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Pulmonary monocytes interact with effector T cells in the lung tissue to drive T_{RM} differentiation following viral infection

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Lung resident memory CD8 T cells (T_{RM}) are critical for protection against respiratory viruses, but the cellular interactions required for their development are poorly understood. Herein we describe the necessity of classical monocytes for the establishment of lung T_{RM} following influenza infection. We find that, during the initial appearance of lung T_{RM}, monocytes and dendritic cells are the most numerous influenza antigen-bearing APCs in the lung. Surprisingly, depletion of DCs after initial T cell priming did not impact lung T_{RM} development or maintenance. In contrast, a monocyte deficient pulmonary environment in CCR2^{-/-} mice results in significantly less lung T_{RM} following influenza infection, despite no defect in the antiviral effector response or in the peripheral memory pool. Imaging shows direct interaction of antigen-specific T cells with antigen-bearing monocytes in the lung, and pulmonary classical monocytes from the lungs of influenza infected mice are sufficient to drive differentiation of T cells in vitro. These data describe a novel role for pulmonary monocytes in mediating lung T_{RM} development through direct interaction with T cells in the lung.

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INTRODUCTION

During respiratory virus infections, effector CD8 T cells are primed in the lung-draining lymph nodes by antigen presenting cells (APCs) that have migrated from the infected lung.^{1–3} Following a program of proliferation and differentiation that is regulated by antigen encounter, co-stimulation receptor engagement, and the local cytokine environment, virus-specific effector CD8 T cells then traffic back to the infected lung to mediate their effector functions.^{4,5} Upon entry into the lung tissue, virus-specific effector CD8 T cells may re-encounter antigen presented by local APCs, including monocytes, macrophages, dendritic cells, and infected epithelial cells. In addition to antigen re-encounter, cues from the local microenvironment can further influence the differentiation of virus-specific CD8 T cells into short-lived effector or long-lived memory cells, ultimately directing cell fate.^{6–8} Despite the significance of these cell fate decisions for pathogen clearance and the establishment of immune memory, the importance of the local microenvironment on T cell differentiation in the tissue and the roles of individual tissue-resident APC subsets that provide these signals for the development of T cell memory are not well understood.

Following respiratory virus clearance, subsets of the CD8 memory precursor cells in the lung will differentiate into tissue-resident memory T cells (T_{RM}). Lung T_{RM} have been shown to be critical for protective cellular immunity against secondary heterosubtypic respiratory infections, enabling the rapid detection of the invading pathogen and thereby limiting pathogen replication and immunopathology.^{9–11} The programming of CD8

T_{RM} has been extensively investigated in recent years, and key cytokines such as TGF- β and IL-15 have been shown to be important for their development.^{12–14} Although antigen stimulation is required to initiate the effector T cell response in the lymph nodes, its role in T_{RM} development in the tissue had been less well characterized. Recently, it has been demonstrated that differentiation of lung T_{RM} during influenza virus infection requires virus-specific T cells to re-encounter antigen in the lung tissue.^{15,16} However, it is not clear if this requires interaction between effector T cells and specific APC subsets in the lung. Previous reports have shown that targeting vaccines to pulmonary CD103⁺ dendritic cells or alveolar macrophages promotes the establishment of lung T_{RM}, but it is unclear whether this was due to the ability of these APC subsets to promote T_{RM} programming during initial priming in the lymph node, or whether these APC subsets regulate lung T_{RM} establishment through antigen re-encounter in the lung itself.^{17,18} Given the importance of T_{RM} for protective cellular immunity in the lung, it is critical to define the cellular and molecular requirements for their establishment and identify new approaches for optimizing vaccines against respiratory pathogens.

To better define the factors that promote T_{RM} in the lung, we investigated the role of different lung-resident APC subsets in virus-specific CD8 T_{RM} development. Although essential for initial virus-specific CD8 T cell activation, depletion of CD11c⁺ dendritic cells during the peak of the effector T cell response did not impact the number of virus-specific lung CD8 T_{RM} following influenza infection. Surprisingly, analysis of pulmonary APC

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subsets around the time of viral clearance showed that monocytes were among the most numerous lung APC subsets harboring influenza antigens, and this correlated with the initial appearance of virus-specific lung CD8 T_{RM}. While monocytes have been investigated for their pro-inflammatory roles in innate immunity, their ability to influence T cell responses through direct interactions with virus-specific CD8 T cells has not been as extensively investigated.^{19–21} Using CCR2-deficient mice, which are defective in monocyte trafficking to the lung during influenza infection,²² we observed a significant decrease in the number of virus-specific lung CD8 T_{RM} in both the parenchyma and airways, but there was no effect on the number of circulating virus-specific memory CD8 T cells in the spleen. Notably, there were no differences in the number of virus-specific effector CD8 T cells or virus-specific memory CD8 precursor cells generated in the lung at the time of viral clearance when comparing wild-type and CCR2^{-/-} mice, demonstrating the role for monocytes in T_{RM} development was restricted to antigen re-encounter in the tissue and not initial T cell priming. In support of this, imaging of the lung revealed a close interaction between virus-specific CD8 T cells and monocytes. Furthermore, pulmonary monocytes sorted from infected lungs were sufficient to activate naive antigen specific T cells, as well as induce their expression of CD103 on a subset of highly-divided cells *in vitro*. Together, these data define a novel role for lung tissue-resident monocytes as critical mediators in the establishment of lung CD8 T_{RM} but not circulating T cell memory following respiratory infection, through presentation of viral antigens to T cells in the infected lung.

RESULTS

Lung CD8 T_{RM} develop immediately following viral clearance
While there is growing appreciation for the role of tissue resident memory T cells in protection against viral challenges at mucosal surfaces, less is known about their ontogeny. To determine the cellular interactions critical for lung T_{RM} development, we first sought to define the kinetics of the appearance of CD8 T_{RM} in the lung following influenza infection. Flu-specific CD8 T cells were identified in both the lung vasculature and lung parenchyma following intravital labeling. The appearance of cells with a tissue-resident phenotype (CD69⁺ CD103⁺) was observed in the lung parenchyma beginning on day 8 post infection (Figs. 1a, b), just preceding the peak of the effector CD8 T cell response at day 10 post infection, as well as influenza viral clearance.^{23,24} Over the course of the effector T cell response in the lung, the frequency of vascular versus tissue-resident flu-specific CD8 T cells remains relatively constant. However, the frequency of CD69⁺ CD103⁺ flu-specific CD8 T cells steadily increases within the resident population from day 8–14 post infection (Fig. 1c). We see a similar pattern in the number of lung T_{RM}, with a peak at D10 and a decline following viral clearance that mimics the kinetics of the total effector response (Fig. 1c). Similar to polyclonal flu-specific T cells, transgenic OT-I T cells resident in the lung show an increase in the frequency of CD69⁺ CD103⁺ cells from days 8–14 following infection with influenza x31-OVA (Fig. 1d), and numbers of lung OT-I T_{RM} matched the kinetics of the overall effector T cell response. Coincident with the expansion of CD69⁺ CD103⁺ cells we observed continued antigen stimulation, as measured by Nur77-GFP expression, in flu-specific CD8 T cells resident in the lung, but not in the lung vasculature or the spleen, on day 10 post infection (Fig. 1e, f). Increased Nur77-GFP expression in lung flu-specific CD8 T cells continued through day 14 post infection, but was largely absent by day 30 post infection. In addition, there were increased numbers of flu-specific CD69⁺ CD103⁺ cells in the lung at day 14 post infection (Fig. 1g). Together, these data show the seeding of the lung-resident T cell pool occurs rapidly following the resolution of influenza infection and is associated with continued antigen recognition by flu-specific T cells in the lung tissue.

Depletion of lung dendritic cells after initial T cell priming does not alter lung CD8 T_{RM} establishment

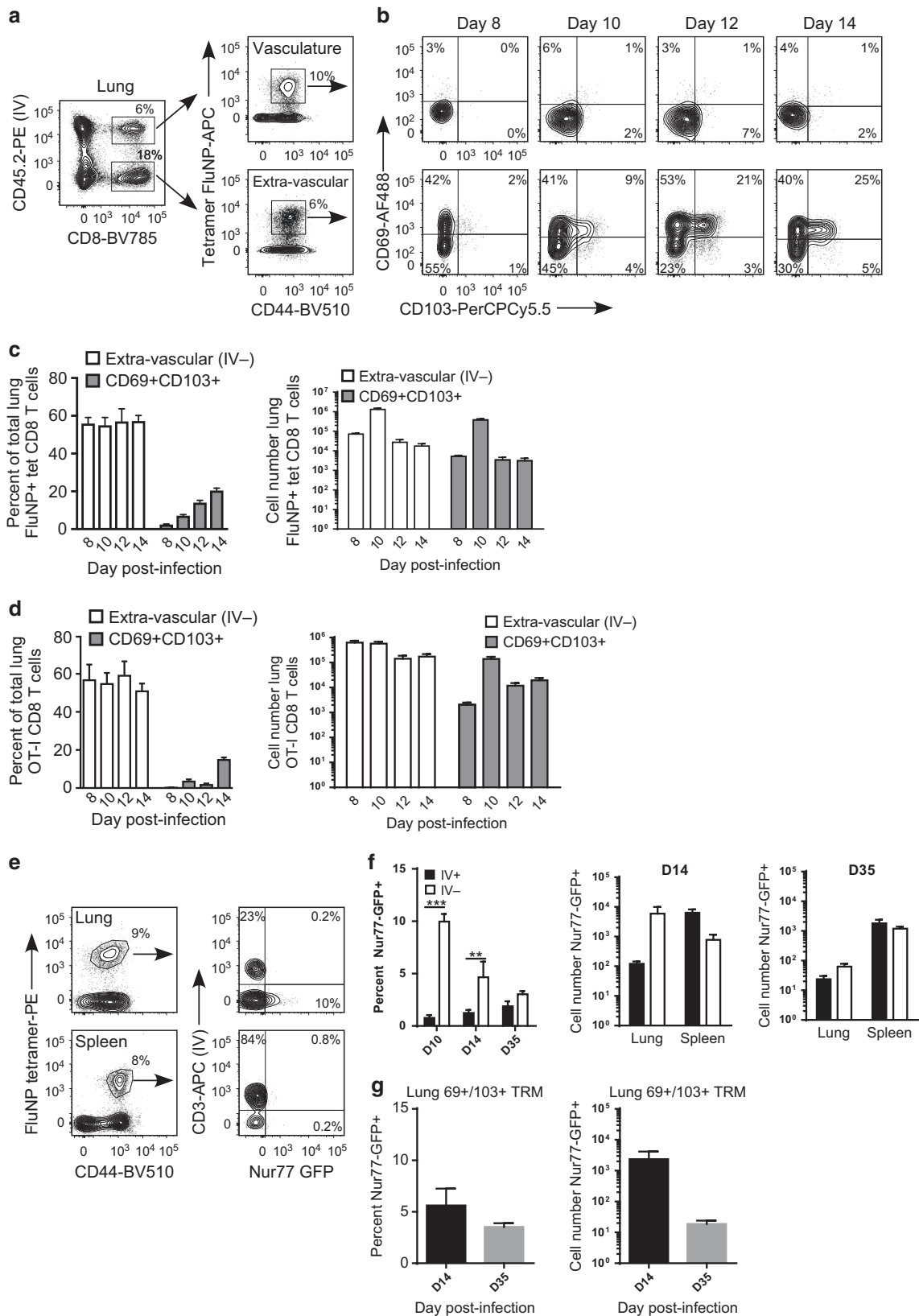
Previous studies have shown that the establishment of lung-resident T cell memory requires antigen re-encounter in the lung,¹⁵ and flu-specific T cells continue to receive antigen stimulation in the lung following viral clearance.^{25,26} Dendritic cells, specifically, are appreciated as efficient mediators of T cell activation and differentiation. As well, the importance of dendritic cells in the initiation of a T cell response against influenza infection has been well documented,^{4,22,27–29} but the role pulmonary DCs may be playing in the differentiation of flu-specific T_{RM} in the tissue after initial priming is less well understood. To address this question, we investigated the establishment of lung T_{RM} following depletion of DCs after initial T cell priming (Fig. 2a). Treatment of CD11c-DTR (Itgax-DTR) chimeras with diphtheria toxin (DTx) beginning on day 5 post infection was sufficient to deplete the majority of CD11c⁺ cells in the lung, which include both DCs and alveolar macrophages, while leaving the CD11b⁺ cells intact (Fig. 2b, c, and S1). Surprisingly, depletion of CD11c⁺ cells before T_{RM} generation but after initial T cell activation had no effect on the phenotype or numbers of flu-specific lung T_{RM} in the airways or parenchyma on day 14 post infection (Fig. S1D and S1E), or at memory (Fig. 2d, e). Therefore, despite their importance for the initial priming of naive CD8 T cells, interactions between DCs and virus-specific CD8 T cells in the tissue are not required for the establishment of lung T_{RM}.

Lung-resident dendritic cells and monocytes are among the most numerous cell types presenting influenza antigens during the initial appearance of lung T_{RM}

This finding led us to broaden our approach to identify additional APC subsets potentially involved in lung T_{RM} differentiation. We investigated which APC subsets in the lung were displaying antigen to flu-specific T cells, particularly at the conclusion of the effector T cell response, when resident memory is established. CD8 T cells in the lung make direct contact with many different APC subsets, including dendritic cells, macrophages, and monocytes (Fig. 3a). To determine which APC subsets were capable of presenting influenza antigens in the lung, we performed intravital labeling to identify different lung extra-vascular APC subsets, (Fig. S2) and assessed intracellular influenza nucleoprotein (FluNP) content.³⁰ This staining approach was specific for the influenza nucleoprotein, as isotype staining, as well as staining of lung APCs following Sendai virus infection, failed to give a positive signal (Fig. 3b). Intracellular FluNP staining was detected in pulmonary monocytes, monocytic respiratory dendritic cells (MoRDCs), CD11b^{hi} DCs, and CD103⁺ DCs, and this staining was restricted to the lung-resident (CD45.2⁻) population of each subset (Fig. 3c). The number of cells containing FluNP in each subset was highest at day 6 post infection, and substantial numbers of pulmonary APCs containing FluNP were detected through day 15 post infection (Fig. 3d).³⁰ Although, as expected, substantial numbers of FluNP⁺ DCs were present in the lung, it was surprising to observe that pulmonary monocytes were also among the most numerous FluNP⁺ APC subset from days 10–15 post infection. Thus, we began to assess what role these monocytes may play in the differentiation of lung T_{RM}, as well as any differences in the roles of the classical and non-classical monocyte subsets.

Inhibiting monocyte recruitment to the lung significantly decreases lung CD8 T_{RM} development

To investigate the role pulmonary monocytes may be playing in the development of lung T_{RM} *in vivo*, we used mice deficient in C-C chemokine receptor type 2, or CCR2, which lack the ability to efficiently traffic monocytes from the circulation into sites of mucosal inflammation. Previous studies characterizing influenza infection in CCR2^{-/-} mice observed no defect in the flu-specific effector CD8 T cell response or viral clearance,^{31,32} but the mice do



show decreased monocyte-driven immunopathology.²² To test this, we seeded WT and *CCR2*^{-/-} mice with naïve OT-I T cells, infected the mice with x31-OVA, and tracked the OVA-specific, as well as endogenous fluNP-specific T cell response (Fig. 4a). As

expected, we observed a significant decrease in the number of monocytes recruited to the lung in *CCR2*^{-/-} mice following influenza infection, but no difference in the numbers of other lung APC subsets, including MoRDC, CD103⁺, and CD11b^{hi} DC subsets

Fig. 1 Rapid appearance of lung resident CD8 T cells following influenza infection. **a** Gating strategy and representative flow plots for the intravital labeling of FluNP-specific CD8 T cells in the lung. **b** Representative flow plots for CD69 and CD103 staining on FluNP-specific CD8 T cells in the lung vasculature (top row) or lung extra-vascular compartments (bottom row). **c** Frequency and number of total extra-vascular FluNP-specific CD8 T cells and CD69⁺ CD103⁺ resident FluNP-specific CD8 T cells among total lung FluNP-specific CD8 T cells over time. **d** Frequency and number of total extra-vascular OT-I CD8 T cells and CD69⁺ CD103⁺ resident OT-I CD8 T cells among total lung OT-I CD8 T cells over time. **e** Representative staining of Nur77-GFP expression in FluNP-specific CD8 T cells. **f** Frequency and number of Nur77-GFP⁺ FluNP-specific CD8 T cells that are circulating (IV⁺) or extra-vascular (IV⁻) in the lung and spleen on days 10, 14, and 35 post infection. **g** Frequency and number of Nur77-GFP⁺ FluNP-specific tissue-resident CD8 T cells expressing both CD69 and CD103 in the lung extra-vascular population. ****p* < 0.001 (two-tailed Student's *t*-test) All graphs error bars are S.E.M. Data are representative of 3 independent experiments with 5 mice per time point (**a-c** and **g**) or 2 independent experiments with 4 mice each (**e** and **f**)

(Fig. S3). Similar to previous reports, at day 10 post infection there were no differences in the number of OT-I effector T cells in the BAL, lung interstitium (lung extra-vascular, LEV), or spleen between WT and CCR2^{-/-} mice (Fig. 4b, c).³¹ In addition, there was no difference in the number of CD69⁺ CD103⁺ lung-resident OT-I cells at this peak of the acute CD8 T cell response (Fig. 4c). To determine whether CCR2^{-/-} mice showed a defect in overall memory T cell development, we assessed the number of memory precursor cells (MPECs) in the lung and spleen (Fig. 4d). Similar to our observations of the overall effector T cell pool, there was no difference in the number of CD127^{hi} KLRG1^{lo} MPECs in the lung or spleen. Thus, CCR2^{-/-} mice showed no defect in the flu-specific effector CD8 T cell response, even within the lung tissue and airways (BAL).

In contrast to the effector T cell response, CCR2^{-/-} mice showed a significant decrease in the number of OT-I lung T_{RM} in both the airway (BAL) and interstitium (LEV) at memory (Fig. 4e, f). Importantly, there was no difference in the number of OT-I memory T cells in the spleen, indicating that the role of pulmonary monocytes in the generation of CD8 T cell memory was restricted to the lung-resident pool. In addition to a significant decrease in the total number of lung extra-vascular T cells, there was a significant decrease in the number of CD69⁺ CD103⁺ T_{RM} OT-I cells in both the airways and interstitium in CCR2^{-/-} mice (Fig. 4f). Although there is some overlap between the number of lung T_{RM} observed in WT and CCR2^{-/-} mice, the compiled data show a significant defect in the average number of flu-specific lung T_{RM} in CCR2-deficient mice.

To differentiate between the potential role classical and non-classical monocytes may be playing in the CCR2^{-/-} model, we generated CX3CR1-DTR bone marrow chimeras to allow for the selective depletion of CX3CR1⁺ non-classical monocytes (Fig. S4A).³² This model showed no difference at memory between the PBS and DTx treated mice, with similar numbers of memory OT-I being generated in the lung extra-vascular compartment, the airways, and the spleen (Fig. S4B and S4C). This indicated that the classical monocytes, but not non-classical monocytes, had a prominent role in driving the differentiation of lung T_{RM} following influenza infection.

To confirm the results of the transferred OT-I cells in the CCR2^{-/-} model, we also investigated the endogenous polyclonal flu-specific T cell response. Similar to the OT-I response, there was no difference in the number of effector fluNP-specific CD8 T cells in the airways and lung on day 10 post infection, but a significant decrease in the number of fluNP-specific lung extra-vascular T cells in the airways and interstitium in CCR2^{-/-} mice at memory (Fig. 4g). Furthermore, in CCR2^{-/-} mice, the overall decrease in fluNP-specific lung extra-vascular T cells was paralleled by a significant decrease in the number of CD69⁺ CD103⁺ T_{RM} in the airway and interstitium (Fig. 4h).

Assessing the impact of reduced numbers of lung T_{RM} on cellular immune protection in CCR2^{-/-} mice is complicated due to the reduced monocyte recruitment, which results in decreased immunopathology and weight loss.^{22,33} To overcome this issue, we treated (H3N2) influenza x31-immune WT and CCR2^{-/-} mice with FTY-720 to sequester circulating immune cells in secondary

lymphoid organs and limit the response to the virus to lung T_{RM} and challenged the mice with heterologous (H1N1) influenza PR8 to assess the protective efficacy of the lung T_{RM} population. Similar to previous reports, mock FTY-720-treated CCR2^{-/-} mice showed decreased weight loss compared to WT mice following PR8 challenge (Fig. 4i). In contrast, treatment with FTY-720 resulted in significantly greater weight loss in CCR2^{-/-} mice compared to WT mice, indicating that the decreased numbers of lung T_{RM} in CCR2^{-/-} mice are associated with impaired immune protection.

Pulmonary monocytes interact with virus-specific CD8 T cells in vivo and are sufficient to drive CD8 T cell activation and differentiation in vitro

The effect of defective monocyte trafficking on lung T_{RM} establishment suggested two possibilities: that the inflammatory milieu of the lung was altered in a manner detrimental to T_{RM} differentiation, or that T_{RM} development is driven by direct interactions between antigen-specific T cells and monocytes in the lung tissue. Given that antigen re-encounter in the tissue is necessary to establish lung T_{RM} following an influenza infection,¹⁵ we focused on the potential role of pulmonary monocytes presenting antigens directly to virus-specific CD8 T cells. The lung is a large organ, making the precise localization of individual cells crucial if they are to directly engage one another. As monocytes are among the most abundant APC subsets containing influenza antigens when lung T_{RM} first appear, we examined whether different fluorescently-marked monocyte subsets were interacting with virus-specific CD8 T cells in the lung (Fig. 5a). Influenza antigen-bearing monocytes were observed in close contact with virus-specific CD8 T cells in the lung on day 12 post infection (Fig. 5b, red arrows). To ensure that the monocytes not only contained but were presenting antigen on MHC-I as described by Jakubzick et al.,³⁴ we stained lung monocytes for H-2K^b bound to SIINFEKL, finding that both classical and non-classical monocytes were capable of presenting virus-derived peptide on MHC-I (Fig. 5c). Furthermore, we found that a subset of these H-2K^b-SIINFEKL positive cells also contained intracellular FluNP protein, thus demonstrating both flu antigen uptake, as well as presentation on MHC-I (Fig. 5d). Though the cells were in contact, it was still possible they did not contribute to T cell activation and differentiation. To test this, we sorted classical and non-classical monocytes from the lungs of mice infected with x31, pulsed the monocytes with SIINFEKL peptide, and cultured the monocytes with naive OT-I T cells. Both monocyte subsets were able to induce substantial proliferation of OT-I cells as measured by CTV dilution (Fig. 5d). However, in highly divided cells, only classical monocytes were able to generate a population of OT-I T cells that co-expressed the T_{RM} markers CD69 and CD103 (Fig. 5e, f). Surprisingly, dendritic cell subsets, while able to induce CD103 upregulation, did so to a lesser degree than classical monocytes (Fig. 5h). Both subsets of monocytes were able to induce the expression of CD127 on highly divided cells, indicating that they were capable of generating OT-I cells with the potential to become memory T cells (Fig. S5). Together, these data indicate that lung monocytes, particularly classical monocytes, are present

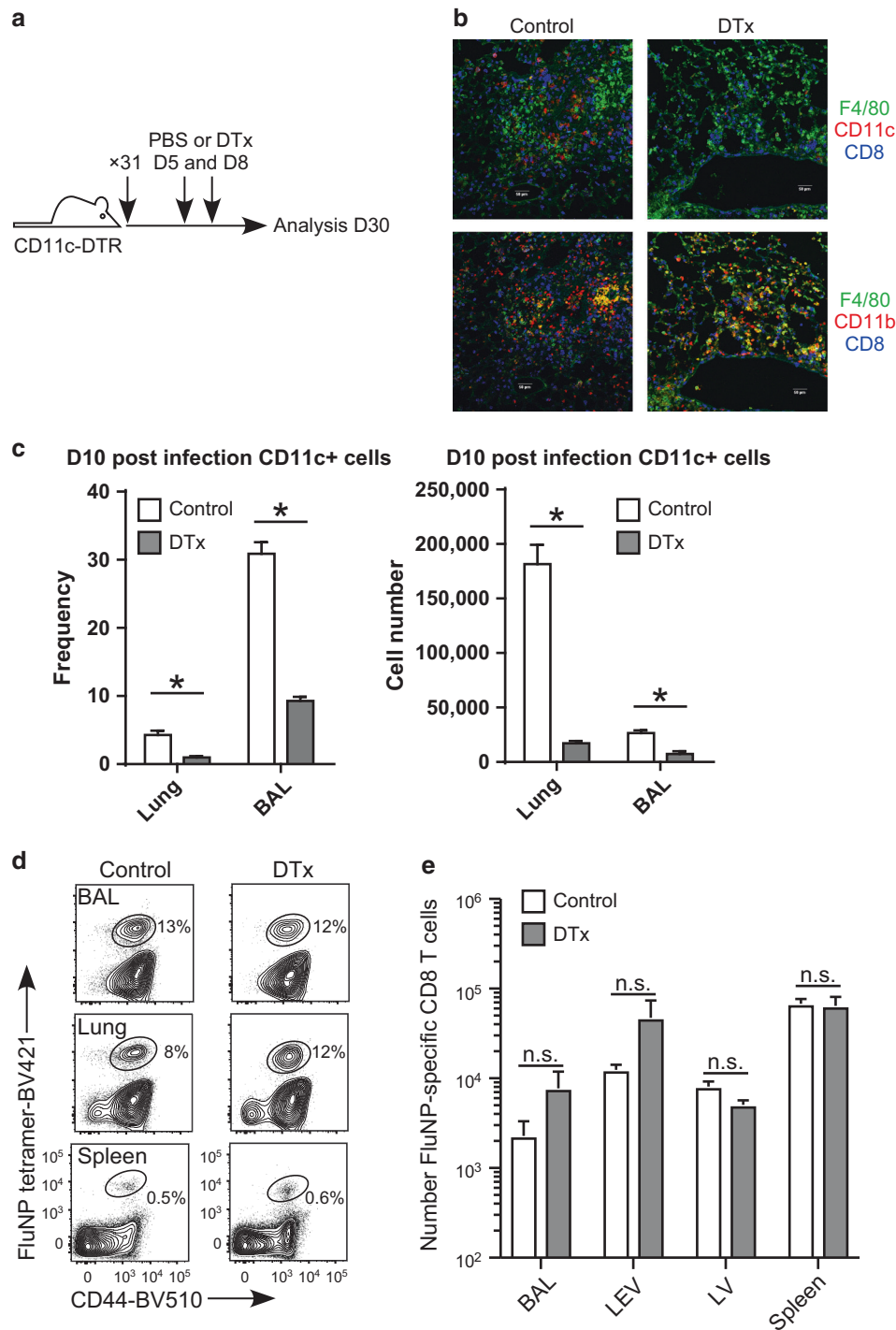


Fig. 2 Depletion of DCs after initial priming does not affect establishment of lung T_{RM} . **a** Experimental diagram of DC depletion using CD11c-DTR (Ilgax-DTR) chimeras. **b** Representative images from the lung on day 11 post infection (day 3 post depletion) demonstrating the specific depletion of CD11c⁺ cells. **c** Frequency and number of total CD11c expressing cells in the lung and BAL at D10 post infection after DTx or PBS treatment. * $p < 0.05$ (two-tailed Student's *t*-test). **d** Representative staining of FluNP-specific CD8 T cells 30 days post infection in control or DTx-treated mice. BAL and lung plots are gated on extra-vascular (IV⁻) cells. **e** Number of FluNP-specific CD8 T cells in BAL, lung extra-vascular (LEV), lung vasculature (LV), and spleen 30 days post infection in control or DTx-treated mice. Data are representative of three independent experiments with 3–4 mice per group. All graphs error bars are S.E.M

in the lung micro-environment, co-localize with CD8 T cells, and are sufficient to drive T cell differentiation, including the expression of CD103 on a subset of highly-divided cells. Overall, these data demonstrate a novel role for antigen presentation by pulmonary monocytes in the establishment of virus-specific lung T_{RM} , but not systemic T cell memory, following influenza infection.

DISCUSSION

Many studies have shown the importance of dendritic cells for the initiation of antiviral T cell responses following influenza infection, with particular subsets such as CD8 α^+ and CD103⁺ DCs playing specific roles in naïve T cell activation and differentiation.^{4,35–37} Given the requirement for antigen re-encounter in the tissue for

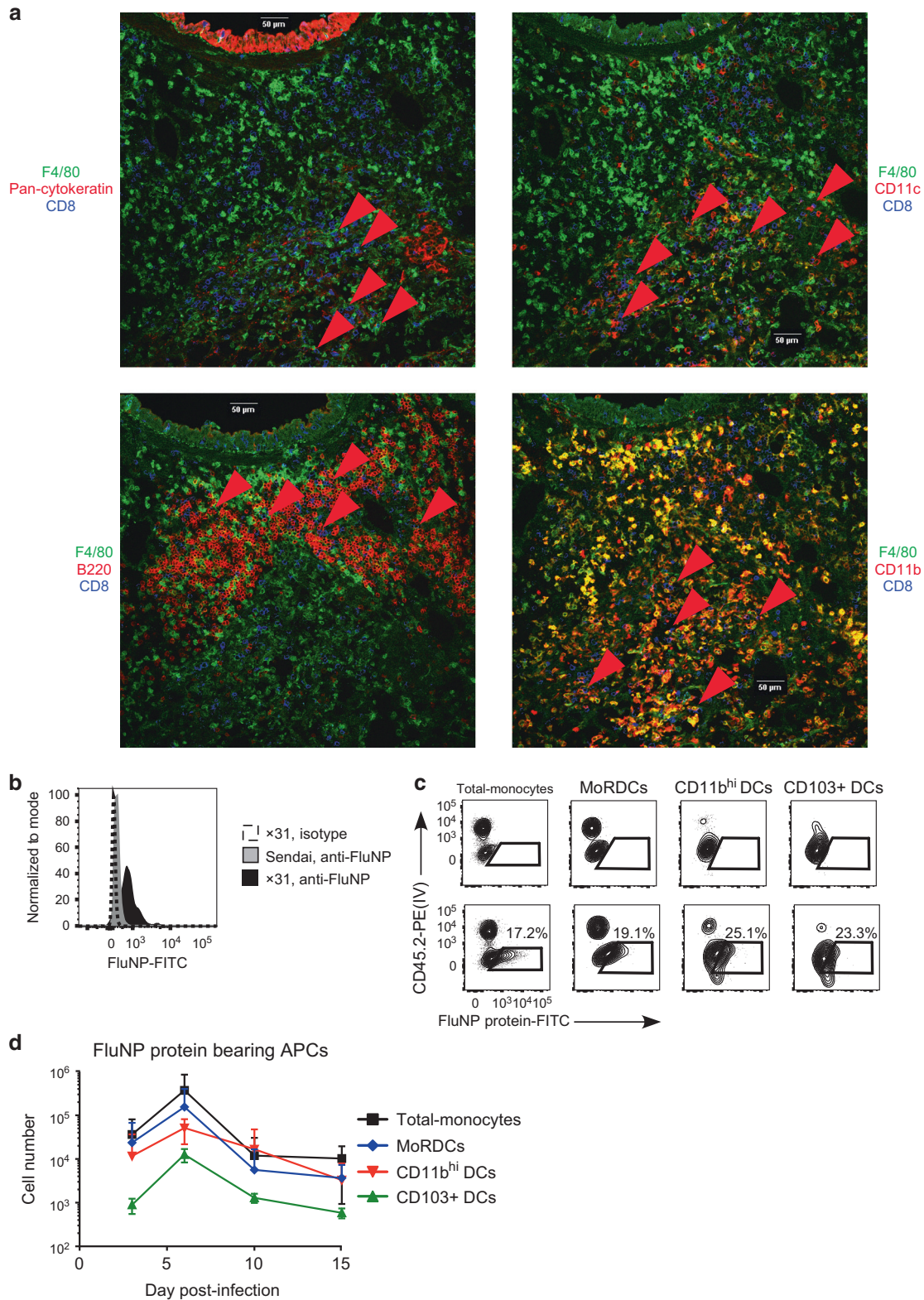


Fig. 3 Large numbers of pulmonary DCs and monocytes contain influenza antigens following viral clearance and during the initial appearance of lung T_{RM}. **a** Representative confocal microscopy images showing the localization of CD8 T cells and various APC subsets in the lung of influenza infected mice. Red arrows highlight areas of T cell and APC co-localization. **b** Representative staining showing the specificity of intracellular FluNP staining in pulmonary monocytes following infection with influenza x31 (black histogram) or Sendai virus (gray histogram). Isotype control staining is shown by the dashed line. **c** Representative staining of intracellular FluNP in extra-vascular (CD45.2 IV⁻) lung APC subsets on day 7 post infection. **d** The number of FluNP-containing total monocytes (black), MoRDCs (blue), CD11b^{hi} DCs (red), and CD103⁺ DCs (green) in the lung over the course of influenza infection. Data are representative of 3 independent experiments with 5 mice per time point. All graphs error bars are S.E.M.

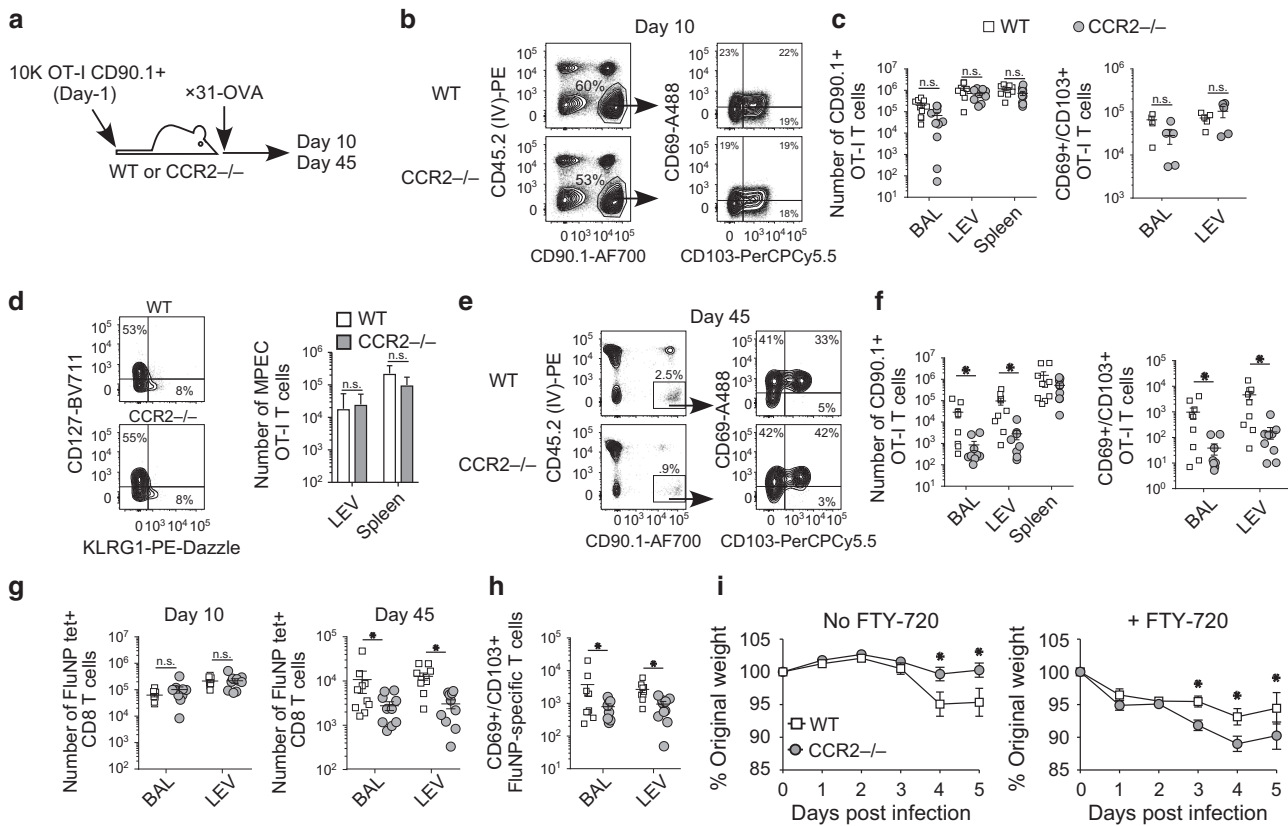


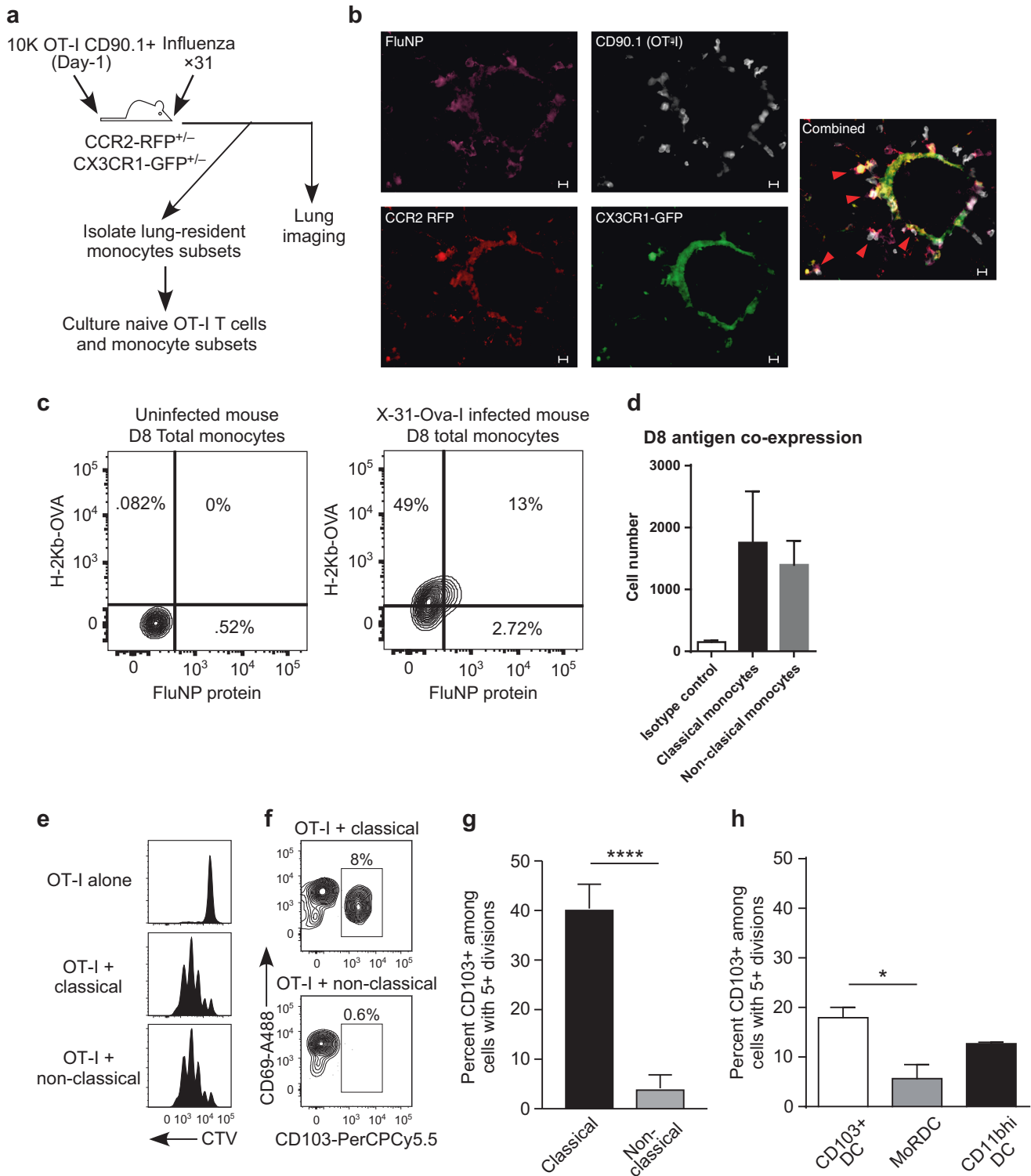
Fig. 4 Inhibiting monocyte recruitment to the lung significantly reduces the number virus-specific lung extra-vascular and lung T_{RM} following influenza infection. **a** Experimental design for investigating the role of pulmonary monocytes in lung T_{RM} establishment. **b** Representative staining and **(c)** numbers of total and $CD69^+$ $CD103^+$ OT-I $CD8^+$ T cells in the airways (BAL), lung extra-vascular (LEV), and spleen on day 10 post infection in WT and $CCR2^{-/-}$ mice. **d** Representative staining and numbers of $CD127^+$ $KLRG1^-$ MPEC OT-I T cells in the lung and spleen on day 10 post infection. **e** Representative staining and **(f)** numbers of total and $CD69^+$ $CD103^+$ OT-I $CD8^+$ T cells in the airways (BAL), lung extra-vascular (LEV), and spleen on day 45 post infection in WT and $CCR2^{-/-}$ mice. **g** Number of FluNP-specific $CD8^+$ T cells in the airways (BAL) and extra-vascular in the lung (LEV) on days 10 and 45 post infection in WT and $CCR2^{-/-}$ mice. **h** Number of $CD69^+$ $CD103^+$ FluNP-specific $CD8^+$ T cells resident in the airways (BAL) and lung (LEV) on days 10 and 45 post infection in WT and $CCR2^{-/-}$ mice. **i** Weight loss of WT and $CCR2^{-/-}$ influenza x31-OVA-immune mice challenged with PR8-OVA in the presence (right graph) or absence (left graph) of FTY-720. Data represent 3 independent experiments with 5 mice per group (**b–h**), or 3 independent experiments with 6 mice per group (**i**). All graphs error bars are S.E.M. * $p < 0.05$ (two-tailed Student's *t*-test)

establishing lung T_{RM} , it was surprising that depletion of $CD11c^+$ cells after initial T cell activation showed that DCs were dispensable for lung T_{RM} formation. In contrast, inhibiting monocyte recruitment to the lung had a dramatic impact on the establishment of lung T_{RM} , despite having no effect on the magnitude of the effector T cell response. Thus, the ability of monocytes to promote T cell responses against influenza is not through the initial priming and expansion of antiviral T cells, but through their ability to present viral antigens to effector T cells in the infected lung tissue and drive T cell differentiation.

Classical monocytes have been characterized as innate inflammatory mediators that produce large amounts of IL-1, IL-6, and TNF α , and promote tissue damage,³⁸ but their ability to drive adaptive immune responses through antigen presentation has been understudied.³⁹ Monocytes have been shown to promote T_H1 responses during viral infection through direct priming of naive T cells in the lymph node.^{4,40,41} However, as we observed no defect in the flu-specific effector $CD8^+$ T cell response or the systemic flu-specific memory $CD8^+$ T cell pool in $CCR2^{-/-}$ mice, it is unlikely that the decreased numbers of lung T_{RM} in these mice are due to a defect in the initial priming of the flu-specific T cell response. Rather, our data show that antigen presentation by pulmonary monocytes to effector $CD8^+$ T cells in the lung tissue is important for lung T_{RM} establishment. Consistent with these findings, $Ly6C^+$ inflammatory monocytes

are efficient at cross-presentation to $CD8^+$ T cells in the presence of TLR agonists, especially TLR7.^{34,42–45} Together, these findings support a model where monocytes in the infected lung are activated by viral TLR agonists, promoting efferocytosis of dying cells and enhancing cross-presentation of influenza antigens to flu-specific $CD8^+$ T cells, which ultimately drives the establishment of lung T_{RM} .

Several studies have identified roles for specific APC subsets in the establishment or maintenance of T_{RM} . Cross-presentation by $DNGR-1^+$ dendritic cells was shown to be required for optimal generation of T_{RM} , but not circulating memory T cells, in a model of cutaneous Vaccinia virus infection.⁴⁶ In the lung, targeting vaccines to respiratory dendritic cells or alveolar macrophages was shown to induce local T_{RM} that could protect mice against respiratory challenge.^{17,47} Recently, it was reported that inflammatory monocytes were important for the maintenance of both lung T_{RM} and circulating memory T cell subsets following Vaccinia virus infection, but the mechanism by which monocytes were promoting memory maintenance was unknown.³¹ We have extended these findings to demonstrate a critical role for antigen presentation by monocytes in establishing lung T_{RM} . Although we did not observe a defect in the circulating memory T cell pool in our study, this discrepancy is likely due to the differences in tissue tropism between influenza and Vaccinia viruses. As influenza replication and inflammation is localized to the respiratory tract,



the impact of monocyte antigen-presentation on memory T cell development during a respiratory virus infection would be limited to the lung.

Although our data support a direct role for antigen presentation by pulmonary monocytes in driving lung T_{RM} establishment, there may be additional monocyte-derived factors contributing to this process, such as cytokines or the propagation of tissue repair. For example, monocytes can produce IL-15, which has been implicated in the initial lodgment of T_{RM}.⁴⁸ Monocytes are also

prevalent in areas of tissue repair following viral clearance in the lung, and these sites have been identified as anatomical niches, termed repair-associated memory depots, that promote the maintenance of lung T_{RM}.⁴⁹ Future studies investigating antigen-independent functions of monocytes in T_{RM} differentiation and maintenance, and the interplay between monocytes and other APCs in these processes, will be required to fully understand the contributions of monocytes for the development of resident T cell memory.

Fig. 5 Pulmonary monocytes interact with T cells in the lung during infection, present influenza-derived antigen, and are sufficient to drive the activation and differentiation of T_{RM} -like CD8 T cells in vitro. **a** Experimental design for using CX3CR1^{+/GFP} CCR2^{+/RFP} dual reporter mice infected with x31-OVA for microscopy and monocyte isolation for in vitro culture. **b** Representative fluorescent microscopy images from the lung at day 12 post infection. OT-I T cells interacting with FluNP-containing monocytes are indicated with red arrows. Scale bar is 20 μ m. **c** Representative co-staining of uninfected and x31-OVA-I infected mice with surface H-2Kb-OVA and intracellular FluNP protein, showing specificity of staining (left plot), as well as monocytes with both surface H-2Kb-OVA and intracellular FluNP protein (right plot). **d** Number of lung extra-vascular monocytes co-expressing FluNP protein and H-2Kb-OVA staining day 8 post infection. **e** Cell trace violet dilution of OT-I T cells cultured in the presence or absence of classical or non-classical monocytes for three days. **f** Representative staining of CD69 and CD103 on OT-I CD8 T cells with 5+ cell divisions as indicated by CTV dilution. **g** Frequency of CD103⁺ OT-I CD8 T cells stimulated by classical or non-classical monocytes with more than 5 cell divisions. **** $p < 0.0001$ (two-tailed Student's *t*-test) **(h)** Frequency of CD103⁺ OT-I CD8 T cells stimulated by CD103+ DC, MoRDC, or CD11bhi DC subsets with more than 5 cell divisions. * $p < .05$ (two-tailed Student's *t*-test) Data are representative of 3 independent experiments, with each in vitro culture (**d–i**) run in triplicate. All graphs error bars are S.E.M.

Monocytes and effector CD8 T cells are recruited to discrete sites of inflammation in the lung through CCR2-dependent and CXCR3-dependent mechanisms, respectively.^{3,19} Given the large size of the lung, having a mechanism to recruit both APCs and virus-specific CD8 T cells to the same inflamed local microenvironment where virus is present is an efficient means to ensure T cells have access to antigen and accessory signals such as costimulation and cytokines that will promote their differentiation. The high number of antigen-bearing monocytes in the lung, their co-localization with antigen specific T cells, and their sufficiency to drive T cell differentiation in vitro all point to monocytes being critical mediators of T_{RM} differentiation.

It should be noted that monocytes did not induce robust expression of T_{RM} markers such as CD103 on all OT-I cells in our in vitro culture system, and we did not examine the transcriptional profile of these cells to assess the complete T_{RM} program.^{50,51} As the in vitro culture system cannot recapitulate all the complex interactions that guide CD8 T cell differentiation in vivo, we believe it is unlikely that monocytes alone are sufficient to program lung T_{RM} development following an influenza infection. It seems likely that multiple T cell—APC interactions, separated by time (initial T cell priming and antigen re-encounter) and location (lymph node and infected lung) are required. One potential developmental pathway suggested by our data is that dendritic cells may induce the initial T_{RM} program, and that pulmonary monocytes may provide additional signals that lead to further progression or enforcement of this program. We are currently investigating these possibilities.

While significant technical hurdles remain, if strategies can be devised by which vaccine-derived antigens are presented by monocytes to activated T cells in the lung, this may enhance vaccine efficacy through the establishment of greater numbers of lung T_{RM} . These findings also provide a rationale for antibody-targeted mucosal vaccines, such as used by Villadangos et al.,¹⁸ to directly target vaccine antigens to lung monocytes in order to produce more robust T_{RM} responses. Combined with the previously demonstrated efficacy of dendritic cell targeting vaccines, this could offer an avenue towards a combination vaccine capable of generating robust cellular immunity in the lung. In summary, we have identified a novel role for antigen presentation by pulmonary monocytes in driving the establishment of lung T_{RM} following influenza virus infection. Further exploration of the mechanisms by which monocytes promote T_{RM} differentiation may aid in the development of new strategies for vaccination against respiratory pathogens.

MATERIALS AND METHODS

Mice and infections

C57BL/6J (WT), B6.PL-Thy1^a/CyJ (CD90.1), B6.SJL-Ptprc^a Pepc^b/BoyJ (CD45.1), B6.129S4-Ccr2^{tm1lfc}/J (CCR2^{-/-}), B6.129P-CX3CR1^{tm1Litt}/J (CX3CR1-GFP), B6.129(Cg)-CCR2^{tm2.1lfc}/J (CCR2-RFP), C57BL/6-Tg(Nr4a1-EGFP/cre)820Khog/J (Nur77-GFP),

B6.129P2-Cx3cr1^{tm3(DTR)Litt}/J (CX3CR1-DTR), and B6.FVB-1700016L21Rik^{Tg(Itgax-DTR/EGFP)57Lan}/J (CD11c-DTR) mice from The Jackson Laboratory were housed under SPF conditions at Emory University and Kindai University. B6.129P-CX3CR1^{tm1Litt}/J and B6.129(Cg)-CCR2^{tm2.1lfc}/J mice were crossed to generate F1 dual reporter mice (CX3CR1^{+/GFP} CCR2^{+/RFP}) for imaging. Intranasal infection with influenza A/HKx31 (H3N2) at 30,000 egg infectious doses (EID₅₀), A/HKx31-OVA1 expressing SIINFEKL peptide at 30,000 EID₅₀, and influenza A/PR8-OVA1 expressing SIINFEKL peptide (H1N1) at 6000 EID₅₀ were performed as previously described.⁵² In some experiments, 10⁴ CD90.1⁺ naive OT-I CD8 T cells were injected i.v. into recipient mice one day prior to infection. In protection experiments, mice were injected daily i.p. with 150 μ g FTY720 (Cayman Chemical) suspended in PBS. All experiments were completed in accordance with the Institutional Animal Care and Use Committee guidelines of Emory University.

Generation of CD11c-DTR and CX3CR1-DTR chimeras

Recipient CD45.1 mice were injected i.p. with 600 μ g of busulfan (Otsuka Pharmaceutical). The next day and 5 \times 10⁶ BM cells isolated from CD11c-DTR (Itgax-DTR) or CX3CR1-DTR mice were injected intravenously. Chimeras were rested for 6 weeks for reconstitution, and were bled to confirm the presence of the donor CD45.2⁺ cells prior to virus infection. Some groups of mice were injected i.n. with 60 μ g of Diphtheria Toxin (DTx) (Sigma-Aldrich) or PBS following infection with influenza x31 or x31-OVA.

Tissue collection and flow cytometry

Intravital staining was performed immediately before mouse euthanasia and tissue harvest as previously described.⁵³ Briefly, to identify T cells resident in various tissues, including the lung parenchyma, 1.5 μ g of fluorophore-conjugated α -CD45.2 antibody in 200 λ 1 \times PBS was intravenously injected into the tail vein of mice; five minutes post injection, mice were euthanized with Avertin (2,2,2-Tribromoethanol, Sigma) and exsanguinated prior to harvest of BAL and other tissues. Cells in the lung airways were recovered by lavage with 5 \times 1 ml R10 media. Lung tissues were digested by collagenase D (Roche) for 30 min at 37 $^{\circ}$ C and enriched by centrifugation in 40/80% Percoll gradient. Splenocytes were obtained by straining through nylon mesh, followed by RBC lysis in buffered ammonium chloride. Cells were blocked first with mAbs to Fc γ III/II and then stained with APC-conjugated influenza NP₃₆₆₋₃₇₄/D^b tetramer. Tetramer-labeled cells were washed and stained with fluorophore-conjugated reagents purchased from BD Biosciences (CD103, CD11c, Ly6C, Siglec-F), BioLegend (CD103, CD11b, CD127, CD69, CD8a, CD90.1, KLRG-1, Ly6G, I-A/I-E, CD45.2), eBiosciences (CD4, CD44, H2KB-OVA), anti influenza NP (Abcam), and R&D (CCR2). Intracellular staining was performed using the Cytotfix/Cytoperm Kit according to manufacturer protocol (BD Biosciences). Tetramers were generated by the NIH Tetramer Core at Emory University. Samples were run on LSRFortessa and LSR-II flow cytometers (BD), and data were analyzed using FlowJo software (Tree Star). Cell sorting was performed on an SH800 (Sony) or FACSAria III cell sorter (BD Biosciences).



Fluorescence and confocal microscopy

Mouse lungs were inflated by intratracheal administration of optimal cutting temperature (OCT) media to preserve lung morphology followed by snap freezing in liquid nitrogen. Six or Seven-micrometer-thick cryosections were fixed for 2 min with acetone/ethanol, and blocked with combined rat serum, donkey serum, mouse serum and FcBlock (anti CD16/32 2.4G2) or Blocking One reagent (Nacalai Tesque) followed by blocking with endogenous avidin and biotin blocking system (Abcam). Sections were then stained with antibodies purchased from Biologend (CD90.1, anti-GFP, donkey anti rabbit IgG, CD8a, CD11b, CD11c, B220), F4/80 (Bay Bioscience), pan-Cytokeratin (Bioss Antibodies), Abcam (fluNP), Lifetech (streptavidin Alexa Fluor 405), Invitrogen (a-RFP rabbit polyclonal), and mounted with ProLong™ Diamond Antifade (Thermo Fisher). Images were acquired on an AxioObserver Z1 (Zeiss) using a ×100 oil objective at room temperature or a C2si confocal microscope (Nikon). Images were processed using Zen 2.3 blue edition software.

Monocyte and T cell in vitro co-culture

Classical and non-classical monocytes were sorted from lungs of day 10 × 31 influenza infected mice (CD45⁺, CD11b⁺, MHC-II⁻, Ly6g⁻, Ly6c⁺, CCR2⁺ classical monocytes; CD45⁺, CD11b⁺, MHC-II⁻, Ly6g⁻, Ly6c⁻, CCR2⁻ non-classical monocytes). Sorted monocytes were pulsed for 2 h with 1 μM OVA peptide (SIINFEKL) in round bottom plates at 37 °C. and cultured with naïve CD8 OT-I T cells isolated from spleens using the EasySep™ Mouse CD8⁺ T Cell Isolation Kit (Stem Cell Technologies) and stained with Cell Trace Violet (ThermoFisher) Co-cultures were performed at a 1:2 monocyte to T cell ratio for 3 days prior to analysis.⁵⁴

Statistical analysis

Statistical analysis was performed using Prism 5 (GraphPad Software), and significance was determined by an unpaired two-tailed Student's *t* test unless otherwise noted in the figure legend. *P*-values less than 0.05 were considered significant.

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AUTHOR CONTRIBUTIONS

P.R.D., S.T. and J.E.K. designed the experiments. P.R.D., E.K.C., A.N.W., T.T., Z.-R.T.L., N.K., I.E.U., S.L.H., S.U. and S.T. performed experiments and analyzed data. I.E.U. and S.T. edited the paper. P.R.D. and J.E.K. wrote the paper.

ADDITIONAL INFORMATION

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