Redefining the heterogeneity of peripheral nerve cells in health and autoimmunity

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Peripheral nerves contain axons and their enwrapping glia cells named Schwann cells (SCs) that are either myelinating (mySCs) or nonmyelinating (nmSCs). Our understanding of other cells in the peripheral nervous system (PNS) remains limited. Here, we provide an unbiased single cell transcriptomic characterization of the nondiseased rodent PNS. We identified and independently confirmed markers of previously underappreciated nmSCs and nerve-associated fibroblasts. We also found and characterized two distinct populations of nerve-resident homeostatic myeloid cells that transcriptionally differed from central nervous system microglia. In a model of chronic autoimmune neuritis, homeostatic myeloid cells were outnumbered by infiltrating lymphocytes which modulated the local cell–cell interactome and induced a specific transcriptional response in glia cells. This response was partially shared between the peripheral and central nervous system glia, indicating common immunological features across different parts of the nervous system. Our study thus identifies subtypes and cell-type markers of PNS cells and a partially conserved autoimmune module induced in glia cells.

Results

Single Cell Transcriptomics Dissects the Cellular Composition of Peripheral Nerves. We aimed to better characterize the cellular composition of the PNS. First, we optimized cell extraction from the PNS and achieved highest cell yield and viability by...
combining enzymatic digestion (SI Appendix, Fig. S1A) with myelin depilation and flow cytometry-based sorting for viable cells (Methods) (SI Appendix, Fig. S1 B and C). On average, we thereby obtained 1,520 ± 453 SD viable cells from the combined brachial plexus and sciatic nerves of one mouse. We then pooled cells from multiple mice (n = 12 per batch) and three biological replicates (SI Appendix, Fig. S1D) as input for scRNA-seq. After removal of low-quality cells (Methods), this returned transcriptional information of 5,400 total high-quality PNS cells, with 596 ± 202 SD average genes detected per cell (SI Appendix, Table S1). After normalization (Methods), we identified 12 total PNS cell clusters (Fig. L4).

Overall, 65% of the single cell transcriptomes were assigned to SC and fibroblast cell types, 20% to vascular, and 15% to hematopoietic cell types (Fig. 1B). The unexpectedly high abundance of hematopoietic cells compared to morphological quantification (7) likely reflects their easier extraction and was observed previously (6). One cluster expressed pan-Schwann cell markers (e.g., Erbb3 and Sl100b) and also coexpressed myelin protein genes (e.g., Mbp and Plp1), while another cluster did not express myelin protein genes, but a known SC receptor (Ngrf/p75) (Fig. 1C and SI Appendix, Table S3). We named these clusters mySCs and nmSCs, respectively (Fig. 1A). An additional cluster expressed fibroblast markers (fibro; Ftn1, Pgf1r, Col1a, and Col3a) (8). Detection of marker genes in only a proportion of cells of a cluster (Fig. 1D) is inherent to the method (9). Vectorial clusters expressed canonical markers of vascular smooth muscle cells (vSMCs; Acta2, Tagln, and Tpm2), of pericytes (PCs; Rgs5 and Pdgfrb), of lymphatic endothelial cells (lymph; Lyve1 and Prox1), and of vascular endothelial cells (ECs; Cldn5, Egfl7, and Pecam1) (10) (Fig. 1D and SI Appendix, Table S2). PCs surround endothelial cells in the vessel wall of the microcirculation (11). An additional endothelial cell cluster expressed genes associated with the blood nerve barrier (EC2s; Cldn1 and Scl6a1) (12). Additional transcripts in the EC2 cluster were either novel to endothelial cells (Moxd1 and Nnng1) or had been described in subsets of brain or lung endothelial cells (e.g., Lypd2, Krt19 and Dcun7) although EC2s were transcriptionally distinct from brain/lung ECs (10). We thus identify a unique transcriptional phenotype of a subset of PNS endothelial cells.

Four additional clusters expressed pan-hematopoietic markers (e.g., Pkrm/Cd45) and specifically markers of the myeloid cell lineage (MC; Ly2), macrophages (MP; Cld88), T cells (TCs; Cd3e), and B cells (BCs; Cd79) (Fig. 1D and SI Appendix, Table S2) (9). Blood contamination is unlikely, because mice were intracardially perfused and red blood cells (expressing Hba and Hhb) were absent. We thus define the heterogeneity and provide a census of PNS cells.

To relate our dataset with human diseases, we plotted the average expression of genes associated with hereditary neuropathies against the cell clusters (SI Appendix, Fig. S2A). We found that—except for myelin protein genes—most neuropathy genes were preferentially expressed in non-SC cell types (SI Appendix, Fig. S2A), suggesting a potential relevance of nonglia cell types in inherited PNS disorders. Notably, neuronal cells are not included in our dataset, which may exaggerate the role of these genes in nonglia cells.

**Detailed Transcriptional Characterization of Schwann Cell and Fibroblast Clusters.** Next, we characterized selected clusters in greater detail. The mySC cluster expressed genes encoding PNS myelin proteins (Plp1, Mbp, and Mp2), a key SC lineage transcription factor (Sox10), and other regulators of myelination (e.g., Ptn and Cryab) (13) (Fig. 1 C and D). In addition, genes highly expressed in mySCs were associated with lipid metabolism (e.g., Apoe and Dbi1) (14) and STAT3 (Socs3) (15) signaling pathways (SI Appendix, Tables S2 and S3). Expression of JNK pathway genes (Fos and Junb) (16) in mySCs (SI Appendix, Fig. S3A) has been described (17), but could represent immediate early gene expression resulting from digestion. Gene set enrichment analysis (GSEA) replicated enrichment of signaling pathways (e.g., TGFβ/SMAD) previously associated with Schwann cell function (SI Appendix, Table S3). We next screened for transcripts not previously described in mySCs or in PNS myelination (SI Appendix, Table S4). Such “new-in-mySC” transcripts included metallothioneins (Mt1 and Mt2) and a ferritin chain gene (Fh1) with metal transport and antioxidant functions and unknown relevance in the PNS (18). Also, the transcription factor Btg2 was not previously described in mySCs. We thus identified candidate genes in mySCs (SI Appendix, Fig. S3A).

Next, we further analyzed the nmSC cluster. It highly transcribed a lipoprotein gene (Apol) (Fig. 1D and E and SI Appendix, Table S2), that is known to be expressed in the PNS (19) in Schwann cells (20) with functions in SC-macrophage communication and promoting axonal regeneration (21, 22). The nmSC cluster also expressed ceruloplasmin (Cp) involved in copper metabolism and reported as a potential pan-SC marker (23). When we focused on receptors (Panther class: PC00197) we found that expression of the genes Mactm2, Myoc, Hspg2/Perlecan, Coloa, and Luma2 in nmSCs was in accordance with their known expression and/or function in the PNS (24–27) (SI Appendix, Fig. S3B). Among transcription factors (TFs) (Panther class: PC00218), expression and/or function of Ctf4, Spy2, and Cebpd have been described in SCs (28–30) (SI Appendix, Fig. S3B). Overall, this supports the assignment of the nmSC cluster to the SC lineage.

Coexpression of Ngrf/p75, Csg4/NG2, and Pdgfr/PDGFRβ was previously described in novel pericyte-like cells in the PNS (7). We found coexpression of Ngrf/Csg4/Pdgfr in both the nmSC and PC clusters (SI Appendix, Fig. S3C).

In addition to these known transcripts, we identified a profile of the nmSC cluster that was distinct from mySCs and included specific cell surface molecules (e.g., Coll11 implicated in central nervous system [CNS] myelination) (31), proteases (Mmp2), and the TF Osr2 not previously reported in glia cells (SI Appendix, Fig. S3B and Table S3). Osr2 regulates embryonic mesenchymal cell differentiation (32). GSEA of nmSC marker genes identified pathways related to bone formation (e.g., WP1270 WikiPathway) and neural crest formation (Tcf4 and Sox9) (SI Appendix, Table S6). Notably, some TFs that have been implicated in myelination (Tcf4, Spy2, and Ebf1) (33, 34) were expressed in nmSCs at a higher level than in mySCs (SI Appendix, Fig. S3B).

The fibro cluster expressed a variety of extracellular matrix (ECM) components (Dpt and Gsn) including specific collagen genes (Colla1, Col1a2, Col3a1, and Col14a1) (Fig. 1D and SI Appendix, Table S2). GSEA accordingly identified pathways associated with ECM formation (SI Appendix, Table S7). The cluster also expressed marker genes (Pit1, Clec3b, and Cygb) and TFs (Proxl and Aebp1) (Fig. 1E and SI Appendix, Table S2), that were previously identified in matrix fibroblasts (8). This supports the idea that the fibro cluster represents nerve-associated fibroblasts (3) with a specific matrix fibroblast phenotype. In this cluster we newly identified Sfpr4—a known regulator of the Wnt signaling pathway (35) (Fig. 1D and E and SI Appendix, Table S2)—and multiple members of the IGF signaling pathway (Igfbp6, Igfbp4, and Igfbp5) and a single complement component (C3) that is known to inhibit axonal outgrowth (36). This suggests that nerve-associated fibroblasts could coregulate axonal growth.

In conclusion, we identify a previously unknown transcriptional signature and candidate regulators of nmSCs and nerve-associated fibroblasts.
Fig. 1. Single cell transcriptomics defines cellular phenotypes in the mouse PNS. (A) After multistep purification of peripheral nerve cells, 5,400 total single cell (sc) transcriptomes were generated from adult naive female C57BL/6 mice (three biological replicates, n = 12 mice each, n = 36 total mice), using microfluidics-based scRNA-sequencing. The sc transcriptomes were clustered (Methods) and manually annotated to cell types based on marker gene expression. Each dot indicates one cell and clusters are color coded. (B) The proportion of cells in each cluster is depicted. (C) Feature plots of selected marker genes of the SC lineage. Intensity of red indicates expression level. (D) Dotplot of selected marker genes grouped by cluster. The average gene expression level per cluster is color coded and circle size represents the percentage of cells expressing the gene. Threshold was set to a minimum of 10% of cells in the cluster expressing the gene. (E) Feature plots of genes expressed by the nmSC and fibro clusters.
Confirming Lineage Assignment and Expression of nmSC Markers. We next aimed to confirm and localize cell-type markers combining immunohistochemistry (IHC) and RNA in situ hybridization (ISH). From the top genes identified in the nmSC cell cluster (SI Appendix, Table S2), we selected transcripts with high and specific expression in the nmSC cluster (Apod and Smoc2) (Fig. 1D and E) and stained them using ISH. Mbp was used as positive control of mySCs and showed widespread endoneurial expression (Fig. 2A). As expected, the staining pattern of Mbp protein and Mbp RNA differ (Fig. 2A vs. Fig. 2 C and E). The ISH signal of Apod was located solely endoneurally in either large peri-nuclear aggregates (4.4% of all endoneurial nuclei) or small cytosolic patches (16.9% of all nuclei) (Fig. 2A). Smoc2 was mainly located endoneurally with a similar aggregated morphology (50.9% of all nuclei) (Fig. 2A). Partially epineurial staining of Smoc2 (Fig. 2A) was not reproduced in costainings (Fig. 2B) and is thus likely unspecific. Cells expressing either of the two markers appeared morphologically distinct from mySCs and also did not express Mbp (Fig. 2A and C).

We next aimed to assign the lineage identity of the nmSC cluster. We therefore costained markers of the nmSC cluster (Apod and Smoc2) with three known SC lineage markers (Ngr, S100b, and Sox10) (Fig. 2B and SI Appendix, Figs. S4–S6). As expected for mRNA (Fig. 1C and D), only a proportion of Apod+ and Smoc2+ cells stained positive for Ngr (36.1 ± 9.5%), S100b (57.6 ± 8.5%), and Sox10 (53.9 ± 5.4%) (Fig. 2B and SI Appendix, Fig. S3D). We also used a reporter mouse line to identify the glia cell marker Gfap on protein level (Methods). Both Apod and Smoc2 costained with the four aforementioned lineage markers (Fig. 2B and SI Appendix, Figs. S4–S6), but not with the mySC marker Mbp (Fig. 2C), and also not with fibro markers Vim (SI Appendix, Fig. S7A) and Pdgfra (SI Appendix, Fig. S7B). This supports the idea that Apod/Smoc2-expressing cells in fact represent nmSCs.

Confirming Lineage Assignment and Expression of Fibroblast Markers. We next aimed to confirm and localize selected fibro cluster marker genes (Fig. 1E). RNA ISH of Sfp4 showed expression in some large epineurial cells with patchy cytosolic staining pattern (Fig. 2A). In addition, Sfp4 was expressed by small endoneurial cells (5.7% of all nuclei). This supports the idea that the fibro cluster represents endo- and epineurial fibroblasts.

We then first performed costaining of the fibro cluster markers Sfp4 and Pit1 and found that both transcripts colocalized to individual cells (Fig. 2D and SI Appendix, Fig. S7C). To verify the fibro cluster’s lineage assignment, we combined Pit1 and Sfp4 staining with a reporter mouse (Methods) and detected colocalization of Pit1 and Sfp4 with Pdgfra-driven green fluorescent protein (GFP) in the epineurium (Fig. 2D and SI Appendix, Fig. S8A and B). In contrast, the transcript Sfp4 did not costain with either Sox10 or Mbp (Fig. 2D and E). This supports the idea that the fibro cluster indeed represents nerve-associated fibroblasts and is distinct from nmSCs and mySCs.

PNS Leukocytes Are Distinct and Contain Unique Homeostatic Macrophages. The abundance (15%) of PNS-resident leukocytes in nondiseased mice was surprising and we therefore characterized them in greater detail. Leukocyte clusters separated into T/NK cell (TC), B cell (BC), macrophage (MP), and myeloid cell (MC) lineage clusters (Fig. 3A). The TC cluster (Cd3e and Cd3d) expressed markers of both the helper T cell (Ih7r) and cytotoxic (Cd8a and Cd8b1) subsets (Fig. 3B). T cell subsets and natural killer (NK) cells (Klrk1, Klrk1, and Nkg7) did not separate into subclusters due to their low total cell number (Fig. 3B). The BC cluster (Cd79a and Msa41/Cd20) expressed markers of naive, nonclass-switched B cells (Ighd, negative for: Xbp1, Sdcl1/Cd138) and genes associated with antigen presentation (e.g., H2-Au) (Fig. 3B). Flow cytometry confirmed this overall composition of leukocytes in the murine PNS and indicated the presence of myeloid lineage cells with differing levels of MHC class II expression (two peaks in Fig. 3C). This is in line with two distinct tissue resident macrophage populations that are distinguished based on their MHC class II expression (37).

We next tested whether our findings could be confirmed in humans. We therefore stained sural nerve biopsies of patients without signs of PNS pathology (SI Appendix, Fig. S9A). SOX10 (SI Appendix, Fig. S9B) and MBP protein (SI Appendix, Fig. S9C) served as markers of SCs and mySCs, respectively, while CD34 and ACTA2 are established markers for fibroblasts and vMSC/PCs (SI Appendix, Fig. S9 D and E). Staining for CD45 was used to show the presence of leukocytes (SI Appendix, Fig. S9F), while CD68 specifically stained for macrophages (SI Appendix, Fig. S9G) and CD4/CD8 for T cell subsets (SI Appendix, Fig. S9 H and I). Endoneurial T cells and macrophages were rare, but clearly identifiable, corresponding to the results in naive mice. We were unable to detect BCs in the nondiseased human PNS. The overall composition of endoneurial leukocytes is thus conserved in human PNS.

We next aimed to better understand the potential heterogeneity of myeloid lineage cells in the murine PNS. In our transcriptional data, we identified two myeloid clusters (Ly21 and Ly22) that we named MC and MP (Fig. 3A). The MC cluster expressed markers of nonclassical monocytes (Clec4/CD16) and genes associated with long-lived CNS microglia (Cx3cr1) (38), activation (Csf1r), phagocytosis (Cd300a), and pattern recognition (Clec4e) (SI Appendix, Table S2). The MP cluster (Adgre1/F4/80) expressed markers of classical monocytes (Cd14), nervous system-resident macrophages (Aif1/Iba1), and unique chemokines (e.g., Ccl6, Ccl9, and Ccl4) (SI Appendix, Table S2). In addition, the MP cluster expressed P2y14/Cxcl4 (Fig. 3B) previously described in blood megakaryocytes (9) and some tissue-resident macrophages (39). PNS leukocytes did not resemble megakaryocytes in cytopsins (SI Appendix, Fig. S10) and did not express megakaryocyte markers by flow cytometry (SI Appendix, Fig. S10B). Instead, we found that rare small endoneurial cells expressed P2y14 by ISH (Fig. 3D). Cx3cr1 and P2y14/Cxcl4 thus identify two distinct subsets of nerve-associated myeloid cells, but not megakaryocytes.

Leukocyte Enrichment Confirms Two Distinct Nerve-Associated Macrophage Populations. Low cell numbers prohibited studying nerve-associated leukocytes in greater detail. We therefore used leukocyte enrichment and rats as donor animals (Methods) before subjecting cells to scRNA-seq. We thereby obtained 12,500 total single cell transcriptomes from the rat PNS (SI Appendix, Table S1) out of which 35.3% expressed hematopoietic markers (Ptpre/Cd45) (Fig. 3E). Cell-type identification was again based on marker gene expression (SI Appendix, Table S8). The dataset contained residual nonleukocyte cells which we removed from further analysis (colored in gray in Fig. 3E). Leukocyte transcriptomes separated into TCs (Cd3e and Cd2), BCs (Cd79a and Ighm), and a small cluster of mast cells (Cmna1 and Mep8) (Fig. 3 F and G). The abundant myeloid lineage cells (Lyz2) separated into two apparent clusters (Fig. 3 F and G). One cluster (MPs) expressed features of classical monocytes (Cd14 and Msa47), tissue resident macrophages including markers we had identified in mice (P2y14/Cxcl4 and Adgre1/F4/80), complement components (e.g., C1qg), and specific chemokines (Ccl4, Ccl3, and Ccl2) (SI Appendix, Fig. S10C). The second cluster (MC) expressed high levels of antigen-presenting molecules (e.g., Rtl1-Bb) and alternative trafficking molecules (Ccl17, Ccl6, and Alcam) (SI Appendix, Fig. S10C). Expression of Cx3cr1 was barely detectable (SI Appendix, Fig. S10C). This again supports that the PNS is populated by two subsets of nerve-associated homeostatic myeloid cells.
Fig. 2. Localization and lineage assignment of marker genes in the PNS. (A) Paraformaldehyde (PFA)-fixed cryosections of sciatic nerves of naive adult C57BL/6 mice were stained for Mbp using RNA ISH. Corresponding ISH stainings of Apod, Smoc2, and Sfrp4 are shown with overview (Left) and zoomed (Right) images for each staining. Mbp and Apod were detected with the ViewRNA ISH Tissue Assay Kit (1-plex) (Thermo Fisher). Smoc2 and Sfrp4 were detected with the BaseScope Detection Reagent Kit-RED (ACDbiotech). (B) Fresh-frozen sections of sciatic nerves of naive adult C57BL/6 mice were stained for Apod, Smoc2 together with Schwann cell markers Ngfr, S100b, and Sox10 by the multiplex ViewRNA Cell Assay Kit. SI Appendix, Fig. S1–S3 show additional corresponding stainings. Nerves from Gfap<sup>GFP</sup> mice were costained for Apod and Smoc2 with ISH. (C) PFA-fixed paraffin-embedded sciatic nerves of naive adult C57BL/6 mice were stained for Smoc2 with the BaseScope Detection Reagent Kit-RED, together with an antibody against Mbp. (D) Sections as in B were costained for Sfrp4 and PI16, and for Sfrp4 with Sox10 to show absence of costain. Nerves from PDGFRα<sup>GFP</sup> reporter mice were costained for PI16 by the multiplex ViewRNA Cell Assay Kit. SI Appendix, Fig. S4–S6 show more corresponding stainings. (E) Sections as in C were stained for Sfrp4 with the BaseScope Detection Reagent Kit-RED together with an antibody against Mbp. White dotted line shows the epineurium border of the sciatic nerve. Nuclei were stained with DAPI. (Scale bars: 50 μm, Left; 20 μm, Right; and 10 μm, magnification.) Please note that each dot represents one single RNA molecule. Arrows indicate costaining of all markers, asterisks indicate costain of an identified marker with a known lineage marker, and arrowheads indicate individual staining.
The composition and phenotype of PNS leukocytes is unique and contains specific macrophage populations. (A) The latent space areas of cell clusters identified as hematopoietic cells in Fig. 1 are depicted in a higher magnification UMAP plot. The proportion of leukocyte subsets is depicted. (B) Feature plots of key leukocyte subset markers are depicted. Insets show higher magnification of smaller clusters of interest. Adgre1 encodes F4/80. Intensity of red indicates expression level. (C) Peripheral nerve cells were purified from two female C57BL/6 mice, pooled, and analyzed by flow cytometry after staining for leukocyte markers. The gating strategy is indicated. The proportion of viable CD45⁺CD11b⁺F4/80⁺ macrophages expressing MHC class II was quantified. One representative out of three independent experiments is shown. (D) Paraformaldehyde fixed cryosections of sciatic nerves of naive female adult C57BL/6 mice were stained for F4/80 using RNA ISH. The tissue was stained with the 1-plex ViewRNA ISH Tissue Assay Kit (Thermo Fisher). (Scale bars: 50 μm, Left and 10 μm, Right.) (E) Peripheral nerve cells purified from n = 10 female Lewis rats were enriched for leukocytes using gradient centrifugation (Methods), and processed by scRNA-seq. The resulting 12,500 total rat sc transcriptomes were clustered and the corresponding UMAP plot is shown. Nonhematopoietic cell clusters (Ptprc/CD45 negative) are colored in gray to deemphasize. (F) Feature plots showing selected leukocyte markers in the latent space as in E. Intensity of red indicates expression level. (G) Dotplot of selected marker genes of leukocyte clusters. Average expression level per cluster is color coded and circle size represents the percentage of cells expressing the gene. Threshold was set to a minimum of 10% of cells in the cluster expressing the gene. (H) Sciatic nerves from CX3CR1-GFP reporter mice were processed as in D and stained for Cxcl4, Cxcl6, and DAPI using IHC. (Scale bars: 50 μm, Left and 20 μm, magnification.) SI Appendix, Fig. S11 shows additional corresponding stainings. (I) Sections as in D were stained for F4/80 and Cd169 (Top) or SIGIRR1 and Cd11b (Bottom) using IHC. (Scale bars: 50 μm, Left and 20 μm, magnification.) Arrows indicate costaining of all markers, asterisk indicates costain of the marker of interest with a known myeloid marker, and arrowheads indicate individual staining.

Fig. 3. The composition and phenotype of PNS leukocytes is unique and contains specific macrophage populations. (A) The latent space areas of cell clusters identified as hematopoietic cells in Fig. 1 are depicted in a higher magnification UMAP plot. The proportion of leukocyte subsets is depicted. (B) Feature plots of key leukocyte subset markers are depicted. Insets show higher magnification of smaller clusters of interest. Adgre1 encodes F4/80. Intensity of red indicates expression level. (C) Peripheral nerve cells were purified from two female C57BL/6 mice, pooled, and analyzed by flow cytometry after staining for leukocyte markers. The gating strategy is indicated. The proportion of viable CD45⁺CD11b⁺F4/80⁺ macrophages expressing MHC class II was quantified. One representative out of three independent experiments is shown. (D) Paraformaldehyde fixed cryosections of sciatic nerves of naive female adult C57BL/6 mice were stained for F4/80 using RNA ISH. The tissue was stained with the 1-plex ViewRNA ISH Tissue Assay Kit (Thermo Fisher). (Scale bars: 50 μm, Left and 10 μm, Right.) (E) Peripheral nerve cells purified from n = 10 female Lewis rats were enriched for leukocytes using gradient centrifugation (Methods), and processed by scRNA-seq. The resulting 12,500 total rat sc transcriptomes were clustered and the corresponding UMAP plot is shown. Nonhematopoietic cell clusters (Ptprc/CD45 negative) are colored in gray to deemphasize. (F) Feature plots showing selected leukocyte markers in the latent space as in E. Intensity of red indicates expression level. (G) Dotplot of selected marker genes of leukocyte clusters. Average expression level per cluster is color coded and circle size represents the percentage of cells expressing the gene. Threshold was set to a minimum of 10% of cells in the cluster expressing the gene. (H) Sciatic nerves from CX3CR1-GFP reporter mice were processed as in D and stained for Cxcl4, Cxcl6, and DAPI using IHC. (Scale bars: 50 μm, Left and 20 μm, magnification.) SI Appendix, Fig. S11 shows additional corresponding stainings. (I) Sections as in D were stained for F4/80 and Cd169 (Top) or SIGIRR1 and Cd11b (Bottom) using IHC. (Scale bars: 50 μm, Left and 20 μm, magnification.) Arrows indicate costaining of all markers, asterisk indicates costain of the marker of interest with a known myeloid marker, and arrowheads indicate individual staining.

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The Composition, Origin, and Phenotype of Nerve-Associated Myeloid Cells Is Unique. We next studied nerve-associated myeloid lineage cells using IHC and reporter mice. One myeloid lineage cluster (named MC) expressed Cx3cr1 in mice. To confirm this population, we used CX3CR1-GFP reporter mice. We found that Cx3cr1-driven GFP was expressed by a proportion of endoneurial mononuclear cells (Fig. 3H and SI Appendix, Fig. S11). This population coexpressed Cd68, while it did not coexpress for Cxcl4 (encoded by Pf4) by IHC. This indicated that one of two distinct subsets of nerve-associated macrophages can be identified by Cx3cr1 and absence of Cxcl4 (Fig. 3H and SI Appendix, Fig. S11). Conversely, we found that another population of endoneurial cells coexpressed the Cxcl4 protein (encoded by Pf4; MP cluster) and the myeloid marker Cd68 (Fig. 3H and SI Appendix, Fig. S11). Cxcl4-positive cells were located in close proximity to F4/80+ and Cd169+ macrophages, but did not coexpress these markers (Fig. 3I, Top). When we stained for more specific markers, we found that Cxcl4-expressing cells were positive for Cd11b and Siglr1; both have been associated with pathogen recognition and phagocytosis in macrophages (40) (Fig. 3I, Bottom). This suggests that the Cxcl4-expressing MP cluster represents phagocytosing nerve-associated macrophages. Notably, Cxcl4-expressing macrophages were recently identified in CNS-associated border compartments of healthy mice (41). Overall, we thus identified the transcriptional profile of two subsets of nerve-associated myeloid cells characterized by expression of Cx3cr1 (MC cluster) and Pf4/Cxcl4 (MP cluster), respectively.

We next initially addressed the ontogenetic origin of nerve-associated myeloid cells. In an Flt3Cre-driven switch-reporter mouse, cells originating from yolk sac hematopoiesis (e.g., microglia) express tdTomato (tdT), while hematopoietic lineages derived from fetal liver and bone marrow express GFP (42) (SI Appendix, Fig. S124). As expected, myeloid cells in the brain were mostly tdT+ (89.75%; i.e., yolk sac-derived microglia) and in other organs were >80% GFP+ (bone marrow 87.1% and spleen 81.9%; SI Appendix, Fig. S12B). In contrast, the PNS contained ~35% tdT+ myeloid cells (36.75%; SI Appendix, Fig. S12 C and D) suggesting a partially late embryonic or bone marrow hematopoietic stem cell (HSC)-derived origin and supporting the heterogeneity of PNS myeloid cells.

Autoimmunity Induces Specific Compositional and Phenotypic Changes in PNS Cells. We next aimed to understand how autoimmunity shapes the composition and phenotype of PNS cells. We therefore extracted nerve cells from a mouse model of spontaneous chronic peripheral neuritis (43). We thereby generated single cell transcriptomes from PNS cells of young and clinically unaffected ICAM-1−/−/nonobese diabetic (NOD) mice (5,250 cells, n = 12 female mice) that histologically did not show PNS inflammation (SI Appendix, Fig. S134) and of prediabetic NOD control mice (5,400 cells, n = 24 female mice). Although cell-type clustering clearly reidentified the PNS cell clusters we had identified in healthy mice (Fig. 4A and B and SI Appendix, Table S9), ICAM-1−/−/NOD nerves contained grossly expanded leukocyte clusters while parenchymal cells and myeloid lineage cells (MC/MP clusters) were underrepresented in comparison to NOD control mice (Fig. 4A and B). Due to their underrepresentation, the transcriptionally similar vSMC/PC and the MC/MP clusters no longer separated into distinct clusters, while plasmacytoid dendritic cells (pDCs) were identified in both genotypes (Fig. 4A and B) resulting in 11 total cell clusters. The presence of pDCs was not detected in other datasets and may thus be specific for the NOD genetic background. The expanded clusters were mainly CD4-expressing T cells with a memory phenotype that previously has been described for tissue-resident memory TCs (CD4; Vps37b, Ccr6, Tnfaii3, and Rora) (44), and BCS with an activated phenotype (H2-DMβ2/MHC class II, Ms4a1/Cd20, Cd83, and Cd74). In addition, ICAM-1−/−/NOD nerves contained expanded clusters of cytotoxic CD8 TGs (CD8; Cdb8a, Krc1, and Nkg7), and pDCs (Flt3 and Siglech) (SI Appendix, Fig. S13B and Table S9). Myeloid lineage cells (Adgre1/F4/80 and Pf4/Cxcl4) were not numerically expanded, but showed a more activated phenotype with increased expression of Cx3cr1 and costimulatory molecules like Cd86 (MC cluster) (SI Appendix, Fig. S13B and Table S9). We confirmed this unique phenotype of PNS-infiltrating cells including the presence of pDCs by flow cytometry (Fig. 4C). PNS autoimmunity in ICAM-1−/−/NOD thus induces a local accumulation of specific leukocyte populations including memory and cytotoxic T cells and pDCs, while homeostatic nerve-associated macrophages are underrepresented.

We next aimed to construct an integrated view of how nonimmune cell types respond to autoimmune tissue destruction. We first focused on vascular cells and identified a set of differentially expressed (DE) genes in vascular endothelial cells in ICAM-1−/−/NOD compared to NOD control cells (Fig. 4D and SI Appendix, Table S10). DE genes were characterized by the induction of a chemokine (Ccx5), antigen-presenting molecules (B2m and H2-D1), and signs of cellular stress (mt-CO2) (SI Appendix, Table S10). The neutris-specific transcriptome in the mmSC (Fig. 4E and SI Appendix, Table S11) and mySC (Fig. 4F and SI Appendix, Table S12) clusters also featured an up-regulation of antigen-presenting molecules (H2-K1, H2-D1, and B2m) and immunoproteasome subunits (Pmnb8 and Pmnb10) in line with the known conditional antigen-presenting function of Schwann cells (45) (SI Appendix, Fig. S13C). Both mmSCs and mySCs also down-regulated some of their cluster-defining transcripts, including collagen (ColIa1) (Fig. 4E) and myelin proteins (Pmp22 and Myp2) (Fig. 4F), respectively. In contrast, mySCs from ICAM-1−/−/NOD mice up-regulated Sostdc1 and Zeb2 (SI Appendix, Table S12)—two transcripts previously identified to coregulate posttraumatic Schwann cell differentiation and nerve regeneration (46, 47). In summary, autoimmunity causes induction of immune-related transcripts, loss of structural parenchymal components, and signs of cellular dedifferentiation in the PNS.

We next analyzed whether the response of PNS tissue to autoimmunity was similar or distinct from the CNS. We therefore compared genes DE in the mySC cluster under neuritis conditions with genes DE in oligodendrocytes (oligos; myelin-glia of the CNS) in the experimental autoimmune encephalomyelitis (EAE) animal model of human multiple sclerosis (MS) (48). Using a similar threshold for significance (Methods), we found that 32 (15.4%) of 208 genes DE in oligos were also DE in EAE in NOD nerves (Fig. 4G and SI Appendix, Table S13). Notably, 30 (94%) of these 32 genes were canonical IFN-response genes (SI Appendix, Fig. S14). The transcriptional response of myelinating glia to autoimmunity is thus partially conserved between oligos and mySCs and mimics an IFN-dependent antiviral response.

Autoimmunity Diversifies Cell–Cell Communication in the Peripheral Nerve. We next aimed to identify mechanisms controlling intercellular communication in the PNS. We therefore systematically predicted intercellular signaling by adapting a recently described human tool (49) for mouse data (Methods). In these predicted networks, cell types are defined as nodes (circles) and ligand-receptor pairings predicted from expression data are defined as directional edges (arrows) (SI Appendix, Fig. S15 A and C). The network of nondiseased NOD nerves showed high connectivity of leukocyte clusters (MCS, BCS, pDCs) and of the fibro and mmSC clusters (SI Appendix, Fig. S15 A and B) as illustrated by the high number of edges (SI Appendix, Table S14). Multiple interactions were directed from EC1 to fibro and mmSC clusters (SI Appendix, Fig. S15 A and B). Fewer edges originated...
Fig. 4. Neuritis induces influx of lymphocytes and a partially shared autoimmunity module in myelinating Schwann cells. (A) Peripheral nerve cells were purified from female prediabetic NOD control mice (Left, n = 24 mice, two biological replicates) and ICAM-1−/−NOD mice (Right, n = 12 mice, one biological replicate) and processed by scRNA-seq. The resulting NOD control (n = 5,400) and ICAM-1−/−NOD (n = 5,250) sc transcriptomes were clustered and are shown in UMAP plots. (B) The percentage of cells in each cluster in NOD control and ICAM-1−/−NOD samples is depicted in a dotplot with circle size representing proportion of cells in each cluster. (C) Peripheral nerve cells were purified from presymptomatic female ICAM-1−/−NOD mice (n = 5) and characterized by flow cytometry. The proportion CD8+ cytotoxic T cells (Top Right) and NK1.1−/−NCR1−/− NK cells (Top Middle) was quantified. B220+ B cells (Lower Middle Left), B220+Ly6C−/−Ccr9−/−CD137− plasmacytoid dendritic cells (Lower Middle Right), and CD11b+F4/80+CD14+ macrophages (Lower Far Right) was quantified. One out of two independent experiments is shown. (D–F) Volcano plots depicting DE genes between ICAM-1−/−NOD vs. NOD control samples in endothelial cells (EC1) (D), nmSCs (E), and mySCs (F). Only DE genes with an average (avg) log fold change (FC) > ±0.25 are plotted. Genes with an avg log FC above ±0.5 and P values <0.001 are marked in red and the gene names are provided. The y axes represent the negative log10 of the adjusted P value. (G) Venn diagram showing genes DE in ICAM-1−/−NOD vs. NOD mice within mySC compared with DE genes within oligodendrocytes (oligo) in EAE vs. control samples in an available dataset (48). EC1: endothelial cells cluster 1, EC2: endothelial cells cluster 2, TC(CD4): CD4 T helper cells, TC(CD8): cytotoxic CD8 T cells and natural killer cells, NS: not significant, P: adjusted P value, pct: percentage expressed.
from the mySC cluster, suggesting low intercellular connectivity of mySCs in the uninflamed peripheral nerve.

We also applied the network analysis to neuritis and—as expected—found that leukocyte clusters increased in size and the total number of intercellular interactions increased in ICAM-1−/−NOD nerves (edge count in SI Appendix, Fig. S15 C and D and Table S14). Surprisingly, the “betweenness centrality” as a measure of the amount of control that one node exerts over the interactions of other nodes (49), did not increase for most leukocyte clusters (MC, pDC, BC), but instead changed for fibro and vSMC/PC clusters (SI Appendix, Table S14). This suggests that fibro and vSMC/PC clusters gain more control over local cell–cell communication in autoimmunity. Comparative network analysis may thus help to prioritize candidate signaling pathways in disease.

Discussion

In this study, we generated an unbiased map of the cellular composition and transcriptional phenotype of PNS cells in health and autoimmunity. We identified and confirmed markers of nonmyelinating SCs and nerve-associated fibroblasts. Such cell-type-specific markers will likely facilitate better characterization of these cell types. We also found an unexpectedly diverse and unique leukocyte repertoire in the PNS with two subsets of nerve-associated homeostatic myeloid cells identified by specific gene sets. And chronic autoimmune neuritis induced an expansion of lymphocytes that altered the local cell signaling circuitry. Notably, the transcriptional tissue response to autoimmunity was partially shared between peripheral and central nervous system glia cells and resembled an IFN response. This suggests a degree of stereotypicity in the response of glia cells to autoimmunity.

We used a specific extraction protocol that was balanced for optimum cell viability at maximum yield and allows identifying all expected PNS cell types. This approach is useful to screen for subsets of PNS cells, but does not allow quantifying the absolute and relative cell abundance in the PNS as observed previously (6) in contrast to morphological approaches (7). For example, SCs are likely more difficult to extract than leukocytes and therefore seem underrepresented.

We found considerable transcriptional differences between mySCs and nmSCs. Although nmSCs clearly expressed SC lineage markers by transcriptomics, ISH, and IHC (Figs. 1D and 2B and C), this cluster expressed multiple transcripts (e.g., Ccl11), including a specific transcription factor (Osr2) that had not been previously associated with PNS glia cells. Interestingly, we also identified a number of genes that were previously associated with myelination, but showed the highest expression in nmSCs rather than mySCs (41, 60) in PNS-derived macrophages (41, 60), ı̇n PNS-derived macrophages (41, 60) and Table S2). Therefore, nerve-associated macrophages are likely transcriptionally distinct from microglia. In contrast to the known yolk-sac origin of microglia, nerve-associated macrophages also seem to have a mixed developmental origin with broad, slowly-derived hematopoietic progenitors partially contributing to this population. In conclusion, our study opens diverse avenues for better understanding the PNS.

Methods

Animals. C57BL/6J, Icam1tm1grepNOD (named ICAM-1−/− NOD for simplicity), NOD/ShiLtJ, CX3CR1-GFP, HgfAP-GFP, Pdgfrα-Egfp, and Flt3Cre-mt/mG mice and Lewis rats were used. Mice were without signs of neuropathy (ICAM-1−/− NOD) or diabetes (NOD/ShiLtJ).

Cell Extraction and Purification. Sciatic nerves and the brachial nerve plexus were dissected from intracardially phosphate buffered saline (PBS)-perfused animals and finely chopped. Enzymatic digestion was optimized (SI Appendix, SI Material and Methods) and adapted from a previous study (61). Myelin was depleted using anti-myelin beads (Miltenyi Biotec). Single cells were subsequently sorted (BD FACSAria III) for intact viable cells using three viability markers: Zombi NIR APC C17, Calcein-AM FITC, and DAPI (Biolegend) (SI Appendix, Fig. S1).

Single Cell RNA-Sequencing and Analysis. Single cell RNA-sequencing was performed using the Chromium Single Cell 3′ Kit with v2 chemistry (10x Genomics) according to manufacturer instructions. Sequencing was done either on a local Illumina Nextseq500 High-Out 75 Cycle Kit with a 26-8-0-57 read setup or commercially on a NovaSeq6000 (300 Cycle Kit) with paired end 150 read setup. Details are provided in SI Appendix, Table S1.

Processing of raw sequencing data was performed with the cellranger pipeline v3.0.2. Subsequent analysis steps were carried out with the R-package Seurat v3.0.0 (62) using R v3.6.0 as recommended. Unique molecular identifier (UMI) data were normalized using an approach with regularized negative binomial regression (63). Dimensionality reduction was done by uniform manifold approximation and projection (UMAP) with default parameters. DE genes were identified with “FindMarkers” function in Seurat. The threshold was set to 0.25 average log fold change and using Wilcoxon rank sum test, unless otherwise stated. To annotate the clusters, genes differentially expressed in a one vs. all cluster comparison were queried for known expression in a literature search and plotted in feature plots.

DE genes were identified between ICAM-1−/−NOD and NOD conditions after alignment using Harmony (64). Interactions between the cells were predicted using CellPhoneDB (49) with normalized and filtered scRNA-seq and conservation of murine ensemble IDs to human using biomartR. Statistical significance of the cellular interactions were calculated as described (49). Interactions between clusters were visualized in a heatmap and clustered with complete linkage and Euclidean distance measure using the R package pheatmap. Network visualization was performed with Cytoscape v.3.7.1 and GSEA with the Enrichr tool.

Comparison with Published Datasets. We compared DE genes in ICAM-1−/−NOD vs. NOD control mice in specific clusters with a published dataset of DE genes in EAE vs. control mice (48). To improve comparability, DE genes in our dataset were identified using MAST instead of Wilcoxon rank sum test with a lower average log fold change of 0.095. All DE genes with an adjusted P value greater than 0.05 were removed. The top up- and downregulated DE genes of the Falcao dataset (48) (cutoff gene expression >4) were compared with our top DE genes using the Venn-Diagram package (SI Appendix, Table S13). Intersecting genes were then analyzed using the “Interferome” database.

Immunohistochemistry. Fixed frozen slides were used for histology-based methods. Slides were stained with CD68, F4/80, Cxcl4, Cd169, Cd11b, and Sign1. Secondary Alexa Fluor (AF)-conjugated antibodies were used. Slides were mounted in Fluoromount G with DAPI (InVitrogen). Images were taken using a three laser fluorescence microscope (Bioerevo BZ-900 microscope with BZII Viewer software, Keyence) and processed in ImageJ.
Five total human samples were selected according to lack of pathological findings in sural nerve biopsies. The study received ethical approval by the ethical committee of the University Clinic Leipzig, Germany. The study was performed by the automated immunostainer Benchmark XT (Roche). The following antibodies were used: CD45, CD68, CD8, CD4, CD34, SMA/ACTA2, SOX10, and MBP.

RNA in situ hybridization. RNAISH was performed on fixed/fresh frozen and paraffin-embedded sections of sciatic nerves from intracardial PBS-perfused C57Bl/6 mice, and POFP-GFP/GFP mice. Three different ISH kits were used according to manufacturer protocol. The thermo Fisher ViewRNA ISH Tissue Assay Kit (1-plex) was used to detect Mm-Mbp, Mm-Apod, Mm-Smoc2, Mm-Sfrp4, and Mm-Pigf as single stains. The thermo Fisher ViewRNA Cell Assay Kit (multiplex) was used, in combination with the first steps of the previously mentioned tissue assay kit, to detect Mm-Apod, Mm-Smoc2, Mm-Nifg, Mm-S100b, Mm-Sox10, Mm-Sfrp4, and Mm-Pigf in different single and costain settings. The ACDBio BaseScope Detection Kit ACDbio BaseScope Detection Red (1-plex) was used to detect Mm-Smoc2 and Mm-Sfrp4 in costain with antibodies for Mbp. All images were obtained with an Axio Observer Z1 (Zeiss) and processed in AxioVision and ImageJ.

Flow Cytometry of Leukocytes. Flow cytometry analysis was performed on isolated PNS cells. The following viability dye and murine antibodies were used: Zombie NIR, CD45, CD11b, B220, CD3, CD4, CD8, NK1.1, NK2G2B4, NKp46, F4/80, CD14, Ly6c, D317, CCR9, CD11c, and MHCI. Samples were measured on the Gallios (10 colors, 3 lasers, Beckman Coulter) and analyzed with FlowJo V10.

Data Availability. The raw scRNA-seq data supporting the findings in this study have been deposited in the GEO repository with the accession code GSE142541, available at https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE142541. All processed scRNA-seq data are included as Datasets S1–S14. Additional figures supporting the main text, are provided in SI Appendix.

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