

ARTICLE

Single-cell transcriptomic analysis of human colonic macrophages reveals niche-specific subsets

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Macrophages are a heterogeneous population of cells involved in tissue homeostasis, inflammation, and cancer. Although macrophages are densely distributed throughout the human intestine, our understanding of how gut macrophages maintain tissue homeostasis is limited. Here we show that colonic lamina propria macrophages (LpMs) and muscularis macrophages (MMs) consist of monocyte-like cells that differentiate into multiple transcriptionally distinct subsets. LpMs comprise subsets with proinflammatory properties and subsets with high antigen-presenting and phagocytic capacity. The latter are strategically positioned close to the surface epithelium. Most MMs differentiate along two trajectories: one that upregulates genes associated with immune activation and angiogenesis, and one that upregulates genes associated with neuronal homeostasis. Importantly, MMs are located adjacent to neurons and vessels. Cell-cell interaction and gene network analysis indicated that survival, migration, transcriptional reprogramming, and niche-specific localization of LpMs and MMs are controlled by an extensive interaction with tissue-resident cells and a few key transcription factors.

Introduction

In recent years, it has been shown that tissue macrophages, residing in virtually all organs, are extremely heterogeneous. Originally, macrophages were described as professional phagocytes that were fundamental to our immune defense. However, it is now clear that macrophages have a broad range of both immune and nonimmune functions. Moreover, many macrophage functions are tissue-specific, such as electrical conduction in the heart (Hulsmans et al., 2017), synaptic pruning in the brain (Hong et al., 2016), alveolar surfactant clearance in the lung (Guilliams et al., 2013), and limb regeneration in salamanders (Godwin et al., 2013). These functions are thought to be controlled by tissue-specific gene modules regulated by specific transcription factors (TFs) imprinted by the microenvironment. Interestingly, tissue-resident macrophages (TRMs) can be reprogrammed when transferred to a new microenvironment (Lavin et al., 2014), indicating that mature TRMs retain their plasticity. In addition to the local microenvironment, macrophage functions also depend on their ontogeny (embryonic precursors or adult monocytes), intrinsic factors, and time spent in the tissue (Bleriot et al., 2020).

The gut is an organ composed of several compartments with unique functions. The mucosa is covered by a one-cell-thick epithelial layer that separates our body from the outside world. This compartment stretches the local immune system to its limits: it has to rapidly and efficiently eliminate pathogens and toxins and at the same time tolerate harmless molecules (e.g., food proteins) and the gut microbiota. Below the mucosa is a thin layer of tissue termed submucosa, which overlies a thick layer of smooth muscle, termed muscularis propria, consisting of a circular and longitudinal muscular layer. The muscular layers are responsible for segmental contractions and peristaltic movement of the intestinal tract. The gut has an extensive enteric nervous system with a plexus of ganglia cells in both the muscular layers (Auerbach's plexus) and the submucosa (Meissner's plexus).

Macrophages populate all layers of the gut wall, and studies in mice suggest that they have niche-specific functions that are essential to maintain tissue homeostasis (De Schepper et al., 2018). Under steady-state conditions, lamina propria macrophages (LpMs) constantly phagocytose apoptotic epithelial cells and are critical for gut microbiota composition (Arandjelovic

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and Ravichandran, 2015; Earley et al., 2018). Recently, LpMs in distal colon were shown to maintain epithelial integrity by limiting fungal product adsorption (Chikina et al., 2020). Muscularis propria macrophages (MMs) interact with both neurons and vessels, and loss of MMs leads to loss of enteric neurons, vascular leakage, impaired secretion, and reduced intestinal motility (De Schepper et al., 2018; Muller et al., 2014). Moreover, Matheis et al. (2020) showed that MMs protected against post-infectious neuron damage. At birth, the mouse gut is populated with embryonic-derived macrophages. However, the LpMs are rapidly (within weeks) replaced by bone marrow-derived circulating monocytes (Bain et al., 2014). Monocytes also emigrate to muscularis propria, but in this compartment embryonic-derived macrophages appear to be more persistent (De Schepper et al., 2018). However, all studies cited above were performed in mice, and translation of these results to understand human macrophage biology should be made with caution.

Gut macrophages may also be detrimental to the host. Aberrantly activated macrophages have been shown to play a key role in inflammatory bowel disease pathology (Martin et al., 2019; Smillie et al., 2019), and tumor-associated macrophages are often associated with worse prognosis (Katzenelenbogen et al., 2020; Zhang et al., 2020). Because of their heterogeneity and plasticity, there is currently great interest in the search for strategies to reprogram macrophages as a therapeutic tool to treat both inflammatory disorders and cancer (Jahchan et al., 2019). However, to target macrophages in human diseases, a deeper understanding of their heterogeneity and tissue-specific functions is necessary.

Our current knowledge about macrophages in the human gut is limited. Using bulk RNA sequencing (RNA-seq), we have shown that LpMs in the small intestine consist of several transcriptional states (Bujko et al., 2018), and by following their turnover kinetics in transplanted duodenum, we found that host circulating monocytes rapidly entered the graft and differentiated into TRMs that completely replaced donor LpMs, within months after transplantation. Here we extend these findings by performing a detailed characterization of both LpMs and MMs from adult human colon, applying single-cell RNA-seq (scRNA-seq) and multicolor immunostaining in situ. Spatiotemporal analysis shows that both compartments contain multiple coexisting TRM subsets with subtissular-specific functions. We furthermore identify a limited number of TFs that may control TRM diversity, and we reveal an extensive cross talk between TRM subsets and tissue-resident cells, including stromal cells, epithelial cells, neurons, and immune cell lineages. These cell-cell interactions are most likely responsible for the imprinting of tissue-specific macrophage identity.

Results

Macrophages in colonic mucosa and muscularis propria comprise transcriptional states associated with monocytes and TRMs

Samples from the large intestine were obtained from colorectal cancer resections ($n = 4$). Macroscopically and microscopically normal tissue ≥ 10 cm from the tumor tissue was used. Mucosa

and muscularis propria were separated longitudinally, and pieces of tissue from each compartment were enzymatically digested to obtain single-cell preparations. Cells from mucosa and muscularis were analyzed separately (Fig. 1 A). Flow cytometric analysis revealed that $CD45^+CD3^-CD19^-HLA-DR^{int/+}$ cells separated into either $CD14^+$ macrophages or $CD14^-$ classic dendritic cells (cDCs), including $CD141^+$ cDC1 and $CD1c^+$ cDC2 (Fig. S1 A), as previously shown in the small intestine (Bujko et al., 2018). Thus, tissue-derived macrophages were sorted as $CD45^+HLA-DR^+CD14^+$ (Fig. S1 B) and processed on a 10X Genomics Chromium platform. A total of 63,970 cells with high-quality mRNA were analyzed. To present high-dimensional data in low dimension, we constructed Uniform Manifold Approximation and Projection (UMAP) plots of each compartment. We found that sorted $CD14^+$ cells expressed the macrophage markers *CD163* and *CD68*, further demonstrating that cells included in the analysis were all macrophages (Fig. S1 C).

We have previously shown that the macrophage population in the small intestine contains several transcriptionally distinct subpopulations (Bujko et al., 2018), including “transient” monocyte-like macrophages expressing high levels of calprotectin (heterodimer of *S100A8* and *S100A9*) and long-lived calprotectin-negative macrophages expressing a TRM phenotype (Bujko et al., 2018). To examine whether colonic LpMs and MMs contained similar phenotypes, we first assessed the expression of *S100A8* and *S100A9* and the complement component 1 chains (*CIQA*, *CIQB*, and *CIQC*); the latter were highly expressed on small intestinal TRMs (Bujko et al., 2018). UMAP visualization of both compartments showed that most LpMs and MMs expressed either *S100A8/S100A9* or *CIQA/CIQB/CIQC* genes (Fig. 1 B). Further transcriptomic analysis of *S100A8/S100A9*⁺ macrophages showed a high number of differentially expressed genes (DEGs; e.g., *FCN1*, *VCAN*, *AQP9*, and *TREMI*) that are preferentially expressed in blood monocytes and small intestinal monocyte-like macrophages (Fig. 1 C). *CIQA/CIQB/CIQC*⁺ macrophages, on the other hand, expressed many genes associated with TRMs (e.g., *MRC1*, *APOE*, *SELENOP*, *CSFIR*, *MERTK*, and *LYVE1*; Fig. 1 C). Flow cytometry of dispersed tissue macrophages confirmed that LpMs and MMs consisted mainly of calprotectin⁺*CIQ*⁺ and calprotectin[−]*CIQ*⁺ macrophages (Fig. 1 D). Together, these findings demonstrated that both LpMs and MMs contained monocyte-like macrophages and mature macrophages with a TRM phenotype.

Mucosal macrophages comprise transcriptionally distinct and niche-specific subsets

To characterize LpMs further, we performed clustering analysis using Seurat 3.0 (Stuart et al., 2019). Based on DEGs, 13 clusters were identified (Figs. 2 A and S2 A) that encompassed macrophages from all patients included in the study (Fig. S3 A). DEGs in clusters 0–4 (LpM₀–LpM₄) were reminiscent of small intestinal monocyte-like macrophages (Bujko et al., 2018), and enriched gene ontology (GO) terms were typical for innate immune responses such as responses to bacterium, fungus, and LPS (Fig. 3). LpM₀ expressed the highest levels of *S100A8*, *S100A9*, and *S100A12* (Fig. 2 B), indicating that this cluster constitutes the most recently elicited monocytes. LpM₂ expressed

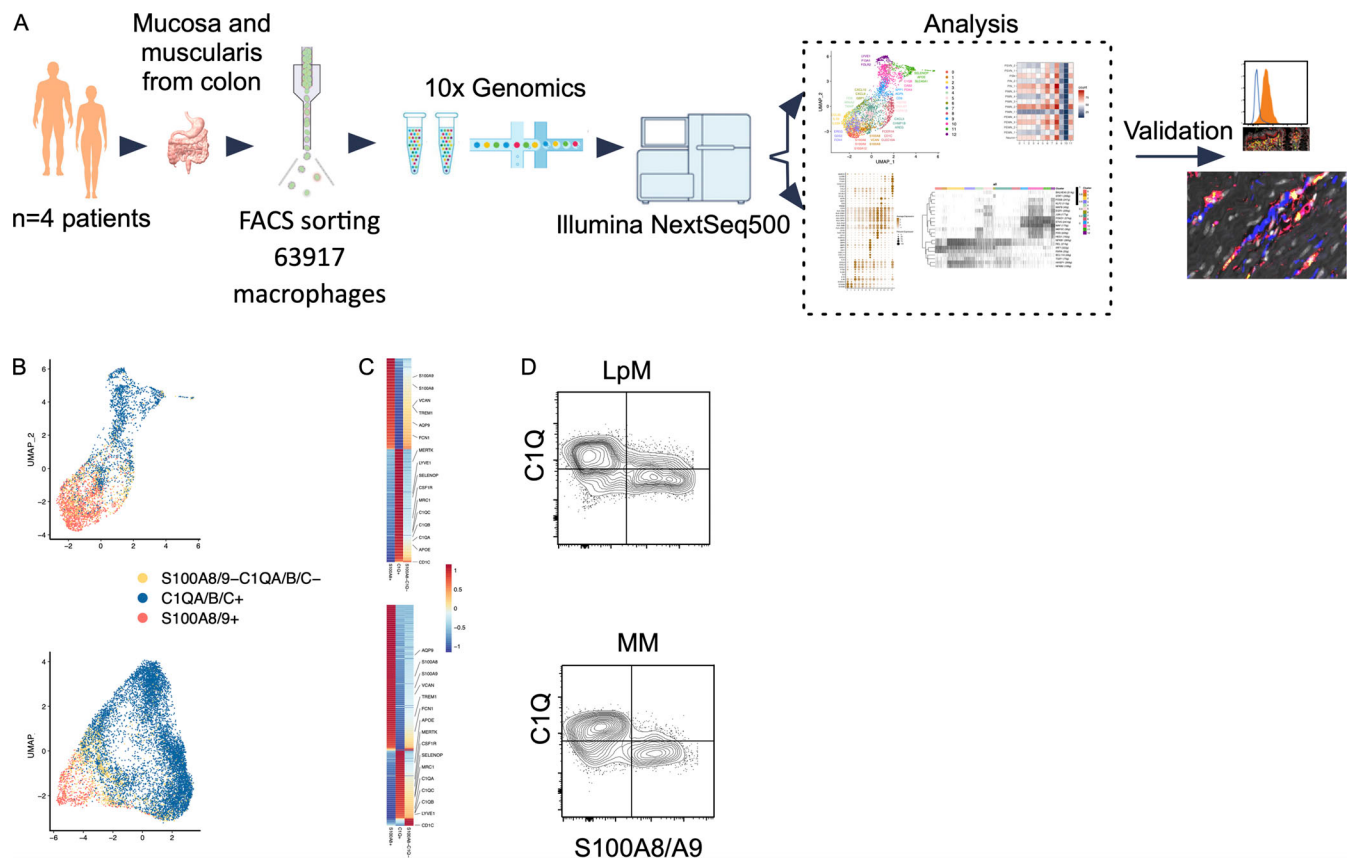


Figure 1. Schematic overview of scRNA-seq analysis and identification of monocyte-like and TRMs in colonic mucosa and muscularis propria. (A) Schematic overview of scRNA-seq analysis. (B) UMAP plots showing expression of *S100A8/A9*⁺ (red), *C1QA/B/C*⁺ (blue), and *S100A8/A9*⁻*C1QA/B/C*⁻ (yellow). The color among compartments is determined by its higher expression level. (C) Heatmap of DEGs by *S100A8/A9*⁺, *C1QA/B/C*⁺, and *S100A8/A9*⁻*C1QA/B/C*⁻ cells. Selected genes are indicated. Expression levels are visualized from low expression (blue) to high expression (red). (D) Flow cytometry plots showing intracellular staining for C1Q and S100A8/S100A9 (calprotectin) in CD45⁺HLA-DR⁺CD14⁺ cells from mucosa (top) and muscularis propria (bottom). Gates were set based on staining with irrelevant isotype-matched antibodies. Representative of more than four experiments.

high levels of many proinflammatory genes (e.g., *IL-1B*, *IL-1A*, *IL-6*, *IL23A*, *CXCL2*, *CXCL3*, and *CXCL8*) reminiscent of inflammatory macrophages found in inflammatory bowel disease lesions (Martin et al., 2019; Fig. 2, B and C). The immunoregulatory cytokine *IL10* showed highest expression in LpM₂–LpM₄ (Fig. 2 B). LpM₅ contained DEGs encoding many heat shock proteins (HSPs; Fig. S2 A), and top GO terms were related to unfolded protein responses (Fig. 3). Interestingly, LpM₆ expressed many IFN γ -inducible genes virtually absent in the other clusters; (e.g., *CXCL9*, *CXCL10*, *CXCL11*, *IDO1*, *GBP1*, *GBP2*, *GBP4*, and *GBP5*; Fig. 2, B and C). The top GO terms were antigen processing and presentation of exogenous peptide antigen via MHC class I, and type I IFN-, IFN γ - and TNF-signaling pathway (Fig. 3). This phenotype has recently been reported to promote antitumor immunity in colorectal cancer of mice (Qu et al., 2020). Cluster 8 expressed low levels of both *S100A8/S100A9* and *C1Q* genes (Fig. 1 B), but many DEGs were associated with DCs (e.g., *FCER1A*, *CD1C*, *CLEC10A*, and *CD1E*; Fig. 2 B). This cluster also expressed high levels of MHC class II genes (Fig. 2, B and C), and the top GO term was antigen processing and presentation of exogenous peptide antigen via MHC class II (Fig. 3). Flow cytometric analysis confirmed the presence of CD14⁺FCER1⁺CD1C⁺ cells in dispersed

mucosa (Fig. S1 D). This phenotype is reminiscent of a recently identified DC population, termed DC3, which originates independently of both cDCs and monocytes (Cytlik et al., 2020; Bourdely et al., 2020; Dutertre et al., 2019).

LpM₉ and LpM₁₀ expressed high levels of HLA class II genes (Fig. 2, B and C). Accordingly, enriched GO terms were antigen processing and presentation of exogenous peptide antigen via MHC class II (Fig. 3). GO terms for LpM₁₀ also included wound healing and receptor-mediated endocytosis (Fig. 3). Interestingly, LpM₉ (and to a lesser extent LpM₁₀) expressed *CD9*, *TREM2*, *SPPI*, and *ACP5* (Fig. 2, B and C), genes that were recently shown to identify monocyte-derived macrophages enriched in liver fibrosis (Ramachandran et al., 2019) and in adipose tissues of obese patients (Jaitin et al., 2019). LpM₁₂ selectively expressed many typical TRM genes, such as *LYVE1*, *COLEC12*, *F13A1*, and *FOLR2* (Fig. 2, B and C), and several chemokine genes (e.g., *CXCL2*, *CXCL3*, *CXCL8*, *CCL3*, and *CCL4*; Fig. 2, B and C). GO terms included chemokine-mediated signaling pathway, synapse pruning, apoptotic cell clearance, and receptor-mediated endocytosis (Fig. 3). To validate the results, we integrated our data with a recently published dataset of macrophages derived from human intestinal mucosa (Elmentaite et al., 2021). Clustering

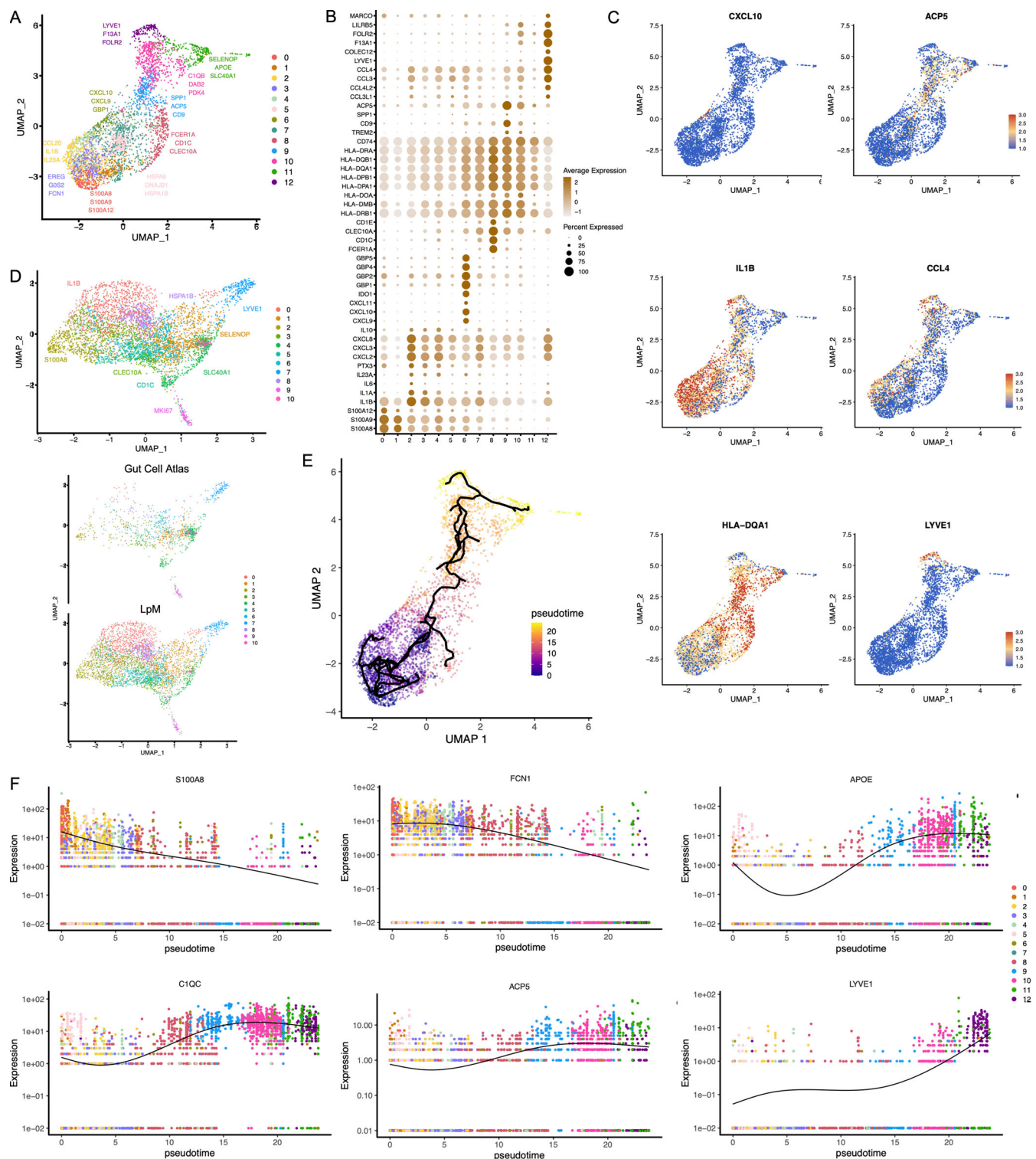


Figure 2. Cluster analysis, differential gene expression identification, and pseudotime trajectories for LpMs. (A) UMAP representation of 13 clusters (different colors) with annotated highly expressed gene identity within selected clusters. (B) Dot plot of average expression of selected genes. Expression levels are visualized from low (white) to high (brown) expression. The percentage of positive cells is indicated by circle size. (C) UMAP visualization of selected genes. Expression levels are visualized from low (blue) to high (red) expression. (D) UMAP of integrated analysis of LpMs and mucosal macrophages from the Gut Cell Atlas (Elmentaite et al., 2021). (E) Reconstructed developmental trajectory of LpMs showing pseudotime colored from blue to yellow. (F) Expression for some selected gene expression through pseudotime. Cell colors represent clusters.



Figure 3. GO terms significantly ($P < 0.05$) enriched at the transcriptomic level for selected clusters of LpMs and MM.

analysis showed that the macrophage transcriptomes from the two datasets were largely overlapping (Fig. 2 D). Moreover, interestingly, the integrated analysis identified a small cluster of DEGs associated with proliferation (e.g., *MKI67*), suggesting that some of the macrophages had the capacity to proliferate.

Other reports (Bain et al., 2014; Bujko et al., 2018) have suggested that the vast majority of LpMs originate from bone marrow-derived monocytes, which constantly emigrate to the mucosa and differentiate into TRMs. To further understand the process of monocyte-to-macrophage differentiation, we reconstructed their developmental trajectories using the Monocle 3 algorithm (Trapnell et al., 2014). We excluded DC3 from the analysis because this subset is reported to originate independently of the monocyte lineage (Cytlak et al., 2020; Bourdely et al., 2020; Dutertre et al., 2019). Selecting random cells in LpM₀ as root, we identified at least two distinct branches. A short branch ended in cluster 2, compatible with a distinct proinflammatory trajectory, whereas a major branch followed from LpM₀ to LpM₉₋₁₂ as a function of pseudotime (Fig. 2 E). As expected, genes typical for monocytes were downregulated following this trajectory, whereas typical TRM genes were increased (Fig. 2 F). Together, the findings suggested that LpMs consisted of multiple transcriptional cell states, indicating that incoming monocytes constantly differentiate into multiple distinct TRM subsets with time (Bujko et al., 2018).

Studies in mice have shown that TRMs occupy distinct tissue niches, where they display niche-specific functions (Chakarova et al., 2019; Guillemins et al., 2020). To determine the anatomic localization of LpM subsets, we performed multicolor immunofluorescence stainings in situ. LpMs were present in high numbers (median 1,080/mm², $n = 5$), and of those, calprotectin-expressing LpMs constituted a median of 6%, scattered in the lower part of the mucosa between crypts (Fig. 4 A), whereas a median of 45% LpMs strongly expressed CIQ, positioned in the subepithelial region (Fig. 4 B). To further determine the localization of distinct TRM clusters, we found that a median of 47% of LpMs stained for ACP5 (expressed by LpM₉ and LpM₁₀), whereas LYVE-1 and COLEC12, which were not expressed in the mucosa, decorated >90% of submucosal macrophages (SmMs; median 243 mm²; Figs. 4 D and S4 A), indicating that LpM₁₂ represented the SmM population. Several studies have shown that LYVE-1 is associated with perivascular macrophages, whereas macrophages expressing COLEC12, a scavenger receptor for uptake of myelin, has been associated with neurons (Bogie et al., 2017). Here we found that the vast majority of SmMs expressed both LYVE-1 and COLEC12.

Collectively, our analyses indicated that the colonic mucosa contains monocyte-derived LpMs that differentiate into multiple LpM subpopulations; some subsets had proinflammatory properties (e.g., LpM₂ and LpM₆), and other subsets had high antigen-presenting and phagocytic capacity and were

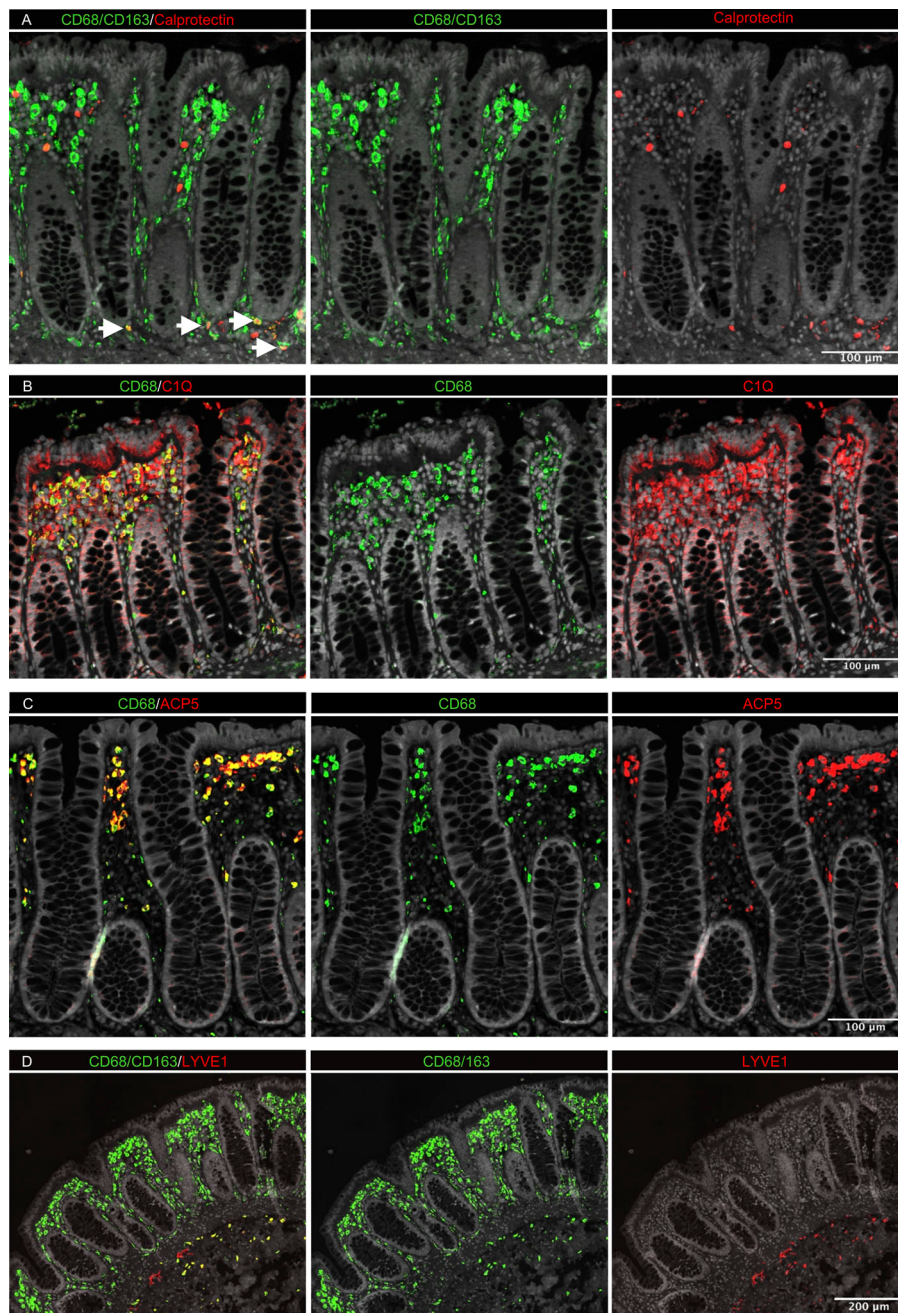


Figure 4. In situ localization of LpMs in human colon. (A–D) Sections stained for CD68/CD163 (green) and calprotectin (S100A8/S100A9; red; A); CD68 (green) and C1Q (red; B); CD68 (green) and ACP5 (red; C); and CD68/CD163 (green) and LYVE-1 (red; D). All sections were counterstained with Hoechst DNA-stain (gray). Arrows indicate CD68/CD163⁺ calprotectin⁺ LpMs (yellow; A). Calprotectin⁺ CD68/CD163[−] cells (red, left upper panel) are granulocytes. Representative of $n \geq 3$.

strategically positioned in the subepithelial region (LpM₉ and LpM₁₀). In contrast, LYVE1⁺ SmMs displayed a transcriptomic profile indicating low antigen-presenting capacity but high chemotactic and tissue-protective properties.

MMs comprise transcriptionally distinct subsets displaying different developmental trajectories

Next, we analyzed macrophages isolated from the muscular compartment. Following high-resolution clustering, the cells were separated into 12 transcriptionally distinct clusters (Figs. 5 A and S2 B) encompassing cells from all donors (Fig. S3 B). In three of the patients, we sampled muscularis propria from two different sites. Clustering analysis revealed that MMs from the sites were very similar (Fig. S2 C), indicating that MM

subpopulations were distributed homogeneously throughout the muscular compartment. Cluster 0 (MM₀) expressed high levels of *S100A8*, *S100A9*, *S100A12*, *IL1B*, *IL1A*, and *CXCL* chemokines (Fig. 5, B and C), and enriched GO terms were cellular response to bacterium and LPS as well as type I- and IFN γ -mediated signaling pathway, together with MHC class I-mediated antigen presentation (Fig. 3). MM₁, MM₂, and MM₄, which clustered adjacent to MM₀, expressed genes associated with immune activation (e.g., HLA class II genes; Fig. 5, B and C), and enriched GO terms were IFN γ -mediated signaling pathway and antigen processing and presentation of exogenous peptide antigen via MHC class II (Fig. 3). Cluster 3 was reminiscent of DC3 (Cytlak et al., 2020; Bourdely et al., 2020; Dutertre et al., 2019), demonstrating that this DC subset also resided in muscularis propria

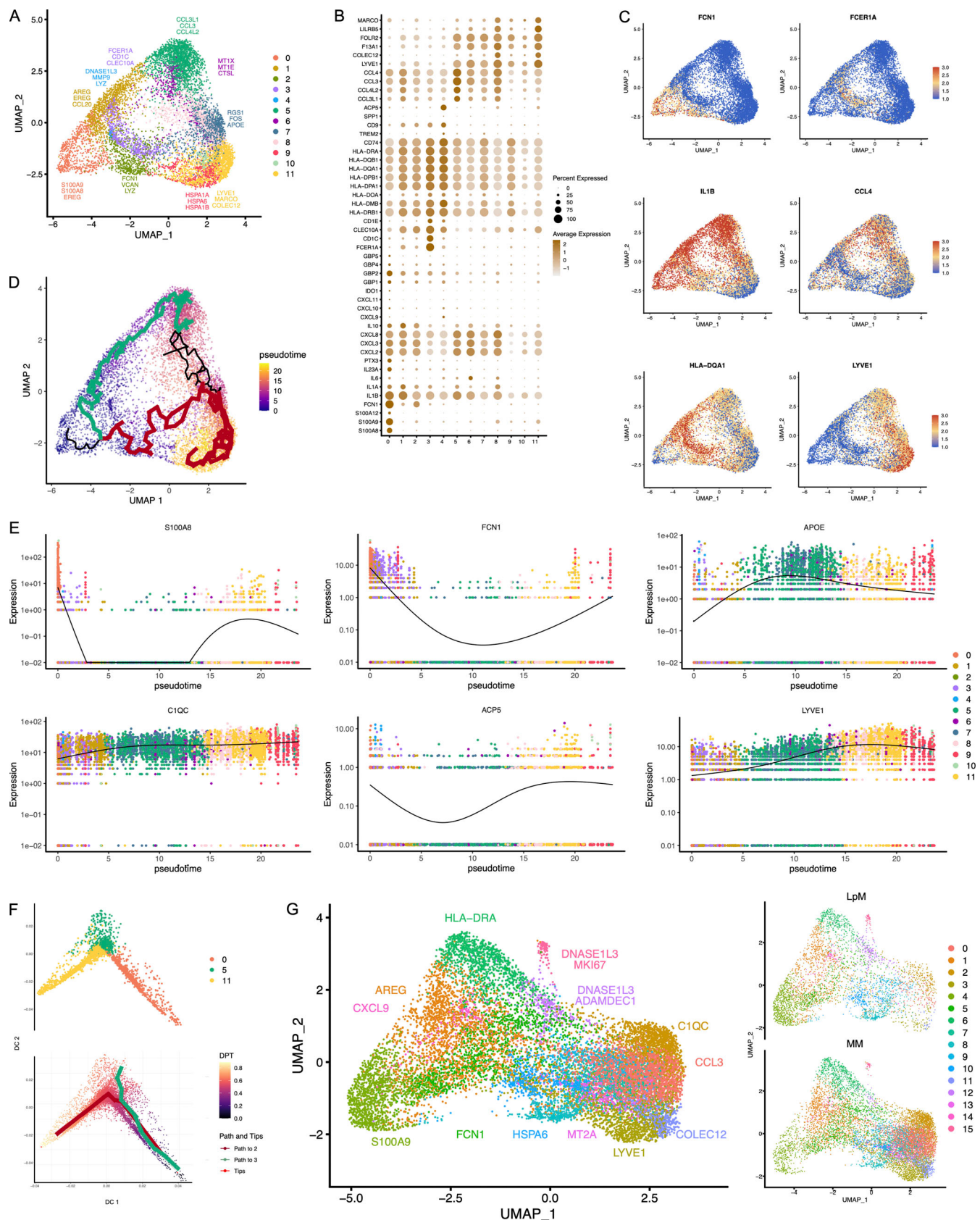


Figure 5. Cluster analysis, gene expression identification, and pseudotime trajectories for MMs. (A) UMAP representation of 12 clusters (different colors) with annotated highly expressed genes within selected clusters. (B) Dot plot of average expression of selected genes in every cluster. Average expression levels are visualized from low (white) to high (brown) expression. The percentage of positive cells is indicated by circle size. (C) UMAP visualization of

selected genes. Expression levels are visualized from low (blue) to high (red) expression. **(D)** Pseudotime trajectory analysis of MMs using monocle. Cells are labeled by pseudotime colored from blue to yellow. The main branches in the trajectory are colored (green and red). **(E)** Expression for selected gene expression through pseudotime. Cell colors represent clusters. **(F)** UMAP representation of clusters 0, 5, and 11 and pseudotime trajectory analysis (lower) of MMs using *destiny* (Angerer et al., 2016). **(G)** UMAP of integrated analysis of LpMs and MMs.

(Fig. S1 D). The clusters transcriptionally most distant from MM₀ could be broadly divided into clusters with proinflammatory and homeostatic properties. MM₅ (and MM₈) expressed high levels of proinflammatory cytokines (e.g., *IL1A* and *IL1B*) and multiple chemokines (e.g., *CXCL2*, *CXCL3*, *CXCL8*, *CCL3*, and *CCL4*), whereas MM₁₁ expressed low levels of proinflammatory cytokines and chemokines but high levels of genes such as *LILRB5*, *MARCO*, *LYVE1*, *FOLR2*, and *COLEC12* (Fig. 5, B and C). Consistently, enriched GO terms for proinflammatory MM₅ were cellular response to LPS and chemokine-mediated signaling pathway, whereas top GO terms for homeostatic MM₁₁ were receptor-mediated endocytosis, synaptic pruning, and apoptotic cell clearance (Fig. 3). Interestingly, high expression of *PMP22* and *EMPI* genes was observed in MM₈ and MM₁₁ (Figs. S2 B and S4 C). These genes are mainly expressed in Schwann cells (Taylor et al., 1995). *PMP22* protein is part of the myelin sheath that protects neurons. High expression of *PMP22* and *EMPI* genes suggests that MMs are phagocytosing Schwann cells and thus are in intimate contact with enteric neurons. Cluster 9, similar to LpM₈ in mucosa, expressed high levels of HSP genes (Fig. S2 B), with enriched GO terms such as unfolded protein responses (Fig. 3). HSPs protect against cellular stress, and it was recently shown that increased expression of HSPs in macrophages concomitant with downregulation of IL-1 had antiinflammatory effects in response to change of diet in experimental mice (Brykczynska et al., 2020). In agreement, HSP⁺ MMs expressed low levels of proinflammatory cytokines and chemokines (Fig. 5 B).

Pseudotime trajectory analysis, using random cells in MM₀ as root, showed a trajectory following two distinct branches (Fig. 5 D). One branch followed through MM₁, MM₅, and MM₆, clusters expressing many proinflammatory genes. Interestingly, all three subsets shared the GO term “positive regulation of angiogenesis” (Fig. 3). Conversely, the homeostatic branch expressed lower levels of proinflammatory genes (Fig. 5 B) and ended in MM₁₁, the cluster most strongly associated with enteric neurons. As for LpMs, typical monocyte-related genes were rapidly downregulated along the trajectory, whereas TRM genes (e.g., *LYVE1*, *CIQC*, and *APOE*) were rapidly upregulated and found to be more broadly expressed by MMs (Fig. 5 E) than by LpMs (Fig. 2 E). To give further support to these findings, we analyzed the data using *Destiny* (Angerer et al., 2016; Haghverdi et al., 2015), an efficient R implementation of the diffusion map algorithm. Here we also found as a function of pseudotime that the trajectory divided into two branches that ended in clusters 5 and 11 (Fig. 5 F).

To directly compare the transcriptomic profile of LpMs and MMs, we performed an integrated analysis. Clustering of the two datasets clearly showed that the transcriptomic profile differed between the two compartments. MMs were dominated in clusters expressing genes associated with

homeostatic functions in muscularis (e.g., *CIQC*, *COLEC12*, *LYVE1*, and *CCL3*), whereas LpMs more dominated by clusters expressing proinflammatory genes (e.g., *SI00A9* and *CXCL9*) and genes involved in antigen presentation (e.g., *HLA-DRA*; Fig. 5 G). Interestingly, a small cluster of cells derived from both compartments expressed genes (e.g., *MKI67*) associated with proliferation.

To determine the anatomic localization of MMs, we performed multicolor immunostaining in situ (Fig. 6). A median of 191 MMs/mm² ($n = 5$) were detected in muscularis propria, and as in the submucosa, most MMs (median >90%) expressed both *LYVE-1* and *COLEC12*. They were distributed throughout the muscularis tissue but were enriched adjacent to neurons (Fig. 6, C and D) and vessels (Fig. 6, A and B). In addition, a minor fraction of calprotectin⁺ MMs (median 6%) was found scattered throughout the tissue (Fig. S4 B).

Together, the results showed that the MM population was very heterogeneous, consisting of multiple functionally distinct subsets. Transcriptomic profiling and pseudotime trajectory analysis revealed significant differences between MMs and their LpM counterparts, indicating that tissue-specific signals from the local microenvironment are important for macrophage differentiation and diversity.

A subpopulation of mucosal and muscularis macrophages express genes compatible with an embryonic origin

Studies in mice have reported that gut macrophages are composed of both monocyte-derived and embryonic-derived macrophages (De Schepper et al., 2018; Shaw et al., 2018). To analyze whether embryonic-derived macrophages also occurred in adult human colon, we examined genes reported to be differentially expressed between lineages. Interestingly, MM₄ and LpM₉ (and MM₁₀) showed higher expression of several genes related to embryonic ontogeny, such as *CD63*, *ADAMDEC1*, and *DNASEIL3* (Fig. 7, A and B; De Schepper et al., 2018; Shaw et al., 2018). Using ClusterMap (Gao et al., 2019), a method to determine cluster similarity across biological samples, we found that MM₄ and LpM₉ showed the highest similarity index (0.21) when all clusters were compared to each other. Moreover, interestingly, in the integrated analysis, cluster 12 (located closest to proliferating macrophages) expressed the highest levels of *ADAMDEC1* and *DNASEIL3* (Fig. 5 G). Transcriptomic analysis of the mucosal macrophage dataset from Elmentaite et al. (2021) showed a similar expression profile (Fig. 7 C). To investigate further the possibility that these clusters contained embryonic-derived macrophages, we examined the transcriptomic profile of intestinal macrophages in human embryos. Fawcner-Corbett et al. (2021) recently published a study analyzing human intestinal development using scRNA-seq. Reexamining the immune cell data of embryos at postconception weeks 12–22, we found that intestinal macrophages contained three distinct subpopulations

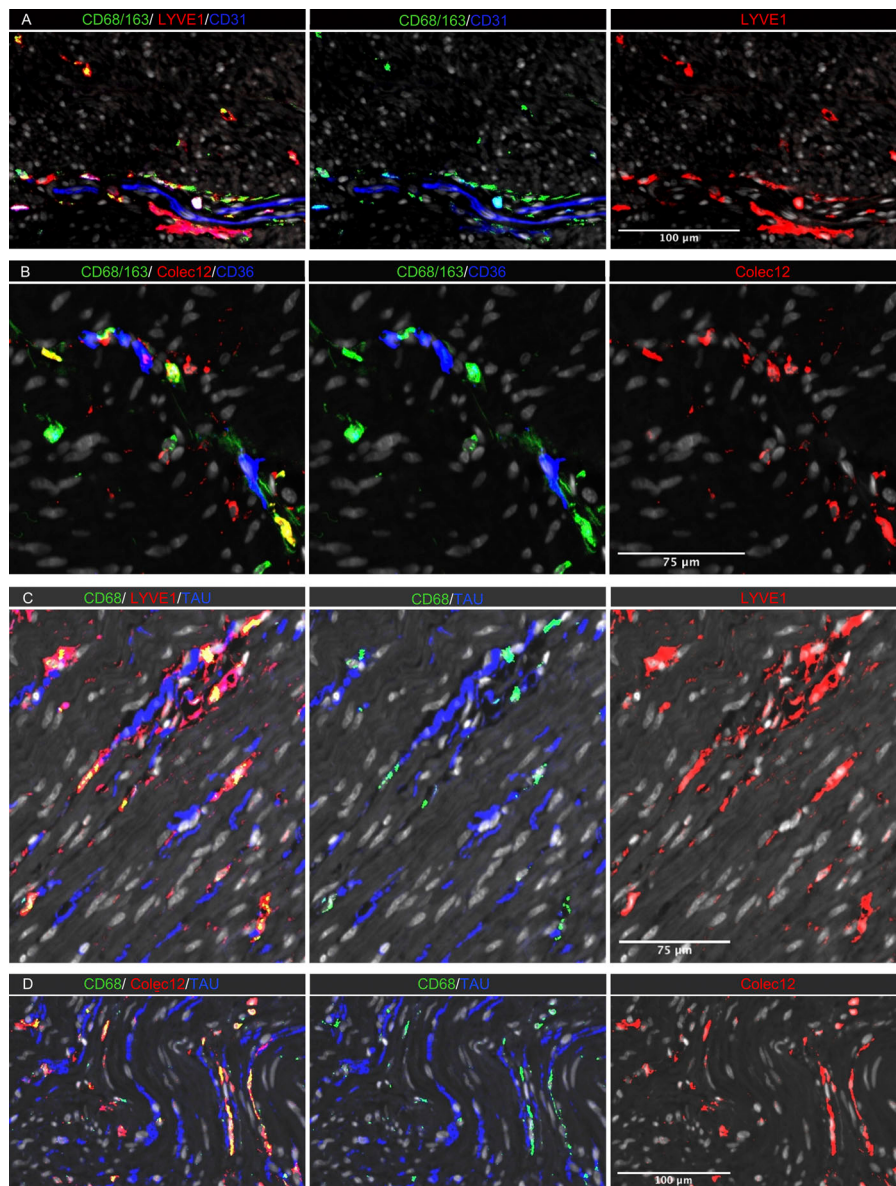


Figure 6. In situ localization of MMs in human colon. (A–D) Sections stained for (A) CD68/CD163 (green), LYVE-1 (red), and endothelial cell marker CD31 (blue); (B) CD68/CD163 (green), COLEC12 (red), and endothelial cell marker CD36 (blue); (C) CD68 (green), neuron marker tau (blue), and LYVE-1 (red); and (D) CD68 (green), tau (blue), and COLEC12 (red). All sections were counterstained with Hoechst DNA-stain (gray). Representative of $n \geq 3$.

(Fig. 7 D). Cluster 3 expressed high levels of monocyte-related genes such as *S100A8*, *S100A9*, *VCAN*, and *FCN1*, whereas clusters 0 and 4 expressed higher levels of genes associated with TRMs such as *CIQA* and *SELENOP* (Fig. 7 D). Interestingly, only cluster 4 expressed high levels of *DNASE1L3* and *ADAMDECI*, similar to embryonic-derived macrophages in mouse colon (Fig. 7 D). To examine whether this phenotype could be retrieved in other tissues with a low component of bone marrow-derived cells, we analyzed a recently published dataset of microglial cells isolated from the human brain (Olah et al., 2020). Integration of datasets from LpMs and microglial cells showed that their transcriptomic profiles were very different, with almost no overlap (Fig. 7 E). As expected, few microglial cells expressed monocyte-like genes (e.g., *S100A8*), and most microglial cells expressed the TRM marker *CIQA*. However, expression of *DNASE1L3* and *ADAMDECI* was not detected (Fig. 7 F).

Together, these results may suggest that a minor population of gut macrophages in human adults are of embryonic ontogeny.

However, the gut “embryonic” signature was not expressed by human microglial cells.

Mucosal and muscularis macrophages interact extensively with resident tissue cells and immune cells

Cell differentiation in tissues is triggered by contacts with other neighboring cells through receptor–ligand interactions. Thus, we interrogated microenvironmental signals that could be involved in the transcriptional reconfiguration process observed in both compartments. We determined such interactions by applying CellPhoneDB2.0 (Efremova et al., 2020), combining our scRNA-seq datasets of LpMs and a published scRNA-seq dataset covering all stromal and immune cells from normal colonic mucosa (Smillie et al., 2019). Numerous statistically significant interactions were found between LpMs and subtypes of fibroblasts, endothelial cells, and epithelial cells (Fig. 8 A). Among receptor–ligand pairs that are particularly important for macrophage survival and differentiation, we found that the

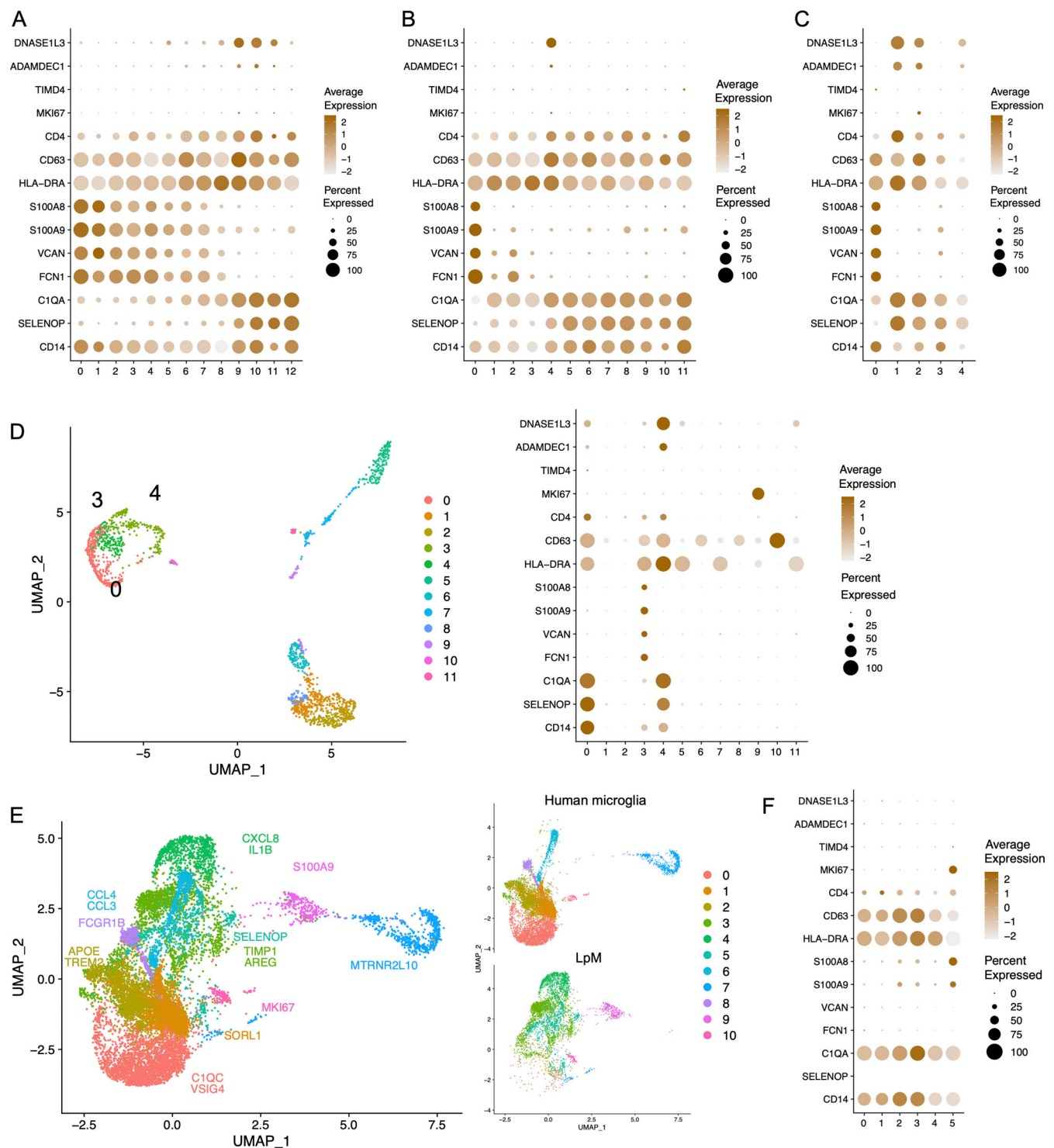


Figure 7. **Expression of genes associated with embryonic-derived macrophages.** (A–C) Dot plot of average expression of selected genes in LpMs (A), MMs (B), and macrophages from Gut Cell Atlas (Elmentaite et al., 2021; C). (D) UMAP plot of embryonic intestinal-derived immune cells (left, macrophage clusters indicated [0, 3, 4]) and dot plot of average expression in fetal macrophage clusters (right). Intestinal samples are from postconception weeks 12–22 (Fawcner-Corbett et al., 2021). (E) UMAPs of integrated analysis of human microglia (Olah et al., 2020) and LpMs. (F) Dot plot of average expression in microglia cell clusters.

CSF1R-ligand cytokines *CSF1* and *IL34* (Lavin et al., 2015) were expressed by postcapillary venules and activated fibroblasts, whereas genes involved in the Notch signaling pathway (e.g., *DLL4/JAG1:NOTCH2*) were expressed by epithelial and stromal

cells (Fig. S5 A). Endothelial cells and fibroblasts expressed many adhesion molecule and chemokine genes that interacted primarily with LpM₀–LpM₈. These results agree with the concept that endothelial cells and fibroblasts are involved in the

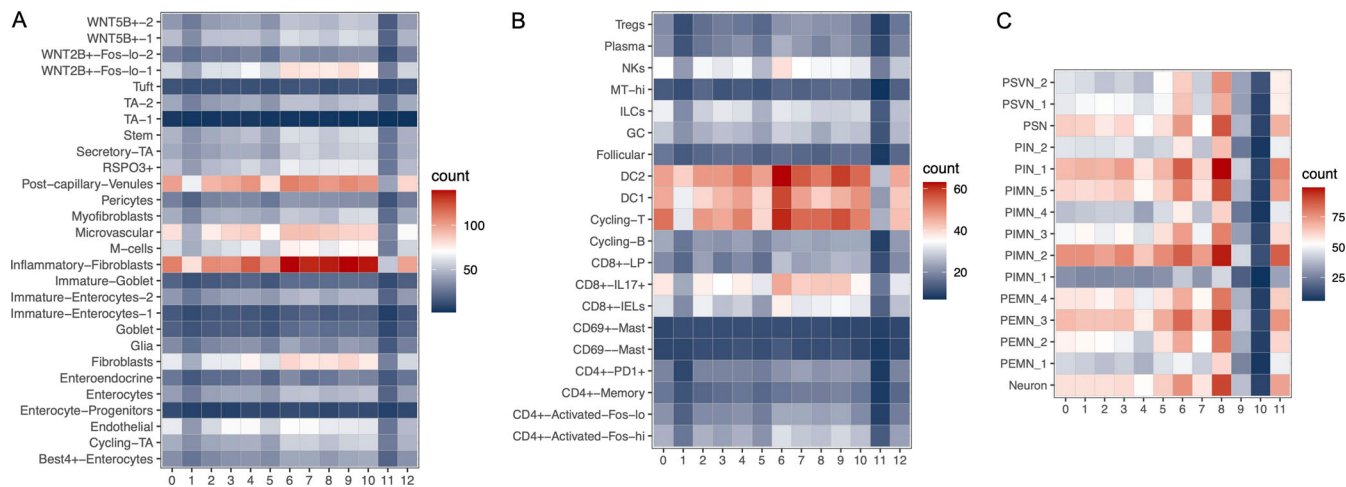


Figure 8. **Cell-cell interaction analysis.** (A–C) Heatmap representing numbers of receptor-ligand pairs between LpM clusters and subtypes of epithelial and stromal cells (A), immune cell lineages (B), and subtypes of enteric neuronal cells (C) from normal colon.

recruitment, migration, and localization of LpMs (Fig. S5 A). LpM₉ and LpM₁₀, on the other hand, displayed interactions with epithelial cells (CDH1:aEb7, DSG2:DSC2; Fig. S5 A), consistent with their subepithelial localization (Fig. 4 C). Importantly, stromal and epithelial cells showed several interactions with LpMs that regulate macrophage functions. This included interactions associated with negative regulation of macrophage activation (LGALS9:HAVCR2, HLA-G:LILRB2, HLA-G:LILRB1, and TNFSF10:RIPK1; Chen et al., 2018; Hartwig et al., 2017; Ocana-Guzman et al., 2016), M2-like polarization (GAS6:PROS1:AXL and GAS6:MERTK; Myers et al., 2019; Myers et al., 2019), and “don’t eat me” signals (CD47:SIRPA and CD52:SIGLEC10; Barkal et al., 2019; Li et al., 2021). On the other hand, LpMs expressed ligands (EGFR and ERBB3) for receptor tyrosine kinases expressed by epithelial cells and activated fibroblasts, suggesting a bidirectional regulation of survival and function between LpMs and tissue resident cells. LpMs also showed numerous interactions with immune cell subtypes (Fig. 7 B). In particular, LpM₆, LpM₇, and LpM₉ (as well as DC3-like LpM₈), which expressed high levels of MCH class II genes (Fig. 2 B), showed interactions with cycling, memory, and regulatory T cells (e.g., CD28:CD80, CLTA4:CD80, CD28:CD86, and CLTA4:CD86; Fig. S5 B), suggesting that LpM subsets play a role as regulators of T cell responses in the colonic mucosa. Moreover, selective expression of several CXCR3-binding chemokines were expressed by LpM₆ (Figs. 2 B and S5 B), indicating a role in the recruitment of CD8 T cells.

Studies in mice have shown that MMs cross talk with enteric neurons (De Schepper et al., 2018; Muller et al., 2014). To interrogate macrophage–neuron interactions in muscularis propria, we analyzed our MM scRNA-seq dataset together with a published scRNA-seq dataset of the human enteric neuron system (Drokhlyansky et al., 2020). A very high number of interactions between multiple neuron subtypes and MMs was found (Fig. 8 C). Central to MM survival and differentiation, several neuron subtypes expressed Notch ligands (DLL1, DLL3, and JAG2) and IL34 interacting NOTCH2 and CSF1R on MMs, respectively. Furthermore, numerous receptor-ligand pairs were involved in

macrophage migration, localization, and activation/regulation (Fig. S5 C). Finally, MM–neuron interactions were associated with synapse pruning (C3:C3AR1 and C5:C5AR2; Hong et al., 2016) and neuron stimulation (BMP2:BMP2R), strengthening the notion that there is an extensive cross talk between MMs and enteric neurons in the human colon that controls gastrointestinal motility (De Schepper et al., 2018; Muller et al., 2014). Together, our results indicated that macrophages interact extensively with tissue-resident cells in both compartments, compatible with the concept that the local microenvironment is important for imprinting of macrophage specialization and niche-specific localization.

Gene regulatory network analysis indicates that a limited number of TFs control macrophage differentiation and diversity

To identify TFs that may control transcriptional programming of LpMs and MMs, we used single-cell regulatory network inference and clustering (SCENIC; Van de Sande et al., 2020) to determine sets of genes coexpressed with their associated TFs (regulons). We found a total of 185 and 187 regulons (active in >1% of the cells) for LpMs and MMs, respectively, with significantly enriched motifs for the corresponding TFs. By ordering macrophage clusters along the pseudotime trajectory, we identified a restricted number of regulons corresponding to specific clusters. LpM₀ showed increased regulon activity corresponding to RXRA and IRF7, whereas LpM₂ was associated with several NF- κ B family members (NFKB1, NFKB2, and REL; Fig. 9 A). LpM₆ was associated with STAT1, whereas LpM₈–LpM₁₁ and LYVE1⁺ SmMs showed regulon activity driven by TFs such as MAF, MAFB, HES1, and EGRI. MAF and MAFB are known for their role in driving terminal macrophage differentiation (Aziz et al., 2009).

The gene regulatory network in MMs showed similarities with LpMs (Fig. 9 B). MM₀ showed regulon activity corresponding to NF- κ B family members (NFKB1, RELB, and BCL11A), whereas regulons corresponding to MAF and MAFB were upregulated in MM₁₁. On the other hand, MM₅ (and to a lesser extent

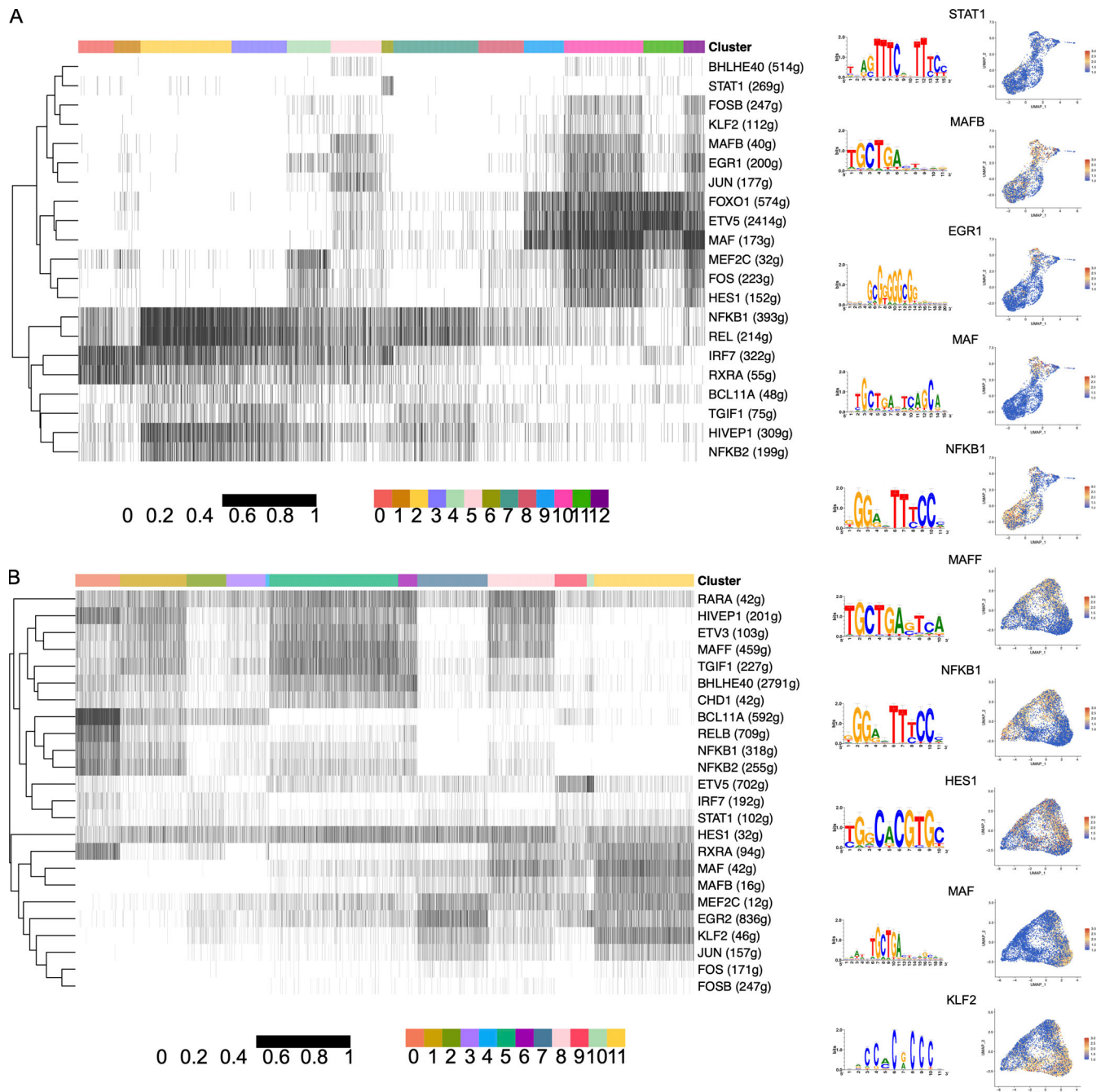


Figure 9. **Identification of TF modules based on regulon activities in LpMs and MMs.** (A and B) Binarized heatmaps of regulon activity in LpMs (A) and MMs (B) with the representative TF and part of their binding motifs. Numbers in parentheses indicate the number of genes enriched in the regulons, and UMAP plots show expression of selected TF.

MMs) expressed regulons corresponding to TFs such as *MAFF*, *RARA*, and *BHLHE40*, not found in the mucosal compartment. Interestingly, regulon activity with corresponding *HES1* was broadly expressed by MMs and by subsets of LpMs (Fig. 8, A and B). *HES1* is a classic TF downstream of the Notch signaling pathway (Sharma et al., 2020). Together with our cell-cell interaction data (Fig. S5, A and C), expression of *HES1* suggests that the Notch signaling pathway is involved in reprogramming of LpMs and MMs, as shown for monocyte-derived macrophages in the liver (Ramachandran et al., 2019; Sharma et al., 2020). In

summary, we found that regulon activity corresponds to distinct macrophage clusters following the pseudotime trajectory, indicating that a limited number of key TFs control the process of monocyte-to-macrophage differentiation observed in both compartments.

Discussion

Using high-resolution spatiotemporal single-cell analysis, we show that macrophage populations in mucosa and muscularis

propria of the human colon contain multiple transcriptionally distinct subsets that display niche-specific localizations and functional properties. Our results also reveal tissue-specific cell-cell interactions and a limited number of TFs that may be key players in the extensive monocyte-to-macrophage reprogramming and subtissular-specific localization observed.

Studies in mice show that macrophages in different organs display functional differences imprinted by tissue-specific cues from the local microenvironment (Guilliams et al., 2020; Okabe and Medzhitov, 2014). Moreover, it was recently shown that heterogeneous macrophages with different transcriptional programs are governed by subtissular niches (Chakarov et al., 2019). Therefore, to understand the functional diversity of macrophages within a tissue, detailed spatiotemporal characterization of the macrophage population and their interactions with neighboring cells is needed. However, the nature of molecular signals that drive macrophage differentiation in the human gut is poorly characterized. Studies in humans and mice have clearly shown that mucosal macrophages in the intestine largely originate from bone marrow-derived monocytes (Bain et al., 2014; Bujko et al., 2018). Thus, probing colonic mucosal macrophages in histologically normal tissue is a unique possibility to study the monocyte-to-macrophage differentiation process *in situ* under steady-state conditions.

Integration of our data suggests that circulating monocytes are elicited through postcapillary venules in the crypt area, after which some cells rapidly differentiate into different types of proinflammatory macrophages and others migrate to the subepithelial region, where they upregulate expression of genes related to endocytosis, wound healing, and antigen presentation, while concomitantly downregulating proinflammatory genes. This latter subpopulation is thus ideally equipped to maintain mucosal barrier integrity with minimal collateral damage.

Our cell-cell communication analysis suggested that the survival, migration, and differentiation of LpMs are controlled by extensive interaction with tissue-resident cells (endothelial cells, fibroblasts, and epithelial cells). Stromal cells may provide a supply of macrophage-trophic factors (e.g., CSF-1 and IL-34), which are crucial for macrophage development and survival (Lavin et al., 2015), whereas several cell types expressed Notch ligands, suggesting that the Notch signaling pathway plays a role in monocyte-to-macrophage reprogramming (Sharma et al., 2020). Several receptor-ligand pairs identified have been shown to have inhibitory or antiinflammatory effects on macrophages, in line with the idea that most tissue macrophages are tolerogenic under steady-state conditions. Interestingly, however, many of these macrophage genes have been shown to promote tumor growth and may be targets for cancer immunotherapy (e.g., *SIRPA*, *SIGLEC10*, *AXL*, *MERTK*, and *RIPK1*; Barkal et al., 2019; Li et al., 2021; Myers et al., 2019; Wang et al., 2018). Thus, in strategies to target these genes to treat cancer, possible side effects should be taken into account, such as drug-induced colitis, as observed by current checkpoint inhibition (Luoma et al., 2020).

Our gene regulatory network analysis identified cluster-specific TFs that are likely to control subset-specific transcriptional programs, which strengthens the idea that the LpM

population contains multiple transcriptionally stable macrophage subsets that coexist in the tissue. Interestingly, the transcriptomic profile of several LpM subsets is similar to that of macrophages associated with various inflammatory and fibrotic diseases (Martin et al., 2019; Qu et al., 2020; Ramachandran et al., 2019). This suggests that these disease-promoting macrophages also play a role under steady-state conditions, but that the relative contribution of functionally different LpM subsets is important for maintenance of homeostasis.

Analysis of muscularis propria showed that many MMs were positioned in close contact with nerves, and cell-cell interaction analysis displayed numerous MM–neuron interactions. Such interactions included BMP2:BMP2R and CSF1R:IL34. As previously shown in mouse models (Gabanyi et al., 2016), this finding suggests that MMs expressing *BMP2* regulate peristaltic motility in the colon by activating *BMP2R* on enteric neurons, whereas neurons expressing IL34 feedback on CSF1R⁺ MMs by stimulating their survival and differentiation. Most MMs expressed high levels of C1Q genes, and we identified interactions between genes of the complement system (C3:C3AR1 and C5:C5AR2), suggesting that MMs are involved in synapse pruning (Stephan et al., 2012). Finally, MMs contained transcripts specific for Schwann cells, indicating that MMs phagocytose cellular material from such nerve-protecting cells. Together, these findings are in agreement with the concept that MM–neuron cross talk plays a pivotal role in gut homeostasis (De Schepper et al., 2018; Muller et al., 2014). MMs were also positioned adjacent to vessels, and the GO term “positive regulation of angiogenesis” was enriched in MM₁, MM₅, and MM₆. Pseudotime trajectory analysis showed that these subsets were found along the proinflammatory branch, whereas neuron-associated MM subsets (first of all MM₁₁) were linked to the homeostatic branch. This suggests that functionally specialized neuron- and blood vessel-associated MMs coexist in the human colon to ensure proper functioning of enteric neurons and blood vessels, respectively (De Schepper et al., 2018).

Several studies in mice have suggested that macrophages important for vascular integrity selectively express LYVE-1 (Chakarov et al., 2019; Lim et al., 2018). However, we found that the vast majority of MMs, those associated with both vessels and nerves, expressed LYVE-1 and COLEC12, and we were not able to distinguish these subsets by *in situ* staining. We also identified calprotectin⁺ MMs scattered throughout the muscularis propria. Clustering and pseudotime trajectory analysis strongly suggested that the vast majority of TRMs found in the muscularis propria originated from these incoming monocytes. Macrophages are the dominating leukocyte population in muscularis propria and submucosa. Interestingly, to this end, we found that MM subsets and SmM expressed high levels of several monocyte-attracting chemokines (e.g., *CCL3*, *CCL4*, *CCL3L1*, and *CCL4L2*), suggesting that TRMs in these compartments are important for continuous recruitment of monocytes.

Somewhat surprisingly, we found that subpopulations of LpMs and MMs showed phenotypic similarities with embryonic-derived macrophages in mouse colon (De Schepper et al., 2018) and with colonic macrophages in human fetus (Fawcner-Corbett et al., 2021). As far as we know there are no established markers

to separate embryonic-derived from bone marrow monocyte-derived macrophages in humans, and evidence to suggest that human macrophages in adult life originate from embryonic precursors is sparse. We and others have shown that human macrophages in various tissues may live for several years (Eguiluz-Gracia et al., 2016; Patel et al., 2021). However, the origin of these long-lived cells has not been determined.

The embryonic signature was not detected in a dataset of human microglial cells, and others have shown that a transcriptional program of fetal macrophages reemerges in skin macrophages during inflammation (Reynolds et al., 2021). However, it is intriguing that a small subset of human colonic macrophages under steady-state conditions in adults express markers similar to macrophages in fetal colon, and that this subset is transcriptionally close to “proliferating” macrophages (Fig. 5 G). Thus, it should be studied further whether genes such as *DNASE1L3* and *ADAMDEC1* could be useful markers to distinguish macrophage lineages in the human intestine.

Together, these results show that LPMs and MMs are extremely heterogeneous and consist of several subsets with distinct functional properties. It appears that maintenance of homeostasis depends on coexistence of both proinflammatory and protective/homeostatic subtypes. Our data also give insights into cell-cell interactions and key TFs that are likely to control tissue-specific macrophage reprogramming. This work constitutes an important framework to understand the complexity of macrophage biology in the human gut and to identify potential targets to better treat inflammatory disorders and cancer in the future.

Materials and methods

Patients and tissue samples

Colonic resections were obtained from patients receiving surgery for colon cancer at Akershus University Hospital (Ahus). The resected colon was immediately examined by an experienced pathologist, and macroscopically normal colon tissue, ≥ 10 cm from the tumor, was placed in vials with RPMI 1640 and put on ice for transport. The study was performed in accordance with the Declaration of Helsinki. Written informed consent was obtained from all participants, and the study was approved by the Regional Committee for Medical Research Ethics (REK, 2018/703, Health Region South-East, Norway). For scRNA-seq, specimens from sigmoid colon were obtained (age 62–78 yr, three males, $n = 4$). For immunofluorescence staining, samples from ascending, transverse, descending, and sigmoid colon were included (age 62–78 yr, four males, $n = 8$). None of the patients had autoimmune, infectious, or inflammatory diseases or had received neoadjuvant chemotherapy or radiation therapy before the operation. All patients received surgery according to the national guidelines.

Single-cell dissociation

Resected colonic tissues were processed within 2 h after removal from the patient. Single-cell suspensions of colonic resections were obtained using a modified version of a previously published protocol (Bujko et al., 2018). The intestinal specimens

were opened longitudinally and washed in Dulbecco's PBS. The muscularis propria was first removed with scissors, after which the mucosa was dissected in narrow strips. The mucosal fragments were then incubated with shaking in PBS with 2 mM EDTA (Sigma-Aldrich) and 1% FCS (Sigma-Aldrich) three times for 15 min at 37°C. The remaining tissue was minced and digested with stirring for 60 min in complete RPMI (RPMI 1640 [Lonza] supplemented with 10% FCS, 1% penicillin/streptomycin [Lonza] and containing 0.25 mg/ml Liberase TL [Roche] and 20 U/ml DNase I [Sigma-Aldrich]). Digested cell suspension was passed through a 100- μ m filter and washed.

Flow cytometry and cell sorting

Released tissue cells were stained in aliquots of 1×10^6 cells/100 μ l PBS with 2% FCS (Gibco) and 0.1% NaN₃ for 30 min on ice. Nonspecific staining was blocked with 10 μ l FcR Blocking Reagent (Miltenyi Biotec) before staining. Dead cells were excluded by TO-PRO-1 Iodide (Thermo Fisher Scientific) or Fixable Viability Dye eFluor 780 (eBioscience) staining. Analysis was performed with a BD LSRFortessa X-20 and sorting with a FACS Aria IIIu (BD Biosciences) running BD FACSDIVA 9.0 software. Purity of $>98\%$ was achieved in sorted populations. Data were processed with FlowJo 10.6.1 (TreeStar). Intracellular staining was performed after surface staining, using a Fixation & Permeabilization Buffer Set (eBioscience) according to the manufacturer's instructions. The following antibodies were used for staining: HLA-DR PerCP/Cy5.5 (307630, clone L243), CD45 BV510 (103138, clone 30-F11), CD3 FITC (300406, clone UCHT1), CD19 Ax488 (302219, clone HIB19), CD14 APC (325608, clone HCD14), CD14 PE/Cy7 (325618, clone HCD14), CD1c BV421 (331526, clone L161), and Fc ϵ RI PE (334610, clone AER-37) from BioLegend; FcR Blocking Reagent (130-059-901) and CD141 PE (130-113-318, clone AD5-14H12) from Miltenyi Biotec; CD14 PE/Cy7 (562698, clone M ϕ P9) and CD11c APC (333144, clone S-HCL-3) from BD Biosciences; epithelial antigen FITC (F0860, clone Ber-EP4) from Agilent; C1Q FITC (F025402-2) from Dako; and calprotectin PE (MCA874PE, clone MAC387) from Bio-Rad.

scRNA-seq

Cellular suspensions ($\sim 15,000$ cells, with expected recovery of $\sim 7,500$ cells) of sorted CD45⁺HLA-DR⁺CD14⁺ macrophages from colonic mucosa and muscularis propria were loaded on the 10X Chromium Controller instrument (10X Genomics) according to the manufacturer's protocol using 10X GEMCode proprietary technology. All samples from individual patients were loaded in one batch. The Chromium Single Cell 3' v2 Reagent kit (10X Genomics) was used to generate the cDNA and prepare the libraries, according to the manufacturer's protocol. The libraries were then equimolarly pooled and sequenced on an Illumina NextSeq500 using HighOutput flow cells v2.5. A coverage of 400 million reads per sample was targeted to obtain 50,000 reads per cell. The raw data were then demultiplexed and processed with the Cell Ranger software (10X Genomics) v2.1.1.

Preprocessing scRNA-seq data

In total, we analyzed 63,917 human cells from donors ($n = 4$). We aligned the reads of the input dataset to the GRCh38 reference

genomes and estimated cell-containing partitions and associated unique molecular identifiers using the Cell Ranger Chromium Single Cell RNA-seq v3.0.2. We performed data preparation using Seurat R packages. Genes expressed in fewer than three cells in a sample were excluded, as well as cells that expressed fewer than 200 genes and mitochondrial gene content >5% of the total unique molecular identifier count. We normalized data by using gene counts for each cell that were divided by the total counts for that cell and multiplied by 10,000 and then log-transformed. Subsequently, we identified genes that were outliers on a mean variability plot using the *vst* method with 2,000 genes. For mucosa and muscularis data, we separately found integration anchors and then performed data integration using a precomputed anchorSet with default parameters. Finally, we scaled data and centered genes in the dataset using linear model.

Dimensionality reduction, clustering, and differential expression analysis and data integration

We ran principal component analysis dimensionality reduction with 30 principal components to compute and store (on 2,000 variable genes). We estimated dimensions of reduction parameter (for LpM = 13 and for MM type = 20) and constructed a shared nearest neighbor graph for given datasets. We first determined the *k*-nearest neighbors of each cell. We used this *k*-nearest neighbor graph to construct the shared nearest neighbor graph by calculating the neighborhood overlap (Jaccard index) between every cell and its 20 nearest neighbors. To obtain the resolution parameter, we used the *clustree* R package, with resolution varying from 0.1 to 2.0. We got resolution parameters for LpM = 0.7 and for MM = 0.6. We then ran the UMAP dimensional reduction technique with principal component analysis dimension reduction and found DEGs for each of the clusters in the datasets.

For data integration and batch correction, we used Seurat integration methods *FindIntegrationAnchors* and *IntegrateData*, with default parameters. We applied integration for LpMs and MMs, microglial cells (Olah et al., 2020), and LpMs, and LpMs and macrophages from Gut Cell Atlas (Elmentaite et al., 2021).

All heatmaps, UMAP visualizations, violin plots, and dot plots were produced using Seurat functions in conjunction with the *ggplot2*, *pheatmap*, and *grid* R packages. *ClusterMap* (Gao et al., 2019) was used to compute a similarity metric for subclusters between LpMs and MMs.

Developmental trajectory inference and transcriptional regulation

To generate pseudotemporal dynamics, we used the *Monocle* R package. We ordered cells in a semisupervised manner based on their Seurat clustering, scaled the resulting pseudotime values from 0 to 1, and mapped them onto UMAP visualizations generated by Seurat. DEGs along this trajectory were identified using Moran's *I* test. To perform pseudotime trajectory analysis, we also used diffusion maps with default parameters setup from *Destiny* R package (Angerer et al., 2016).

For TF analysis, we obtained a list of all genes identified as human TFs. To analyze TF regulons further, we adopted Single Cell Regulatory Network Inference and Clustering (SCENIC; Van

de Sande et al., 2020), using default parameters and the normalized data matrices from Seurat as input. SCENIC is a combination of three packages (*GENIE3*, *RcisTarget*, and *AUCell*). For motif visualization, we obtained the highest normalized enrichment score of the motif in the gene set.

Identification of significant ligand-receptor pairs

For comprehensive systematic analysis of interlineage interactions, we used *CellPhoneDB2.0* (Efremova et al., 2020). *CellPhoneDB2.0* is a manually curated repository of ligands, receptors, and their interactions, integrated with a statistical package for inferring cell-cell communication networks from single-cell transcriptomic data. This package searches for ligand-receptor interactions and outputs multiple result files based on curated databases such as UniProt, IUPHAR, and Ensembl.

Each dataset was analyzed using matrices from Seurat and datasets covering epithelial, endothelial, fibroblast, and immune cell (Smillie et al., 2019) subsets and enteric neurons (Drokhlyansky et al., 2020). Significant ligand-receptor pairs identified from datasets, with adjusted *P* value <0.05, were extracted, requiring the ligand and receptor to be expressed in ≥10% of the cells.

Immunofluorescence staining

Sections of formalin-fixed and paraffin-embedded tissue were cut in series at 4 μm, mounted on Superfrost Plus object glasses (Thermo Fisher Scientific), and washed sequentially in xylene, ethanol, and PBS. Heat-induced epitope retrieval was performed by boiling sections for 20 min in citrate buffer (pH 6.0) and cooling to room temperature (RT) before staining. Sections were incubated with mixtures of primary antibodies for 1 h at 37°C, rinsed in PBS, and incubated with secondary antibodies for 1.5 h at RT. The following primary antibodies were used: CD68 (M087601-2, clone PG-M1), C1q Complement/FITC (F025402-2), and CD31 (M082329-2, clone JC70A) from Dako Agilent; CD36 (14347S, clone D8L9T) and Tau (4019S) from Cell Signaling Technology; LYVE1 (ab10278) and ACP5 (ab238033, clone rACP5/1070) from Abcam; CD163 (NCL-L-CD163) from Leica Biosystems; C25H (LS-B14159-50, clone aa142-247) from LS Bio; Clec12/CL-P1 (AF2690) from R&D Systems; and anti-human calprotectin from Calpro. The following reagents served as secondary antibodies: donkey anti-goat IgG Alexa Fluor 555 (A-21432), donkey anti-rabbit IgG Alexa Fluor 647 (A-31573), goat anti-mouse IgG3 Alexa Fluor 488 (A-21151), donkey anti-mouse IgG Alexa Fluor 488 (A-21202), goat anti-mouse IgG1 Alexa Fluor 555 (A-21127), goat anti-mouse IgG2b Alexa Fluor 555 (A-21147), and goat anti-mouse IgG1 Alexa Fluor 647 (A-21240) from Thermo Fisher Scientific; donkey anti-rabbit IgG Cy3 (711-165-152) from Jackson ImmunoResearch; rat anti-mouse IgG3 Alexa Fluor 488 (clone SB76b, ab172328) from Abcam; and rat anti-mouse IgG1-Alexa Fluor 647 (clone SB77e, 1144-31) from SouthernBiotech. Sections were then incubated for 5 min at RT in Hoechst 33342 nucleic acid stain (Thermo Fisher Scientific), and stained sections were mounted with ProLong Glass Antifade mountant (Molecular Probes). Laser scanning confocal microscopy was performed by acquiring tile scans on an Andor

Dragonfly equipped with a fusion stitcher. The Andor Dragonfly was built on a Nikon TiE inverted microscope equipped with a 60×/1.40-NA oil-immersion objective. To determine cell densities, the total number of positive cells for all staining combinations was counted in an average of 1.7-mm² tissue area for every patient in both the mucosa and muscularis propria.

Online supplemental material

Fig. S1 shows the sorting strategy, feature plots of canonical macrophage markers, and flow cytometric detection of DC3. **Fig. S2** shows heatmaps of top DEGs in LPM and MM clusters. **Fig. S3** shows sample representation in clusters. **Fig. S4** shows in situ immunostaining of COLEC12 and Calprotectin, as well as feature plots of Schwann cell markers in MMs. **Fig. S5** shows bubble plots of ligand-receptor pairs between macrophages and other colonic cells.

Data availability

scRNA-seq datasets generated in this study are deposited in the European Genome-Phenome Archive under the following accession numbers: EGAD00001007765 and EGAS00001005377. In addition, all data are available upon request.

Reference datasets used in this study are as follows: human ulcerative colitis scRNA-seq dataset (Smillie et al., 2019) Single Cell Portal: SCP259; human enteric neuron system scRNA-seq dataset (Drokhlyansky et al., 2020) Broad Institute Single Cell Portal: SCP1038; human fetal intestinal scRNA-seq dataset (Fawcner-Corbett et al., 2021) Gene Expression Omnibus: GSE158702; GUT HUMAN ATLAS scRNA-seq dataset (Elmentaite et al., 2021) <https://www.gutcellatlas.org/>; and human microglia scRNA-seq dataset (Olah et al., 2020; <https://www.synapse.org/#!Synapse:syn21438358>).

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Supplemental material

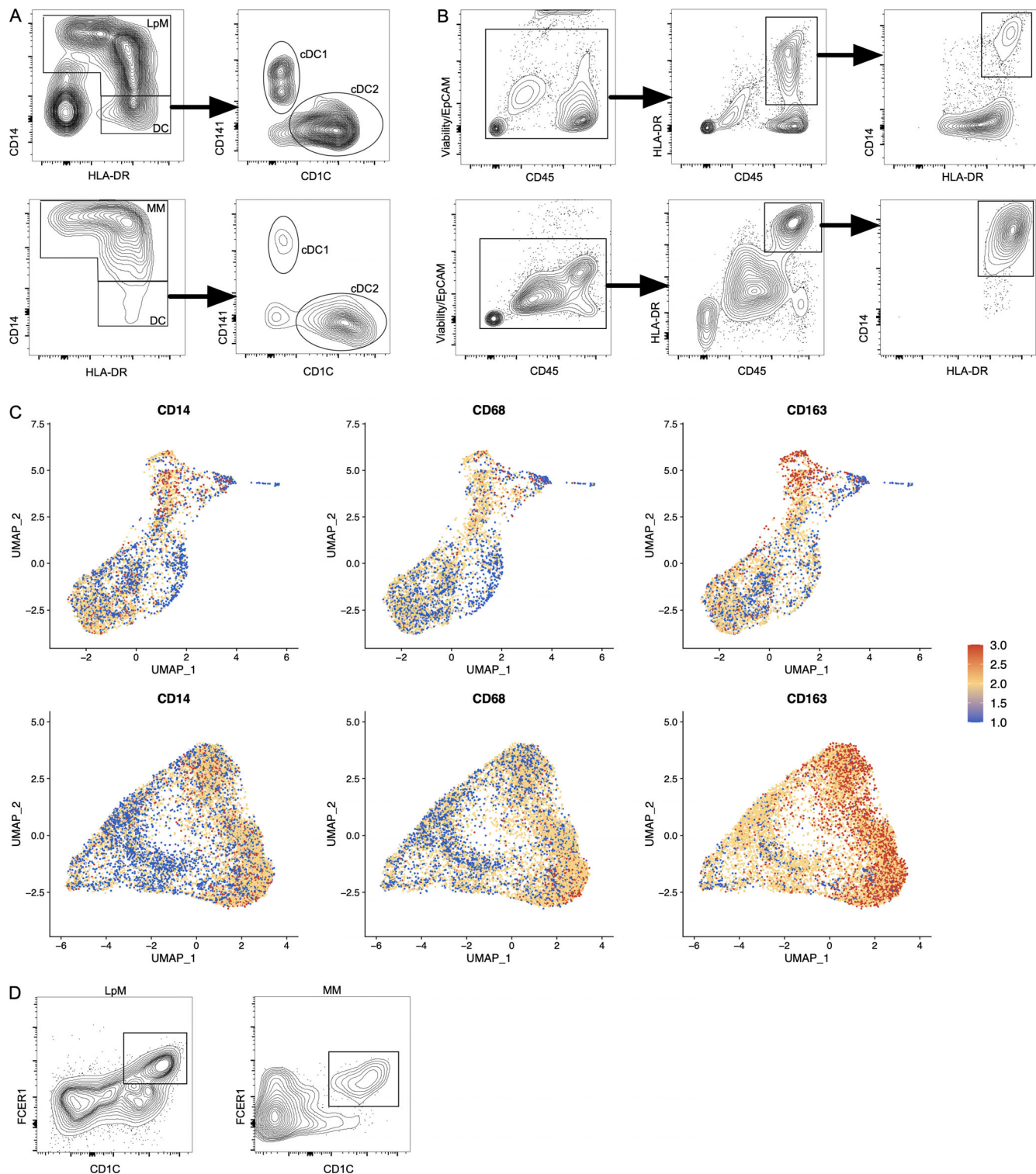
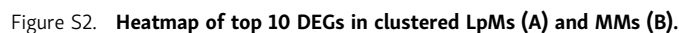


Figure S1. **Characterization and flow cytometry sorting strategy for LpMs and MMs.** (A) Flow cytometry plots of live-gated CD45⁺CD3⁻CD19⁻ cells showing that HLA-DR⁺ cells are separated into CD14⁺ macrophages (LpMs, top left; MMs, bottom left) or CD14⁻ DCs, where the latter comprise CD141⁺ cDC1 and CD1C⁺ cDC2. (B) Flow cytometry gating strategy to target CD14⁺ LpMs (top) and CD14⁺ MMs (bottom). (C) UMAP plots of selected macrophage-related genes in LpMs (top) and MMs (bottom). Expression levels are visualized from low (blue) to high (red) expression. (D) Flow cytometry plots showing CD14⁺CD1C⁺FCER1⁺ cells in mucosa (left) and muscularis propria (right). All flow cytometry plots are representative of more than four donors.



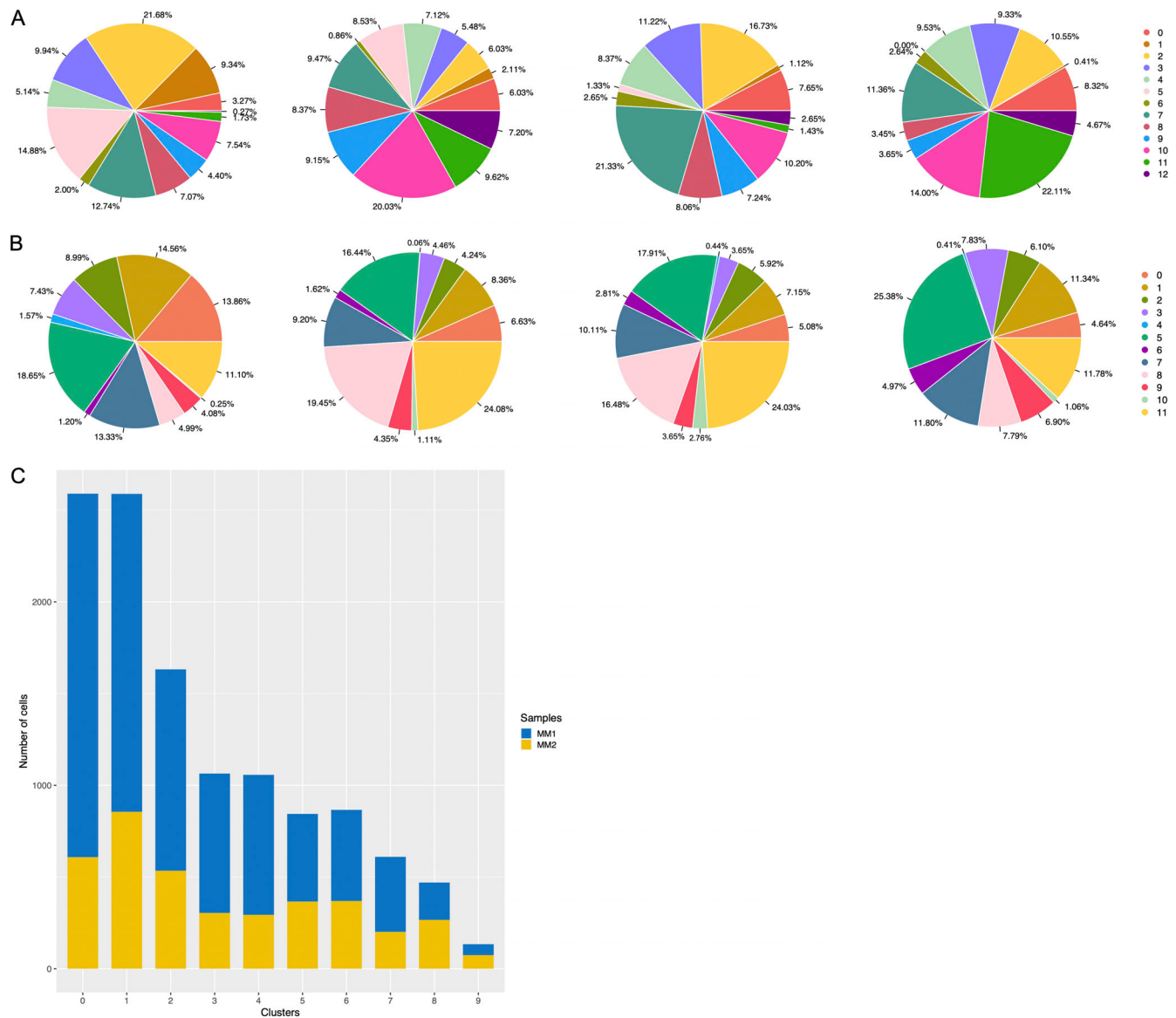


Figure S3. **Representation of cell frequencies from donors and compartments. (A and B)** Cluster size of LpMs (A) and MMs (B) shown for each donor. **(C)** Representation of cells in MM clusters obtained from two sites (MM1, MM2) of muscularis propria ($n = 3$).

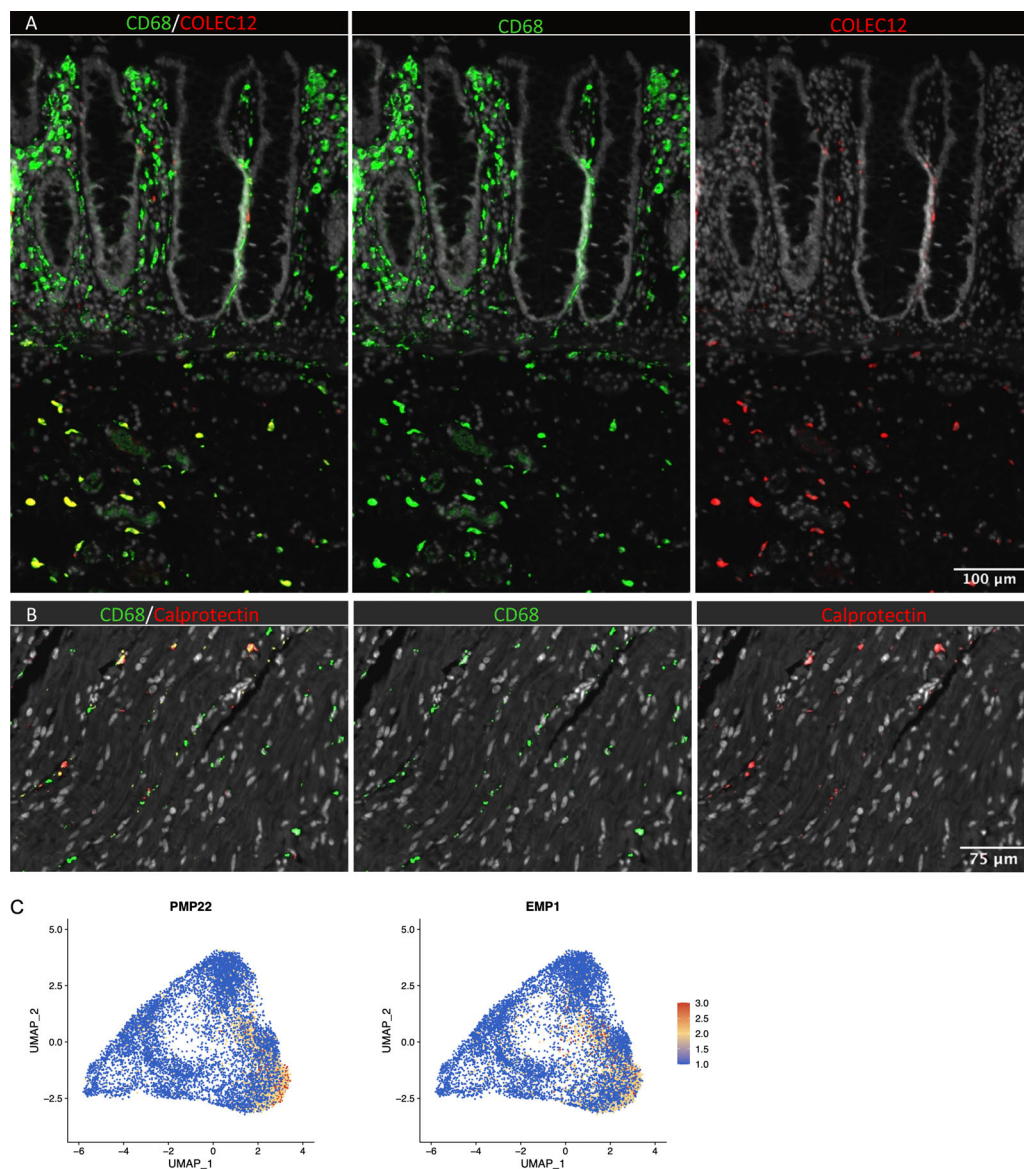


Figure S4. **In situ localization of macrophages in human colon and expression of Schwann cell markers.** (A) Section of mucosa and submucosa stained for CD68/CD163 (green) and COLEC12 (red). (B) Section of muscularis stained for CD68/CD163 (green) and calprotectin (S100A8/S100A9; red). Sections were counterstained with Hoechst DNA-stain (gray). Representative of $n \geq 3$. (C) UMAP plot of MMs showing the expression of PMP22 and EMP1. Expression levels are visualized from low (blue) to high (red) expression.

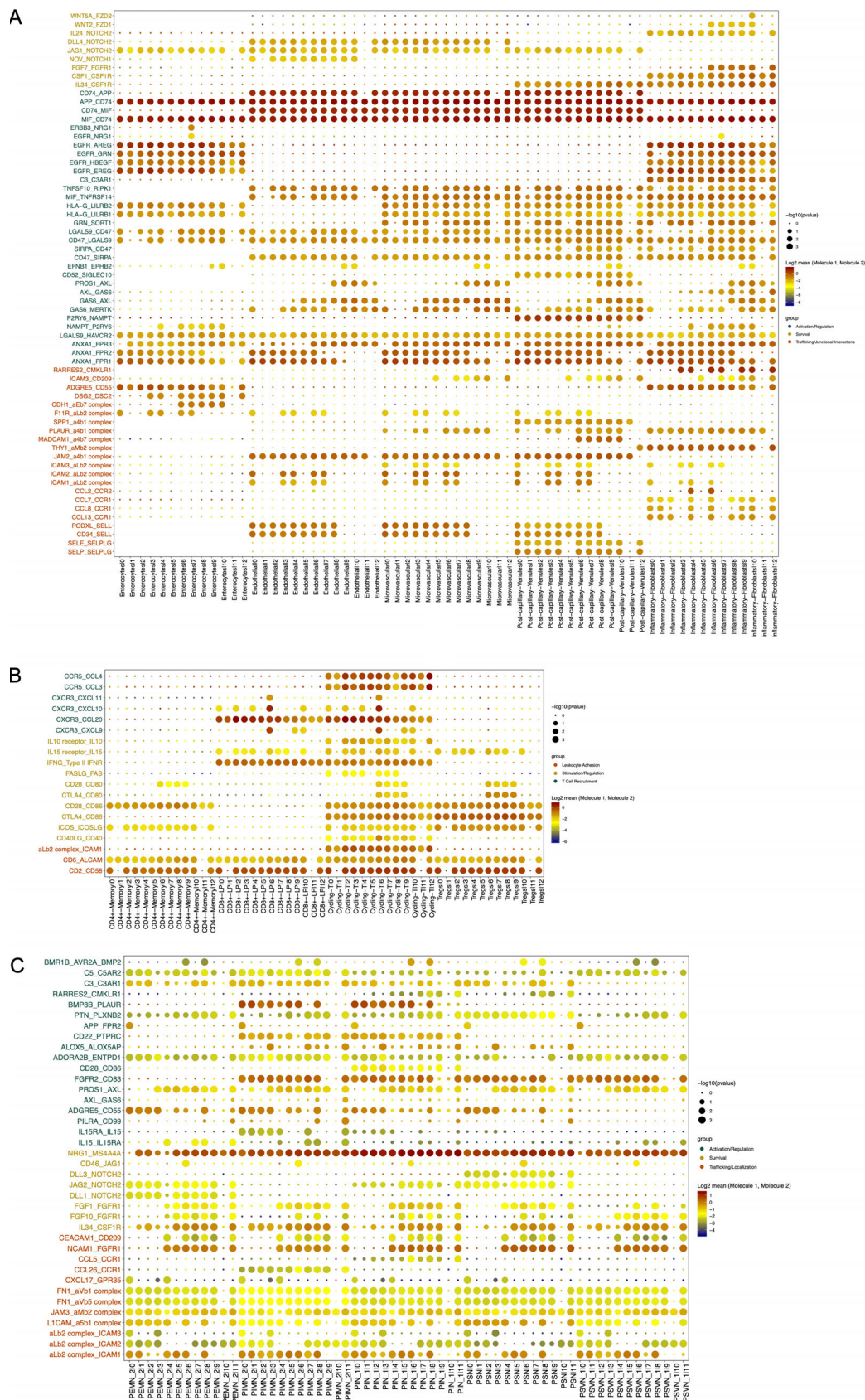


Figure S5. Cellular interactions between colonic macrophages and other cell types in colon. (A and B) Dot plot of interactions between LpMs and epithelial cells and stromal cells (A) and immune cells (B). **(C)** Dot plot of interactions between MMs and subtypes of enteric neurons. Rows represents ligand-receptor pairs, and columns define cell-cell interaction pairs.