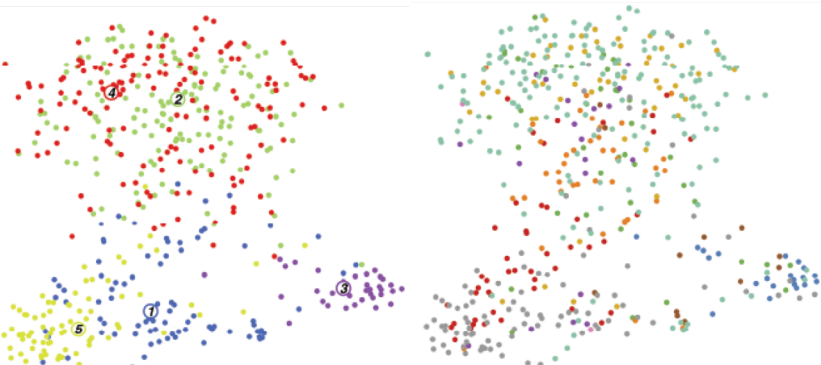
## e BM sorting strategy



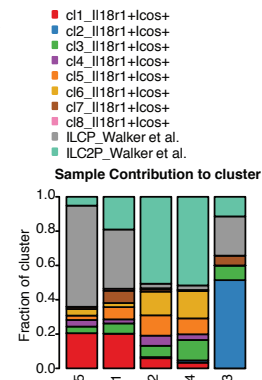
## f

Walker et al., Integrated with Il18r1+ BM Cluster

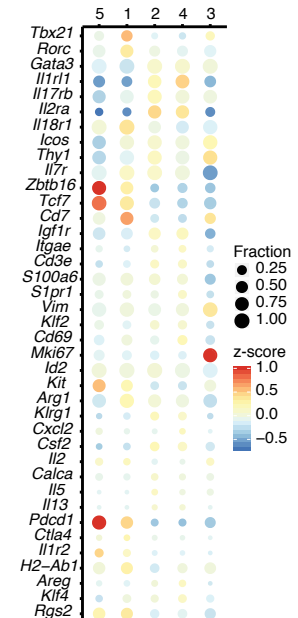
Walker et al., Integrated with Il18r1+ BM Cluster



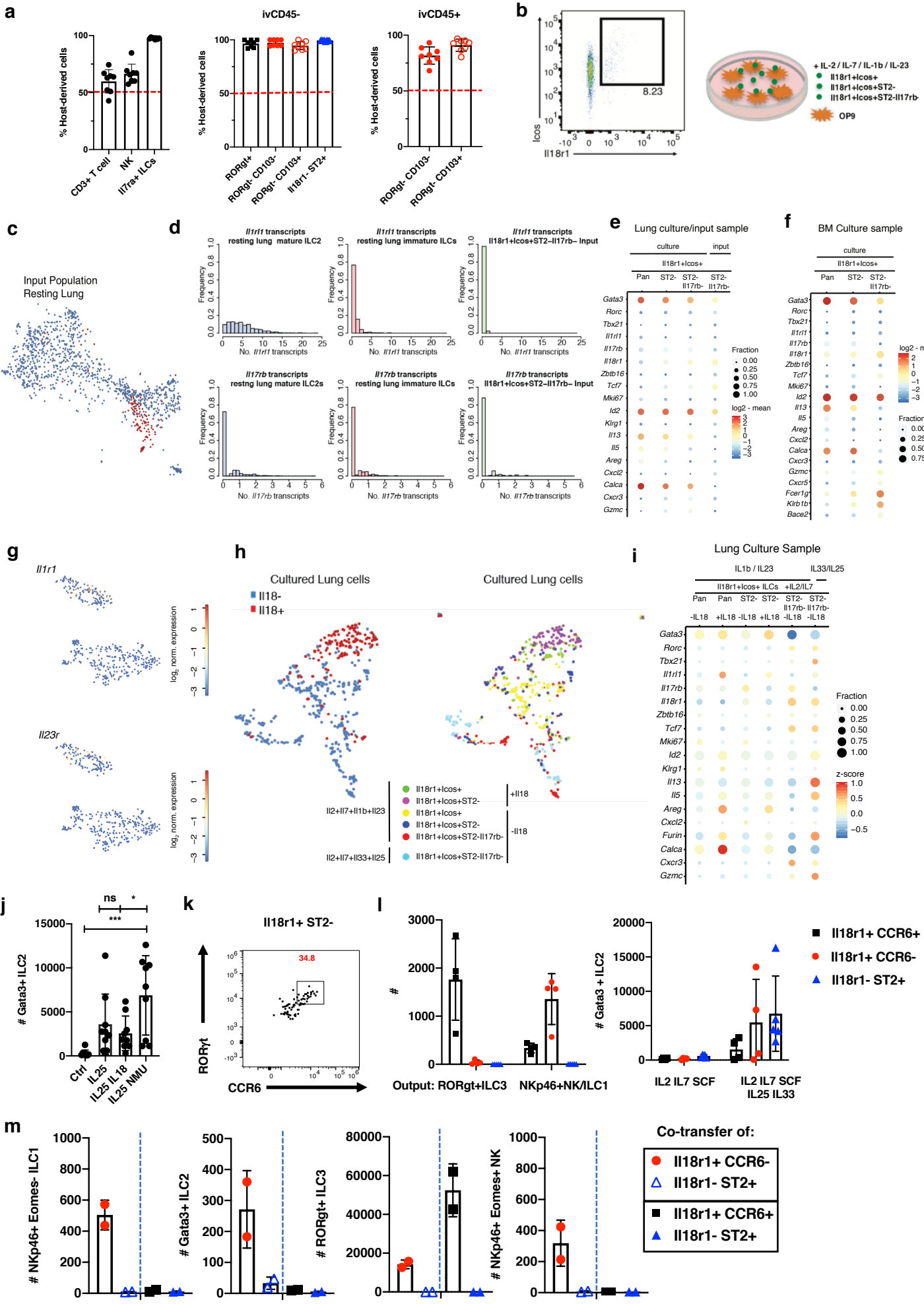
## g



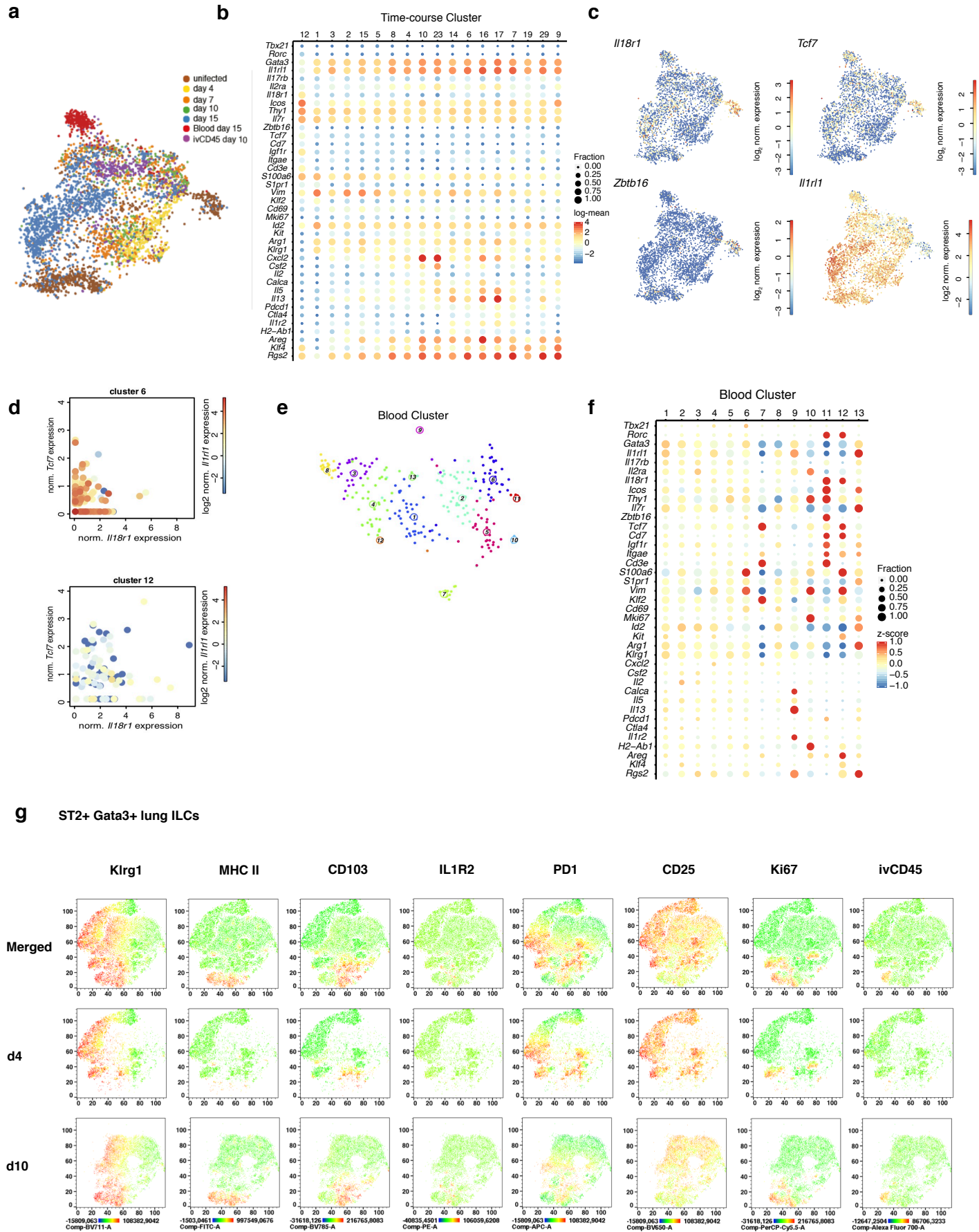
## h



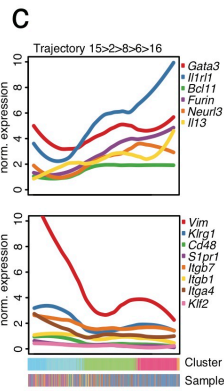
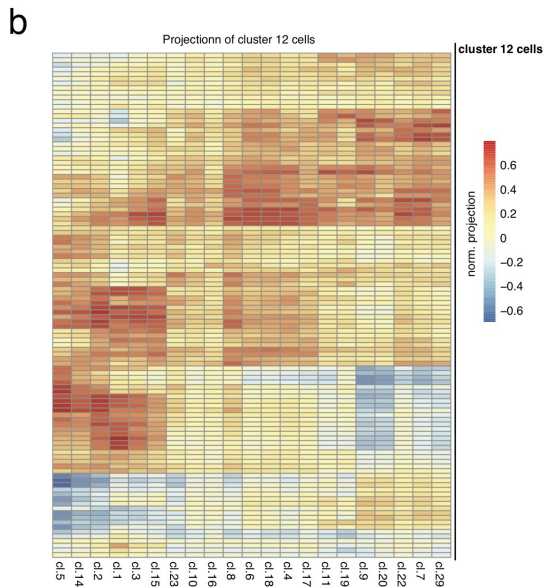
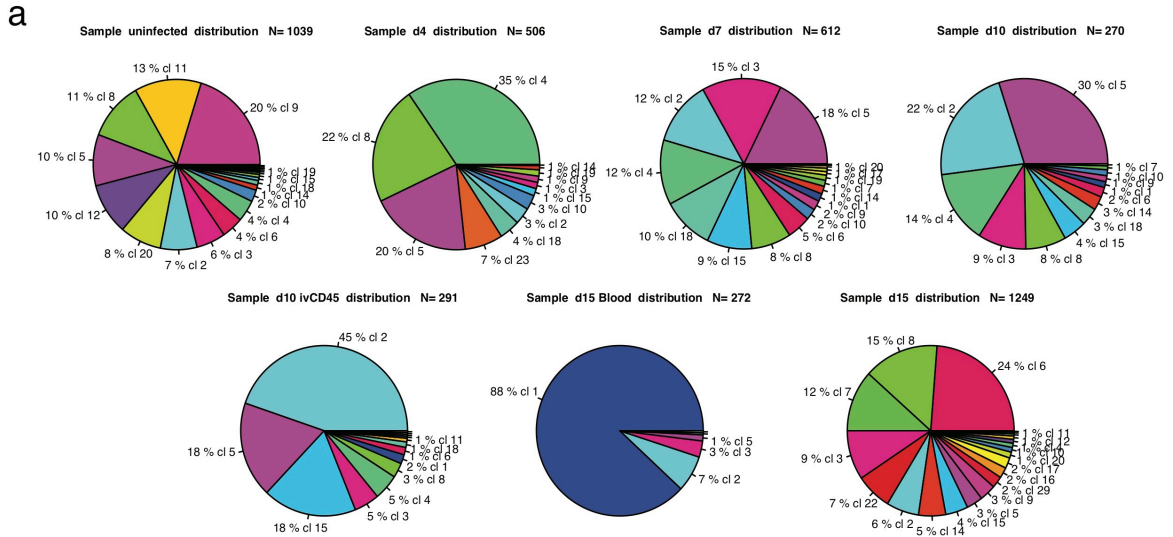
**Figure S4**



**Figure S5**



**Figure S6**



**Figure S7**

