An environment-dependent transcriptional network specifies human microglia identity

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INTRODUCTION: Microglia play essential roles in central nervous system homeostasis and influence diverse aspects of neuronal function, including refinement of synaptic networks and elaboration of neuromodulatory factors for memory and motor learning. Many lines of evidence indicate that dysregulation of microglial functions contributes to the pathogenesis of neurodegenerative diseases, including Alzheimer’s disease and Parkinson’s disease. Emerging evidence from mouse and human studies also suggests that microglia influence neurodevelopmental and psychiatric disorders such as schizophrenia and depression. Most disease risk alleles associated with neurodegenerative diseases reside in noncoding regions of the genome, requiring the delineation of functional genomic elements in the relevant human cell types to establish mechanisms of causation. The recent observation that mouse brain environment strongly influences microglia-specific gene expression has implications for understanding pathogenic responses of microglia in diseases and disorders and modeling their phenotypes in vitro.

RATIONALE: Although dysregulation of microglial activity is genetically linked to neurodegenerative diseases and psychiatric disorders, no systematic evaluations of human microglia gene expression or regulatory landscapes are currently available. In addition, the extent to which mice provide suitable models for human microglia is unclear. The major goals of this study were to define the transcriptomes and DNA regulatory elements of human microglia ex vivo and in vitro in comparison to the mouse and to systematically relate these features to expression of genes associated with genome-wide association study (GWAS) risk alleles or exhibiting altered expression in neurodegenerative diseases and psychiatric disorders.

RESULTS: We used RNA sequencing, chromatin immunoprecipitation sequencing, and assay for transposase-accessible chromatin sequencing to characterize the transcriptomes and epigenetic landscapes of human microglia isolated from surgically resected brain tissue in excess of that needed for diagnosis. Although some effects of underlying disease cannot be excluded, the overall pattern of gene expression was markedly consistent. Microglia-enriched genes were found to overlap significantly with genes exhibiting altered expression in neurodegenerate diseases and psychiatric disorders and with genes associated with a wide spectrum of disease-specific risk alleles. Human microglia gene expression was well correlated with mouse microglia gene expression, but numerous species-specific differences were also observed that included genes linked to human disease. More than half of the genes associated with noncoding GWAS risk alleles for Alzheimer’s disease are preferentially expressed in microglia. DNA recognition motifs enriched at active enhancers and expression of the corresponding lineage-determining transcription factors were very similar for human and mouse microglia. Transition of human and mouse microglia from the brain to tissue culture revealed remodeling of their respective enhancer landscapes and extensive down-regulation of genes that are induced in primitive mouse macrophages following migration into the fetal brain. Treatment of microglia in vitro with transforming growth factor β1 (TGF-β1) had relatively modest effects in maintaining the ex vivo pattern of gene expression. A significant subset of the genes up- or down-regulated in vitro exhibited altered expression in neurodegenerative diseases and psychiatric disorders.

CONCLUSION: These studies identify core features of human microglial transcriptomes and epigenetic landscapes. Intersection of the microglia-specific gene signature with GWAS and transcriptomic data supports roles of microglia as both responders and contributors to disease phenotypes. The identification of an environment-sensitive program of gene expression and corresponding regulatory elements enables inference of a conserved and dynamic transcription factor network that maintains microglia identity and function. The combinations of signaling factors in the brain necessary to maintain microglia phenotypes remain largely unknown. In concert, these findings will inform efforts to generate microglia-like cells in simple and complex culture systems and understand gene-environment interactions that influence homeostatic and pathogenic functions of microglia in the human brain.

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Microglia are the resident macrophage population of the central nervous system (CNS) that perform functions required for typical development, as well as homeostasis, plasticity, immunity, and repair (1–5). Dysregulation of microglia functions contributes to the pathogenesis of neurodegenerative diseases, including Alzheimer’s disease (AD) (6–8), Parkinson’s disease (PD) (9–12), and Huntington’s disease (HD) (13–15). In addition, microglia may play roles in neurodevelopmental and psychiatric diseases such as autism, schizophrenia (Scz), and depression (16, 17). Collectively, these observations provide a compelling incentive to define the mechanisms that control microglia functions and to investigate the potential to alter these functions for therapeutic purposes.

Microglia possess distinct ontogenic, morphologic, gene expression patterns, and functional characteristics and account for 5 to 20% of all glial cells, depending on brain region. Studies in mice indicate that microglia are derived from yolk sac progenitor cells that populate the brain early in development (18, 19). Microglia share many traits with other subsets of tissue-resident macrophages, including dependence on the CSF1 receptor for differentiation and survival (18, 20), a requirement for PU.1 as an essential lineage-receptor for differentiation and survival (18, 20), a determination of transcription factor (LDTF) (21–23), highly efficient phagocytic and tissue debris clearance abilities (2, 5), and the ability to quickly trigger an inflammatory response following detection of pathogens or tissue damage (23).

Microglia also express hundreds of genes at higher levels than other macrophage subsets (24–26) that may be influenced by factors present in the CNS microenvironment (25, 27). Selecting mechanisms by which the brain environment influences the activity and functions of one of its primary support cells. As microglia participate in the developmental refinement of synaptic networks (28) and elaborate neuromodulatory factors for adult motor learning (29), alterations in the brain microenvironment may lead to pathogenic programs of gene expression in microglia.

Insights into mechanisms that control the development and function of specific cell types can be obtained by analyses of their enhancer repertoires and corresponding epigenetic landscapes (30). Microglial enhancers are primed by relatively simple combinations of LDTFs (31, 32) to become sites of activity of broadly expressed signal-dependent transcription factors (SDTFs). These enhancers thereby integrate cellular signaling pathways with DNA. Within cell-specific enhancers, DNA sequences provide information about the identities of key LDTFs and the activity states of signaling pathways that control the functions of SDTFs (31, 33–35). Furthermore, the locations and sequence content of enhancers provide insights into how noncoding genetic variation may affect gene expression and cellular phenotypes (32, 36–38).

Human microglia transcriptomes

Microglia were isolated from brain tissue resected for treatment of epilepsy, brain tumors, or acute ischemic events in 19 individuals. Patient characteristics, medications, and assays performed for each sample are provided in table S1. For patients with epilepsy, the resected tissue provided did not include the brain area displaying the strongest epileptogenic activity. Although most specimens from individuals with epilepsy exhibited some degree of cortical dysplasia, three samples were determined to be histologically normal. For patients with brain tumors (8), samples were considered to be tumor negative by pathological examination. The two brain samples resected to treat acute ischemic injuries were from individuals who had no prior evidence of brain pathology. As judged by neuropathological diagnostic evaluation, none of the samples displayed evidence of pathological microgliosis (i.e., loss of tiled distribution, process retraction, and enlarged cell bodies) despite the occasional presence of reactive cells (fig. S1).

Tissues were immediately placed on ice in the operating room and processed for microglia isolation within a few hours of surgical resection (25). A microglia-rich cell fraction was isolated by Percoll gradient centrifugation and further purified by fluorescence-activated cell sorting. Cell purity was determined by gating on live-CD11b+CD45low/CD64+CX3CR1high single cells (fig. S2A). This strategy excluded cells expressing moderate to high levels of CD45, which were present in variable amounts depending on the specific sample (fig. S2A). Microglia were then triaged to specific analysis workflows depending on yields and depth of existing data sets (table S1). For five individuals, RNA sequencing (RNA-seq) was also performed from a portion of the intact cortical brain tissue used to isolate microglia (hereafter referred to as “cortex”). Isolated microglia exhibited a high degree of viability and RNA integrity. In one case, brain tissue was simultaneously obtained from the occipital and parietal lobes of the same individual, enabling independent isolation of microglia from each region. RNA expression levels for these two samples exhibited a Pearson correlation coefficient of 0.929 (fig. S3A). In a second case, an individual underwent sequential surgical resections approximately 1 year apart. The RNA expression levels in microglia obtained from this individual at these two time points exhibited a Pearson correlation coefficient of 0.905 (fig. S3B). On the basis of these observations, the variance in gene expression noted in these samples likely overestimates...
the variance in the isolation of microglia and implies a high degree of precision in the RNA-seq data set. Comparisons of the RNA-seq profiles of any two individuals for genes expressed >2 log2 transcripts per million (TPM) yielded Pearson correlation coefficients of ~0.8, as exemplified by a comparison of microglia isolated from a 1-year-old female patient diagnosed with atypical rhabdoid tumor to microglia isolated from a 7-year-old female patient with epilepsy (fig. S3C). When all transcripts were considered (i.e., including genes expressed at ≤2 log2 TPM), Pearson correlation coefficients were in the range of 0.94 to 0.98 (fig. S7D), indicating no substantial outliers among samples. RNA-seq data indicated low-to-absent levels of neuronal, astrocyte, oligodendrocyte, and endothelial-specific genes, confirming insignificant levels of contaminating cells (fig. S2B and table S2).

Hierarchical clustering of expressed genes from all microglia samples in comparison to cortex is illustrated in Fig. 1A. Although there are likely to be alterations in gene expression related to disease, no strong patterns emerged that were related to diagnosis or age (Fig. 1A), perhaps in part to exclusion of most reactive cells at the time of sorting (fig. S2A). A small set of genes exhibited significant sex-specific differences in gene expression, and these were mostly related to X- and Y-linked genes (fig. S4). Relative expression values for the 30 most highly expressed genes across each of the patients are illustrated in Fig. 1B, indicating substantial individual variation for some transcripts. This analysis confirmed that human microglia expressed very high levels of numerous genes associated with regulation of microglia ramification and motility.
including the fractalkine receptor CX3CR1, purinergic receptor P2RY2, and genes involved in synaptic remodeling, such as complement factors C3, C1QA, C1QB, and C1QC (table S2).

Nearly all studies of differentially regulated genes in human neurodegenerative or behavioral disorders are based on RNA profiling of intact tissues. As gene expression profiles for individual cell types in the human brain are poorly characterized, it is difficult to assign cell types responsible for disease-related changes. Using a stringent cutoff of 10-fold–increased expression relative to cortex tissue at a false discovery rate (FDR) of 0.05, we defined a microglial gene signature in the human brain comprising 881 transcripts (Fig. 1A and table S2). We then intersected this gene list with 46 publicly available data sets for differentially regulated genes in neurodegenerative and behavioral disorders derived from microarray or RNA-seq analysis of intact tissue (39–53). Two-sided Fisher’s exact tests revealed significant enrichment or depletion of microglia signature genes in 28 of the 46 data sets (table S3). For example, hundreds of genes in the microglia gene signature exhibited significant overlaps with sets of genes that were positively correlated with Braak stage in AD, were up-regulated in frontotemporal lobar degeneration (FTLD) in frontal cortex, were up-regulated in HIV-associated neurocognitive disorder (HAND), or were up-regulated in Scz (Fig. 1C). Notably, different subsets of the microglia signature gene set were up-regulated or down-regulated in cortex in PD (Fig. 1C).

Characterization of the microglia transcriptome, in conjunction with parallel whole-brain...
cortical gene expression profiling, also enabled evaluation of the relative expression levels of genes associated with AD, PD, multiple sclerosis (MS), and Scz risk alleles (54). Many of these genes were preferentially expressed in microglia, with the relative proportion varying by disorder (Fig. 1D and fig. S5). With respect to AD, for example, expression for 28 of 48 AD genes was higher in microglia, including TREM2, SORL1, INPP5D, MEF2C, and CD33 (Fig. 1E). For MS, 48 of 81 genes associated with risk alleles were preferentially expressed in microglia (fig. S5). By contrast, approximately one-third of the genes associated with PD and Scz risk alleles exhibited preferential expression in microglia (fig. S5). As the risk alleles for these disorders are largely nonoverlapping, these findings suggest different roles for microglia in each disorder context.

The extent to which findings in mice and humans are similar can vary depending on the system evaluated (55). We compared average transcript levels of human microglia mRNAs with orthologous mRNAs from microglia isolated from 7- to 10-week-old male C57BL/6 mice using an identical protocol (25) (Fig. 2A, fig. S6, and tables S4 and S5). The microglia transcriptome comparison between the two species revealed overall similarities in patterns of gene expression (Pearson’s r = 0.806), with the majority of orthologous genes pairs (13,253 of 15,788) expressed within a fourfold range. Nonetheless, this analysis also revealed genes with species-specific bias in expression magnitude. At a cutoff of 10-fold difference and an FDR of 0.05, 400 orthologous mRNAs were preferentially expressed in human microglia and 290 mRNAs were more highly expressed in mouse microglia (Fig. 2A).
Notable examples included higher expression in human microglia of regulators of the complement system (e.g., C2, C3, VSIG4, SERPING1) and genes involved in brain structure development, including SYNDIG1, GLDN, CTNNBP2, and ROBO3. By contrast, genes more highly expressed in mouse microglia included Hexp, Sparc, and Sall1. Of the subset of these genes that also exhibited microglia-specific expression, a similar species-specific pattern of expression was observed in the intact cortical tissue used for microglia isolation, indicating that these differences were not a consequence of the isolation procedure (fig. S7).

We compared the levels of mRNA transcripts in C57BL/6 microglia with different tissue-resident macrophage populations (26, 27) and observed 900 mRNAs with levels of expression in microglia that were 10-fold greater than the integrated mean of expression levels in the other macrophage subsets at a FDR of 0.05 (Fig. 2B). This definition captures a mouse microglia gene signature set that is larger but mostly inclusive of previously described microglia signature sets (27, 57, 58). Gene ontology analysis of this extended microglia gene signature returned significant functional enrichment for CNS-related categories such as regulation of neurogenesis, synaptic signaling, CNS development, response to transforming growth factor beta (TGF-β), and regulation of neurotransmitter transport (Fig. 2C). RNA-seq data sets are not available for a sufficient number of human resident macrophage populations to generate a corresponding human microglia signature gene list. However, the relative expression levels of the human and mouse orthologous genes corresponding to the mouse microglia signature genes revealed that ~53% (477 of 900) of the genes that were highly specific to microglia in the mouse were expressed at comparable levels in human microglia (Fig. 2D).

Comparing the expression of genes linked to human neurodegenerative and behavioral disorders, of the 39 AD-associated and expressed genes, 11 were higher in human microglia whereas three were increased in mouse microglia (Fig. 2E). Notably, of the top 25 most expressed, two genes, APOC1 and MPZL1, were essentially not expressed in C57BL/6 microglia; the expression of a third one, SORL1, was nearly 20 times lower in mice than in humans (Fig. 2F). PD-associated genes also displayed similar human versus mouse distributions (fig. S6). PD-associated genes SNC2, LRRK2, CD46, and CXCL12 were expressed to a greater extent in human microglia than in mouse microglia. Corresponding comparisons for all expressed genes linked to AD, PD, MS, and SzC are illustrated in fig. S8. Collectively, these results highlight human versus mouse similarities and differences in the expression profile of core microglia genes.

**Enhancer landscape of human microglia**

We performed assay for transposase-accessible chromatin–sequencing (ATAC-seq) to define open regions of chromatin in and around transcription start sites (fig. S1). ChIP-seq and ATAC-seq data were highly correlated across human individuals and mouse biological replicates (fig. S9). A representative University of California Santa Cruz (UCSC) genome browser track for human microglia data in the vicinity of the BIN1 gene is illustrated in Fig. 3A, and for CX3CR1 and TREM2 in fig. S10A. rs6733839 is a noncoding risk allele for AD (29) that lies within an enhancer-like region ~25 kb upstream of the BIN1 transcriptional start site, suggesting regulatory function. De novo motif analysis of ATAC-seq peaks associated with H3K4me2 was performed to identify the most highly enriched transcription factor recognition motifs associated with poised or active chromatin. These analyses returned very similar motifs for both mouse and human microglia. A dominant signature was observed for PU.1 in both species, with highly significant enrichment also observed for motifs assigned to CTCF, IRF, RUNX, MEF2, C/EBP, AP-1, SMAD, and MAF factors (Fig. 3B and fig. S10B).

Expression levels of transcription factors recognizing the motifs identified in Fig. 3B are illustrated in Fig. 3C. SPI1 (encoding PU.1) was the most highly expressed ETS domain transcription factor, and the top motifs were nearly perfect matches for its consensus recognition sequence. The assignment of the second-most enriched motif to CTCF was unexpected because CTCF is ubiquitously expressed and is thought to primarily play roles in the organization of broad domains of chromatin architecture. However, the identified motif was a near-perfect match for the CTCF consensus recognition site, and CTCF was the only transcription factor expressed in microglia known to recognize this motif. By contrast, the remaining motifs were clearly recognized by multiple related members of transcription factor families that were reliably coexpressed in microglia.

We used H3K27ac as a means of identifying super-enhancers (SEs), which are regions of the genome exhibiting a disproportionately high density of active regulatory marks and transcription factor binding (60–62). Common SEs were associated with genes that are known to play essential roles in the development and function of virtually all macrophages, whereas mouse microglia-specific SEs were associated with genes known to play restricted or selected roles in this cell type (25). Corresponding analysis of the H3K27ac-high regions in human microglia identified ~700 SEs that were also associated with active regions of the genome overlapping or in close proximity to genes highly expressed in specific to microglia, including CX3CR1, P2RY12, and TMEM119 (table S6). The intersection of transcription factor genes associated with SEs in human and mouse microglia captured many of the transcription factors illustrated in Fig. 3C. An additional 35 factors were associated with SEs, some of which have been linked to functional roles in microglia, including SALL1, STAT3, and RELA (Fig. 3D). These factors also exhibited similar expression levels in mouse and human microglia.

Evaluation of mRNAs annotated as encoding orthologous transcription factors indicated similar expression levels in human and mouse for the great majority. However, using a 10-fold cutoff at an FDR of 0.05, 20 factors were preferentially expressed in human microglia, including CIITA, RUNX2, EGR3, and NR4A2. By contrast, 16 factors were preferentially expressed in mouse microglia, including Sall4 and Smart1 (Fig. 3E).

To establish the relationship between enriched motifs and actual transcription factor binding, we performed ChIP-seq for PU.1 in microglia isolated from six individuals. These experiments yielded a highly consistent pattern of genomic binding (Fig. 3A and fig. S10A), with PU.1 being localized at 51.2% of ATAC-seq–defined regions of open chromatin associated with H3K4me2 and at 60.9% of regions exhibiting these features plus H3K27ac. De novo motif enrichment analysis identified the PU.1 motif as the top hit, followed by motifs for IRF, AP-1, MEF2, C/EBP, and RUNX (Fig. 2B and fig. S10B). Similar results were observed for Pu.1 binding sites in mouse microglia. The frequency distribution of many of these motifs is consistent with the corresponding transcription factors forming ternary complexes with PU.1 (e.g., the IRF motif; fig. S10C) or functioning as collaborative binding partners (e.g., CTCF, AP-1, and RUNX motifs; fig. S10C).

We also investigated the relationships between PU.1 binding, open chromatin, and H3K27ac for genes that are highly differentially expressed between human and mouse microglia. H3K27ac at promoters was strongly correlated with differential gene expression (Fig. 3F). Differential gene expression and H3K27ac were also associated with species-specific PU.1 binding and ATAC-seq signal exemplified by the SALL1, SPARC, VSIG4, and APOC1 loci (fig. S11). Collectively, these findings provide evidence that the transcriptional mechanisms that control gene expression are largely conserved between human and mouse microglia and that differences in gene expression are primarily driven by species-specific organization of cis-regulatory elements at target genes. However, as a small number of transcription factors thought to play important roles in microglia are divergently expressed, such as CIITA and RUNX2 proteins, there is also the potential for trans effects at the subsets of genes regulated by these factors.

**Alterations of the human microglia transcriptome in vitro**

Tissue environment has emerged as an important determinant of microglia identity, but many studies of microglia are performed in vitro. Transcriptional analysis of mouse microglia transferred to a tissue culture environment for 7 days documented significant changes in expression of hundreds of genes, with a preferential reduction in expression of microglia-specific genes (25). Similar to what was observed with mouse microglia, transfer of human microglia to the in vitro environment resulted in significant changes in the expression of thousands of genes. Responses of
human microglia isolated from different individuals to the in vitro environment were concordant (Fig. 4A, iv). For the individuals with both ex vivo and 7-day in vitro microglia populations obtained, more than 300 genes were induced greater than 10-fold and more than 700 genes that were repressed more than 10-fold in vitro (FDR 0.05). At a twofold cutoff, 2263 genes were up-regulated in vitro and 3702 genes were down-regulated at a FDR of 0.05. Thirty-three percent of the conserved signature gene set and 31% of the genes associated with AD risk variants exhibited >2-fold reduction in expression in vitro (Figs. 4, B to D). A similar proportion (30%) of genes associated with PD also displayed such reduction in vitro, whereas more limited changes for genes associated with MS and Scz were observed (fig. S12).

We performed RNA-seq in human microglia at 6, 24, and 168 hours following transfer to tissue culture. Highly dynamic responses were observed beginning at 6 hours and lasting throughout the time course (Fig. 4D). Cultured microglia remained positive for Iba1 staining, whereas no expression of markers for astrocytes or neurons was observed, indicating that changes in RNA expression were not due to contaminating cells (fig. S13A). A total of 1957 mRNAs were induced more than fourfold by 6 hours after plating. Gene ontology analysis of highly induced genes indicated strong enrichment for terms related to inflammation and stress responses (fig. S13B, C). Notably, more than 2000 genes were down-regulated more than fourfold at 6 hours after plating. Gene ontology analysis of highly induced genes indicated strong enrichment for terms related to inflammation and stress responses (fig. S13A, C). Notably, more than 2000 genes were down-regulated more than fourfold at 6 hours after plating.

An environment-dependent transcriptional network

The perturbation in gene expression resulting from the transition to an in vitro environment provided an opportunity to gain insight into the transcriptional network controlling human microglia identity. Although SPI1 increased slightly, most of the mRNAs encoding transcription factors that recognize enriched motifs in microglia enhancers exhibited significant reductions in vitro. In particular, AP-1 and MEF2 family members, as well as MAF, RUNX, and SMAD3, were highly sensitive to the environmental perturbation (Fig. 5A). Similarly, the majority of transcription factors associated with SREs were also down-regulated (Fig. 5B).

**Fig. 4. Influence of tissue culture environment on microglia gene expression.** (A) Comparison of gene expression of ex vivo microglia from two individuals (panel i), effect of transition to an in vitro environment in the presence of IL-34 for 7 days on microglia from each individual (panels ii and iii), and comparison of in vitro gene expression profiles (panel iv). (B) Scatter plot illustrating effects of culture environment on the conserved microglial gene signature (red) in human microglia. (C) Pie chart of the genes associated with AD risk variants that are expressed in ex vivo or in vitro microglia above a log2 (TPM +1) value of 2. Genes more highly expressed ex vivo are in blue, and genes more highly expressed in vitro are in purple. (D) Combined bar graphs and dot plots of the 21 most expressed genes, indicating effects of in vitro culture on expression of genes associated with AD risk alleles. Red stars indicate significant differential expression (>2-fold, FDR 0.05). (E) Heat map illustrating changes in microglia gene expression 6 hours, 24 hours, and 7 days after transfer to culture environment.
We performed ATAC-seq and ChIP-seq for H3K27ac in microglia placed into culture for 7 days (Fig. 5C). Overall, of the 31,164 ATAC-seq peaks associated with at least 16 normalized H3K27ac tags (i.e., high confidence sites), ~14% exhibited a loss of H3K27ac and ~20% a gain of H3K27ac (Fig. 5D), indicating remodeling of the regulatory landscape in vitro. Extending this analysis, 424 of the 740 SEs defined ex vivo were lost and 541 new SEs were observed (Fig. 5E). De novo motif analysis of the ATAC-seq peaks associated with a >2-fold decrease in H3K27ac returned recognition elements for PU.1, IRF, CTCF, AP-1, SMAD, and MEF2 motifs as the most significantly enriched (Fig. 5F), consistent with PU.1 establishing microglia enhancers and reduced expression and/or activity of many of the collaborative and signal-dependent factors that recognize these motifs (Fig. 5A). Notably, loss of the SMAD motif is consistent with reduced TGF-β signaling in the in vitro environment.

We compared the expression of the mouse orthologs of transcription factors with those present in primitive yolk sac macrophages and their progeny in developing and adult mouse brain (63). A subset of mRNAs encoding these factors was expressed in primitive macrophages and maintained high levels of expression in brain microglia (Fig. 6A). By contrast, an alternative subset of factors that recognize DNA motifs enriched in microglia enhancers and/or were preferentially expressed in adult human and mouse microglia exhibited increases in expression from the embryonic to the adult stage of brain development (Fig. 6A). Notably, a majority of the adult microglia transcription factors that exhibited reduced expression after transfer to the in vitro environment were induced following migration of primitive macrophages into the developing brain, consistent with their induction by local environmental factors.

We compared the set of genes robustly up-regulated during brain development in mouse microglia to those consistently down-regulated upon transition to the in vitro environment. A highly significant overlap was observed: Of 3517 genes that increased during development, 1459 intersected with the 2398 genes “down” in culture ($P < 1E-308$; Fig. 6B). This finding illustrates that a substantial component of the microglia unique signature that emerges during development is down-regulated in vitro. The correlation between genes up-regulated in fetal microglia compared to yolk sac macrophages and genes down-regulated upon transfer of adult microglia to an in vitro environment provided evidence that the changes observed in

![Fig. 5. Environment-dependent enhancer landscapes.](http://science.sciencemag.org/)
vivo result from loss of signals that are part of the brain environment. We cultured mouse microglia in vitro for 7 days and treated them with TGF-β1 cytokine (Fig. 6C and fig. S15), which is necessary for maintenance of the in vivo mouse microglia phenotype (27, 64). Of the 2398 genes down-regulated >2-fold in vitro, TGF-β1 induced expression >2-fold of 429, including Sall1, Tmem119, and P2ry12. Corresponding studies of TGF-β1 treatment of human microglia in vitro yielded similar results, although variation between individuals limited the number of genes achieving statistical significance after correction for multiple testing. Nonetheless, SALL1 was among the genes exhibiting significant up-regulation (fig. S15B). Because of the relatively modest effects of TGF-β1, we evaluated several other tissue culture conditions to investigate whether any might be more effective in maintaining an ex vivo gene expression signature. Although most treatment conditions produced some effect on gene expression, none resulted in preservation of the ex vivo molecular phenotype (fig. S16). It has not yet been possible to maintain primary human microglia in a serum-free environment. Recent studies have derived microglia-like cells from human induced pluripotent stem cells in a defined media that more efficiently expressed the microglia molecular signature. Therefore, we focused our attention on the tissue culture conditions that yielded the most consistent gene expression changes after 7 days in vitro. Although the TGF-β1-treated cultures resulted in the greatest number of genes with altered expression, all tissue culture conditions resulted in altered expression.

**Fig. 6. A network of developmentally programmed and environment-dependent transcription factors establishes microglia identity and functions.**

(A) Heat map displaying changes in mRNA expression of key microglia transcription factors as a function of development of the mouse brain. (B) Venn diagram illustrating significant overlap of mouse microglia genes repressed (>2-fold, FDR 0.05) in culture and genes that increased (>2-fold, FDR 0.05) in expression during brain development. P-value of the overlap is provided (Fisher’s exact test). (C) Venn diagram illustrating significant overlap of mouse microglia genes repressed (>2-fold, FDR 0.05) in culture after 7 days and genes maintained or induced (>2-fold, FDR 0.05) by chronic stimulation with TGF-β1. P-value of the overlap is provided (Fisher’s exact test). (D) Pie chart of microglia-enriched genes compared to brain cortex overlapping with differentially expressed genes in neurodegenerative or behavioral diseases from Fig. 1C color-coded according to whether they are increased, decreased, or unchanged after transfer to a tissue culture environment. (E) Diagram illustrating main features of transcription factor network regulating microglia cell identity and functions.
closely resembles cerebrospinal fluid (65). However, these cells exhibited a transcriptomic phenotype that was much more similar to human microglia cultured in vitro than to ex vivo microglia (fig. S17).

Lastly, we investigated whether genes expressed at least 10-fold higher in ex vivo microglia compared to brain cortex tissue and that are associated with differentially regulated genes in neurodegenerative or behavioral disorders (Fig. 1C) were sensitive to the environment. Of the nonredundant set of 291 microglia transcripts associated with differentially regulated genes in AD, PD, HD, SCZ, or FTD, 117 were down-regulated in vitro whereas 29 were up-regulated in vitro (Fig. 6D). These observations provide evidence that differential expression of microglia-enriched genes in these disorders is at least partially due to changes in brain environment.

Discussion

The transcriptomes and epigenomic features of human microglia show that disease status, age, sex, and treatment history had a relatively minor influence on the overall pattern of gene expression. It is thus likely that the composite transcriptomic and epigenomic profiles identified largely capture a representative human microglial gene signature and its underlying network of genomic regulatory elements. We find that mouse and human microglia transcriptomes are largely conserved, but they also exhibit large differences in the expression of several hundred genes that appear to be primarily driven by cis-regulatory differences.

Genetic evidence suggests that microglia dysregulation is a core feature of numerous chronic neurodegenerative and psychiatric diseases. To this end, our profiling of the human microglia transcriptome enabled us to estimate a weighted influence of microglia on the development of different brain disorders. In addition, by defining a microglial gene signature in the human brain, it is possible to link changes in gene expression in neurodegenerative and behavioral disorders to alterations in the number and/or function of microglia. The observation that the microglial gene signature overlaps with genes that are both up- and down-regulated in PD provides strong evidence that differential gene expression at the whole-tissue level reflects changes in microglia gene expression and not simply changes in the microglial population number. This interpretation is further reinforced by the observation that many of the microglia-enriched genes that change in the context of neurodegenerative and behavioral disorders also show significant changes in expression upon transfer to an in vitro environment. Although we also observed substantial individual variation in genes linked to disease risk, more individuals will be required to power analyses linking genetic variation to gene expression.

Examining differences in ex vivo and in vitro microglia, we observe rapid and profound alterations in gene expression by 6 hours, characterized by activation of genes associated with acute inflammatory responses and decreased expression of genes associated with diverse immune functions. The incomplete recovery of down-regulated genes at 7 days was at least in part due to the absence of brain-specific signals. At present, the spectrum of brain-specific signals that are necessary to maintain microglia phenotypes remains largely unknown, but this finding highlights the limitations of investigating microglia biology in a tissue culture setting. Notably, current efforts to reprogram various starting cell populations to microglia-like cells largely achieve an in vitro phenotype. It is likely that more complex culture conditions, such as co-culture with neurons and astrocytes in three-dimensional or organoid contexts, will be required to more precisely model the in vivo phenotype. Integration of these findings with recent studies of yolk sac and embryonic microglia (50, 63) suggests a model by which intersecting developmentally determined and environment-dependent transcriptional networks establish microglia identity and function (Fig. 6E). We speculate that regulatory elements are targets of environmental perturbations associated with disease and are also important sites of action of natural genetic variation.

In sum, our work demonstrates that human and mouse microglia display relatively well-conserved transcriptomic and epigenomic phenotypes, and it provides a critical resource that will allow for a better understanding of mouse models of human brain disorders, guide reprogramming efforts for generation of human microglia in vitro, and enable a better interpretation of the functions of DNA variants associated with disorders of the central nervous system.

Methods

Human tissue

Microglia were isolated from brain tissue in excess of that needed for pathological diagnosis (37) and for treatment of epilepsy, brain tumors, or acute ischemia in 19 individuals. For patients with epilepsy (9), the resected tissue displayed epileptogenic activity. However, the epileptic focus exhibiting the strongest epileptogenic activity was not provided and was separated at the time of resection for pathological analysis. For patients with brain tumors (8), samples consisted of incidentally resected brain tissue, i.e., brain tissue resected to gain access to the tumor and considered to be tumor negative by the neurosurgeon intraoperatively. Two brain samples were resected to treat acute ischemic injuries from individuals who had no prior evidence of brain pathology. Brain tissue was obtained with informed consent under a protocol approved by the UC San Diego and Rady Children’s Hospital Institutional Review Board. Resected brain tissue was immediately placed on ice and transferred to the laboratory for microglia isolation within 3 hours after resection.

Human and mouse microglia isolation

Resected human brain tissues and mouse brains were homogenized by gentle mechanical dissociation, as performed in Gosselin et al. (25). Microglia were then enriched by Percoll gradient, washed, and stained with cell surface markers for final purification by flow cytometry, using a BD Influx cell sorter. Human microglia were defined as live/DAPT CD11b+CD45+CD64+ CX3CR1hi single cells. C57BL/6 mouse microglia were defined as live/DAPT CD11b+CD45+ single cells.

Primary culture of isolated microglia

Isolated microglia were maintained in culture for 6 hours, 24 hours, or 7 days. Culture media consisted of Dulbecco’s modified Eagle medium (DMEM)/F12 (Life Technologies, 113300-032) supplemented with 5% fetal bovine serum (heat inactivated, Omega Scientific, FB-12) and 1× Anti-Anti (Life Technologies, 152240-062). No media changes were performed for the duration of the culture experiments. CSF1R ligand Interleukin-34 (R&D, human 5265-IL-010; mouse 5195-ML-010) was supplemented daily at a concentration of 20 ng/ml. For experiments investigating the beta1 on microglia gene expression in culture, recombinant human or mouse TGF-beta1 cytokine (R&D, human 240-B-002; mouse 7666-MB-005) was added daily at a concentration of 10 ng/ml.

RNA-seq library preparation: Isolation and fragmentation of poly(A) RNA and synthesis of cDNA

Isolated microglia were pelleted and put into 150 μl of lysis/Oligo d(T) Magnetic Beads binding buffer and stored at −80°C until processing. For human brain cortical RNA, 500 ng of RNA was diluted with proper volume of 2× lysis/Oligo d(T) Magnetic Beads binding buffer to a final concentration of 20 ng/ml. For experiments investigating the effects of TGF-beta1 on microglia gene expression in culture, recombinant human or mouse TGF-beta1 cytokine (R&D, human 240-B-002; mouse 7666-MB-005) was added daily at a concentration of 10 ng/ml.

Chromatin immunoprecipitation

For ChIP-seq experiments, a cross-linking step with paraformaldehyde was performed immediately after staining but before microglia sorting. Chromatin immunoprecipitation for histone modification H3K4me2, H3K27ac, and transcription factor PU.1 was performed as described in Gosselin et al. (25).

RNA-seq and ChIP-seq final library preparation and sequencing

Sequencing libraries were prepared from recovered DNA (ChIP) or generated cDNA (RNA) by blunt-end, A-tailing, and adapter ligation as previously described in Heinz et al. (31). Prior to final polymerase chain reaction amplification, RNA-seq libraries were digested by 30 min of incubation at 37°C with Uracil DNA Glycosylase. RNA-seq and ChIP-seq libraries were single-end sequenced for 51 cycles on an Illumina HiSeq Gosselin et al., Science 356, eaal3222 (2017) 23 June 2017
**Data analysis**

**Preprocessing**

FASTQ files from sequencing experiments were mapped to either the UCSC genome build hg38 (for human) or mm10 (for mouse). STAR with default parameters was used to map RNA-seq experiments. Bowtie2 with default parameters was used to map ATAC-seq and ChIP-seq experiments. HOMER was used to convert aligned reads into "tag directories" for further analysis.

**RNA-seq**

The "analyzeRepeats" function of HOMER was used to generate a table of read counts for each experiment using the parameters "-raw-count exons -strand both -condenseGenes" and a table of transcripts per million (TPM) values using "-tpm-normMatrix 1e7 -count exons -strand both -condenseGenes." The base-2 logarithm of the TPM values was taken after adding a pseudo-count of 1 TPM to each gene. Within-species differential gene expression analysis was performed using the R package llimma with the voom normalization. Mapping of orthologous genes between species was done using the one-to-one orthologs from the Ensembl (version #84) Compare database.

**ChIP-seq**

Tag directories for the ChIP-seq experiments were combined into ex vivo and in vitro pools for each target in both mouse and human. The tag directories for the input DNA were likewise combined. Peaks were then called on the pooled tags with the pooled input DNA as background using HOMER’s “findPeaks” command with the following parameters for the PU1 ChIP-seq: "-style factor -size 200 -mindist 200"; and the following parameters for the H3K27ac and H3K4me2 ChIP-seq: "-style histone -size 500 -mindist 1000 -region." The "tpm" parameter was set to the number of replicates in the pool.

**Motif enrichment**

Peaks were considered overlapping if their coordinates overlapped by at least one base pair. De novo motif enrichment analysis was performed on ATAC-seq and PU1 ChIP-seq peaks that overlapped either H3K4me2 (ATAC-seq) or H3K27ac (PU1), using HOMER’s "findMotifsGenome" command with the following parameters: "-size given -len 14,12,10,8 -mask."
References


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SUPPLEMENTARY MATERIALS

www.sciencemag.org/content/356/6344/eaal3222/suppl/DC1 Materials and Methods

Figs. S1 to S7

Tables S1 to S6

References (66–76)

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10.1126/science.aal3222
An environment-dependent transcriptional network specifies human microglia identity


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Of mice and men's microglia

Microglia are immune system cells that function in protecting and maintaining the brain. Gosselin et al. examined the epigenetics and RNA transcripts from single microglial cells and observed consistent profiles among samples despite differences in age, sex, and diagnosis. Mouse and human microglia demonstrated similar microglia-specific gene expression profiles, as well as a shared environmental response among microglia collected either immediately after surgery (ex vivo) or after culturing (in vitro). Interestingly, those genes exhibiting differences in expression between humans and mice or after culturing were often implicated in neurodegenerative diseases.

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