Sibling chimerism among microglia in marmosets

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Abstract

Chimerism happens rarely among most mammals but is common in marmosets and tamarins, a result of fraternal twin or triplet birth patterns in which *in utero* connected circulatory systems (through which stem cells transit) lead to persistent blood chimerism (12-80%) throughout life. The presence of Y-chromosome DNA sequences in other organs of female marmosets has long suggested that chimerism might also affect these organs. However, a longstanding question is whether this chimerism is driven by blood-derived cells or involves contributions from other cell types. To address this question, we analyzed single-cell RNA-seq data from blood, liver, kidney and multiple brain regions across a number of marmosets, using transcribed single nucleotide polymorphisms (SNPs) to identify cells with the sibling's genome in various cell types within these tissues. Sibling-derived chimerism in all tissues arose entirely from cells of hematopoietic origin (i.e., myeloid and lymphoid lineages). In brain tissue this was reflected as sibling-derived chimerism among microglia (20-52%) and macrophages (18-64%) but not among other resident cell types (i.e., neurons, glia or ependymal cells). The percentage of microglia that were sibling-derived showed significant variation across brain regions, even within individual animals, likely reflecting distinct responses by siblings' microglia to local recruitment or proliferation cues or, potentially, distinct clonal expansion histories in different brain areas. In the animals and tissues we analyzed, microglial gene expression profiles bore a much stronger relationship to local/host context than to sibling genetic differences. Naturally occurring marmoset chimerism will provide new ways to understand the effects of genes, mutations and brain contexts on microglial biology and to distinguish between effects of microglia and other cell types on brain phenotypes.

Introduction

Chimerism, in which an organism contains cells from genetically distinct animals, happens rarely among mammals. Chimerism is common, however, among marmosets and their close relatives the tamarins: in these primate species, animals usually give birth to dizygotic twins or trizygotic triplets whose blood contains cells from siblings. During development, the siblings share circulation *in utero*, allowing the exchange of hematopoietic stem cells (Gengozian et al., 1969; Wislocki, 1939). Most marmosets then exhibit blood chimerism throughout life: their blood-derived DNA is a mixture of both twins' genomes, with the twin's genome contributing 12% to 80% of the DNA in the blood (Niblack et al., 1977; The Marmoset Genome Sequencing and Analysis Consortium, 2014). This indicates that the twins' hematopoietic stem cells establish permanent residency in one another's bodies and contribute to blood cell populations throughout life.

A longstanding mystery involves whether other tissues and organisms also harbor chimerism. Beyond the blood, Y-chromosome DNA has been detected in the brain and other organs of female marmosets with male twins (Ross et al., 2007; Sweeney et al., 2012), eliciting much speculation about how chimerism might have shaped behavior and natural selection in marmosets. However, it is still not known what cell types harbor this sibling DNA; such observations could in principle be explained by the presence of blood cells within these organs.

Here, we analyze chimerism in the marmoset brain, liver, kidney and blood, using single-nucleus RNA-seq (snRNA-seq) to infer cell types and using combinations of transcribed SNPs (visible in the snRNA-seq data) to determine which marmoset sibling is the source of each cell. This approach makes it possible to determine whether chimerism arises from blood cells, resident immune cells, or other cell types, and to explore what chimerism can teach us about cellular migrations and population dynamics.

Results

Marmoset chimerism can be characterized at single-cell resolution

To identify which individual cells have the genome of the host marmoset, and which have the genome of the host's birth sibling, we used combinations of many transcribed SNPs that were visible in the snRNA-seq data for each nucleus (Wells et al., 2023).

We first determined whether marmosets have sufficient sequence variation to enable the distinction between host and sibling cells. From whole-genome sequences of 123 marmosets, we identified 13 million polymorphic bi-allelic SNPs in the marmoset genome, with individual marmosets harboring 2.3 to 3.8 million (average 3.4 million) heterozygous sites – comparable to levels of heterozygosity in humans. For sibling comparisons, we detected a large number of sites at which any two siblings' genomes differed, ranging from 2.0 to 3.7 million sites (average 2.9 million sites) across 96 sibling comparisons (**Fig. 1A**). To determine how many of these sites were visible in snRNA-seq data, we analyzed snRNA-seq data for several marmoset tissues. The results varied by cell type, reflecting that different cell types' nuclei harbored different quantities of RNA. The hundreds of transcribed variant sites that differed between siblings (median>311 per nucleus) suggested ample power to distinguish between siblings in all cell types (Wells et al., 2023) (**Fig. 1B**).

We next evaluated whether the genome variation visible in snRNA-seq reads was sufficient to distinguish between host and sibling cells. For this we used Dropulation, which identifies the

96 donors of individual cells (from a set of genome-sequenced candidate donors) by using
97 combinations of the transcribed SNPs visible on the snRNA-seq reads of the individual cells
98 (Wells et al., 2023). We first analyzed the blood cells by snRNA-seq of a marmoset (CJ028) born
99 with two birth siblings and used Dropulation (Wells et al., 2023) to assign individual cells to the
100 correct sibling (**Fig. 1C**). The relative likelihoods of the siblings could be strongly differentiated
101 (relative likelihoods of 10^3 to 10^{23}) for >99% of the nuclei (**Fig. 1C**). We found a high level of
102 chimerism: 84% of all nuclei sampled in this marmoset's blood appeared to contain the genome
103 of one (67% - sibling #1) or the other (17% - sibling #2) of its two birth siblings (**Fig. 1C**), consistent
104 with the wide range of chimerism found in previous studies: 4-82% in marmoset T cells and B
105 cells (Niblack et al., 1977); 13-37% in marmoset whole blood ("The Marmoset Genome
106 Sequencing and Analysis Consortium", 2014).
107
108 **Apparent liver and kidney chimerism arises from infiltrating monocytes**
109
110 Earlier studies have identified Y-chromosome-derived DNA sequences in the organs of female
111 marmosets with male birth siblings, suggesting that these organs harbor chimerism (Sweeney et
112 al., 2012). However, such observations could also in principle arise from blood or from blood-
113 derived immune cells that are present in those organs (Sweeney et al., 2012).
114
115 We performed snRNA-seq analysis of the blood (1,741 nuclei), liver (10,877 nuclei) and kidney
116 (9,262 nuclei) of a marmoset (CJ026) with one birth sibling (**Fig. 1D**). The snRNA-seq profiles
117 clustered into groups that were readily recognized (based on the RNAs expressed) as the
118 principal cell types of each organ; we determined the identity of each cluster using scType, a cell-
119 type identification tool that uses a database of known marker genes (Ianevski et al., 2022).
120
121 In kidney and liver, the only clearly twin-derived cells were cells of hematopoietic origin: the
122 resident macrophages in liver (Kupffer cells), lymphocytes in liver, and lymphocytes in kidney
123 (**Fig. 1E,F**). All non-hematopoietic cell types in liver and kidney appeared to contain only the host
124 marmoset's own genome. Chimerism levels for the two chimeric liver immune cell types appeared
125 to diverge, with sibling-derived cells accounting for 15% of Kupffer cells and just 4% of
126 lymphocytes (5/122 vs 57/383; Chi-square test *P*-value=0.003). In the blood, chimerism levels
127 varied across the various cell types: the most abundant cell types, the Naive B cells and Naive
128 CD8+ T cells, were respectively 29% and 32% sibling-derived, while the less-abundant CD8+
129 NKT-like cells were 15% sibling-derived (**Fig. 1F**, Chi-square test *P*-value=0.01).
130
131 These results indicate that, in this animal's liver and kidney, apparent DNA chimerism likely arose
132 from infiltrating immune cells rather than other cell types. These results also indicate that cells
133 with siblings' genomes can differ in their tendency to acquire specific hematopoietic cell fates and
134 in their tendency to infiltrate into organs.
135
136 **Marmoset brain microglia and macrophages exhibit abundant chimerism**
137
138 To characterize chimerism in the marmoset brain, we utilized a large snRNA-seq data set being
139 generated for a marmoset brain cell atlas (Krienen et al., 2022). We first analyzed 497,000
140 single-nucleus RNA-expression profiles from the neocortex, thalamus, striatum, hippocampus,
141 basal forebrain, hypothalamus and amygdala of an adult marmoset with two birth siblings
142 (marmoset CJ028). We clustered the cell types using gene expression similarities (**Fig. 2A**) and
143 identified brain cell types as in earlier work (Krienen et al., 2020), identifying neurons,
144 astrocytes, oligodendrocytes, ependymal cells, endothelial cells, microglia and macrophages
145 (**Fig. 2B**). Microglia (which expressed markers *TREM2*, *LAPTM5* and *C3*) and macrophages
146 (which expressed *LYVE1* and *F13A1*) were a small fraction of all nuclei analyzed (about 3.6%),

but due to the large number of nuclei we profiled (53 brain tissue dissections, 497,000 nuclei), we were able to ascertain sufficient numbers of microglia (18,175 nuclei) and to a lesser extent macrophages (172 nuclei) for many downstream analyses. We found microglia and macrophages in snRNA-seq data from 10 additional marmosets with different genetic backgrounds from 3 different colonies (sample information for 11 marmosets in **Supplementary Table 1**; a total of eleven marmoset brains were analyzed and all are unrelated except for CJ006 and CJ007 which are birth siblings, and CJ025 and CJ026 which are (non-birth) siblings; only CJ026 was assessed for liver and kidney, and 3 marmosets were assessed for blood: CJ026, CJ027 and CJ028). Brain snRNA-seq of all 11 marmosets showed consistently the presence of these two myeloid cell types in the brain (**Supplementary Fig. 1**; number of microglia and macrophages in **Supplementary Table 2**).

Donor-of-origin analysis of snRNA-seq data from 2.2 million nuclei sampled from 137 brain tissue samples from these 11 marmosets showed a clear and consistent pattern: microglia and macrophages, but not neurons, glia or endothelial cells, harbored chimerism (**Fig. 2C**, **Supplementary Fig. 2**). Microglia exhibited abundant chimerism – across the 11 marmosets, the total fraction of cells with the sibling's genome ranged from 20% to 52% (for triplets, sum of two siblings; **Fig. 2D**). Macrophages exhibited a similarly wide range of sibling fractions across marmosets (18% to 64%, **Fig. 2D**).

The quantitative extent of microglial chimerism varied across individuals (**Supplementary Fig. 3A**; test of heterogeneity *P*-value<2.2×10^{-16}), as did that of macrophage chimerism (**Supplementary Fig. 3B**; test of heterogeneity *P*-value=1×10^{-4}). We asked whether microglial and macrophage chimerism were correlated. Intriguingly, only a modest correlation of chimerism levels across 14 host-sibling pairs was observed between the microglia and macrophages (**Fig. 2E**; Pearson correlation 0.31), suggesting that in addition to the cell's genome, other factors such as local host environment play a role in differential recruitment or survival of sibling cells. Neither of these two myeloid cell types showed consistently higher chimerism than the other cell type did (**Fig. 2D**).

Sibling contributions in blood vs. brain

Though microglia (like macrophages) are myeloid cells that derive from hematopoietic stem cells, the ontogenies of microglia and brain macrophages are distinct from those of bone-marrow-derived peripheral blood mononuclear cells (Perdiguero & Geissmann, 2016). As such, differences in the developmental and migration histories of these cell populations could in principle have caused their chimerism fractions to diverge in a systematic way.

We analyzed three marmosets for which snRNA-seq was performed on both blood and brain tissues. Sibling contributions to microglia and brain macrophages were in general quite different from those in blood (**Fig. 3**).

Marmoset CJ028's chimerism (involving two birth siblings) provided a setting in which cells with three different genomes shared the same environment through development until adulthood (to two years of age). Among CJ028's microglia, the fraction of cells from sibling 1 (35%) was greater than that from sibling 2 (13%) (two-sided test of proportionality *P*-value<2.2^{-16}), while in blood, the opposite was true (fraction of cells from sibling 1 across all blood cell types was 18%, fraction of cells from sibling 2 across all blood cell types was 67%; two-sided test of proportionality *P*-value<2.2^{-16}) (**Fig. 3D,E**, **Supplementary Table 3**).

198 **Microglia chimerism fraction varies across brain regions**
199
200 Sibling contributions to the microglial population could in principle be shaped by effects that are
201 local to specific brain areas, including differential response of sibling microglia to local
202 recruitment or proliferation cues. To evaluate whether the sibling contribution to the microglial
203 population varied across brain areas within individual marmosets, we performed chimerism
204 analysis for each of the brain regions profiled in the snRNA-seq datasets: neocortex, thalamus,
205 striatum, hippocampus, basal forebrain, hypothalamus and amygdala (Krienen et al., 2020,
206 2022). Within each marmoset, the fraction of microglia with a sibling's genome diverged across
207 a marmoset's brain regions (**Fig. 4A**). For example, in marmoset CJ025, sibling contributions to
208 microglial populations ranged from 11% (21/193) in the thalamus to 56% (174/310) in the
209 striatum (*P*-value=1.1×10^{-23}, Chi-square test of thalamus vs striatum; *P*-value=1.5×10^{-40}, Chi-
210 square test across all 4 brain regions). For marmoset brains profiled with at least 300,000
211 nuclei, CJ027, CJ028 and CJ029, tests of heterogeneity *P*-values were even more significant:
212 8.8×10^{-83}, $<1 \times 10^{-300}$, and 6.1×10^{-47}, respectively. Analysis of finer brain substructures showed a
213 similar result **(Fig. 4B**). None of the brain regions exhibited consistently higher or lower
214 chimerism levels, suggesting that these divergences did not result from differential physical
215 access of host and sibling microglia to different brain areas (**Fig. 4**).
216
217 **Gene-expression comparisons of host- to sibling-derived microglia**
218
219 Chimerism provides the unusual opportunity to compare cells with different genomes in a
220 shared *in vivo* biological context. We compared RNA expression between host- and sibling-
221 derived microglia of a female marmoset with two birth siblings. Sex differences among the
222 siblings (the host (CJ028) was a female and one of the two siblings was a male) allowed a
223 natural control: the *XIST* gene encodes a non-coding RNA involved in silencing one copy of
224 chromosome X in females and thus exhibits sex-specific expression due to cell-autonomous
225 mechanisms. We found that (as expected) *XIST* transcripts were detected at far higher levels in
226 the snRNA-seq profiles of microglia with the female twin's genome relative to the male twin's
227 (**Fig. 5A,C**). By contrast, *XIST* transcripts were detected at similar levels in two microglial
228 populations with the genomes of female twins (**Fig. 5B**).
229
230 Gene-expression differences between host- and sibling-derived microglia in the same brain
231 could in principle arise from asymmetries in their developmental histories (which would be
232 shared across host animals) or from genomic differences (which would vary from host animal to
233 host animal). In all eleven individual marmosets, analysis identified anywhere from six to
234 hundreds of genes whose differential expression distinguished microglia with host vs. sibling
235 genomes, but aside from sex differences (*XIST* gene) we did not find any gene that consistently
236 distinguished host from sibling microglia across the sibling comparisons (**Supplementary Fig.**
237 **4**, **Fig. 5A-C**).
238
239 **Brain context vs. genetic differences as determinants of microglial gene expression**
240
241 Chimerism presents interesting opportunities to distinguish between cell-autonomous and
242 contextual effects on biology, and to compare the magnitudes of such effects.
243
244 We first considered the difference in contexts provided by pairs of brain areas by analyzing
245 snRNA-seq data from the neocortex and striatum of marmoset CJ027; the resident microglial
246 populations with different genomes make it possible to compare contextual to genetic effects on
247 microglial gene expression (**Fig. 5D**). Genetic effects appeared to elicit very many small-
248 magnitude gene-expression differences; these differences were shared between cortical and

striatal microglia (**Fig. 5E**). Brain-area context elicited much larger-magnitude gene-expression differences, which were experienced in common by microglia with both genotypes (**Fig. 5F**). We obtained similar results for all pairs of brain areas analyzed (52 context vs genetic effect from brain snRNA-seq of 6 marmosets with at least 60 cells available for analysis in each context; **Supplementary Table 4**; **Supplementary Fig. 5**).

We next considered the difference in contexts provided by the same brain area in different marmosets. Two of the marmosets profiled, CJ006 and CJ007, were birth siblings who passed away as neonates (the only birth siblings in our dataset), and thus provided the additional opportunity to distinguish genetic from contextual effects by analyzing the two sibling microglial populations in the cortex, striatum and hippocampus of both marmosets (**Fig. 5G**). The effects of context (host marmoset) in microglia from all three brain areas appeared to be far larger than the cell-autonomous effects of genetic differences (**Fig. 5H,I**).

Discussion

A longstanding debate concerns the extent of chimerism in marmosets and tamarins. Chimerism in these species has been detected in diverse organs but arises from unknown cell types (Ross et al., 2007; Sweeney et al., 2012). Here we found that chimerism in the brain, liver and kidney is present but appears to arise entirely from cells of the myeloid and lymphoid lineages, including infiltrating macrophages, monocytes, and microglia.

Cells of the myeloid and lymphoid lineages derive developmentally from hematopoietic stem cells. We found no strong evidence of chimerism among 2.2 million non-hematopoietic cells in the liver (from one marmoset), kidney (from one marmoset) or brain (from 11 marmosets). Thus, while marmosets share a circulation *in utero*, we found no evidence that non-hematopoietic stem cells or progenitors had been shared via this route in any appreciable number. However, we found that in the marmoset brain, the microglia and macrophages, which also derive from this lineage, routinely harbor abundant chimerism, with 10-50% of a marmoset's microglia containing the genome(s) of birth sibling(s).

Organs in the same marmoset (liver, kidney, brain) differed markedly in the sibling contribution to resident macrophage and monocyte populations, with microglial chimerism fraction (the fraction of cells contributed by siblings) varying by as much as 40 percentage points across a marmoset's brain areas. This phenomenon has more than one potential explanation. First, cells from the host and sibling could in principle respond differently to recruitment or proliferation cues that vary spatially; if this is the case, marmoset chimerism could provide a model for studying the effects of mutations and natural sequence variation on cell migration and recruitment. (Although we found that genetic effects were smaller than contextual effects in shaping microglial gene expression at any moment in time, genetic effects were clear (**Fig. 5E**), and even small effects on proliferation rates would tend to have effects that increase exponentially over time.) Second, beyond such recruitment effects, it is also possible that these differences suggest a substantial role of clonal expansions and population bottlenecks in shaping local microglial and macrophage populations.

We found that the cellular contribution of birth siblings to myeloid cell populations was significantly different in blood than in brain in the modest number of marmosets analyzed (**Fig. 3**). Unlike the differences among brain areas, the blood-brain differences tended to be directional, with more-modest sibling contributions in the brain than in the blood. Though this would need to be confirmed in many more marmosets to be definitive on its own, it is plausibly connected to this aspect of marmoset fetal development: sharing of a blood circulation between

the two fetuses occurs during a temporal window that is more temporally extended than the waves of colonization of the brain by microglia, potentially allowing for greater exchange in the centers of blood hematopoiesis (the liver and then the bone marrow). In microglia and macrophages, due to the defined waves of hematopoiesis in the yolk sac and migration patterns of microglia and macrophages to the developing brain, the opportunity for a progenitor cell from a twin to colonize a host's brain may need to occur during a more-restricted temporal window.

Comparisons of gene expression between microglial cells with host and sibling genomes in a shared brain context may provide many future opportunities to distinguish the cell-autonomous from the non-cell autonomous genetic effects of genetic differences and engineered mutations. Such analyses could become especially useful scientifically as genome editing increasingly enables the utilization of marmosets as a model organism in translational neuroscience (Aida & Feng, 2020; Feng et al., 2020). Our pilot analysis of host-sibling microglial gene expression differences in the brains of two co-twins revealed a large role of animal context (relative to genetic differences) in shaping microglial gene expression. This result points to an important principle: the ability to isolate the effects of a mutation will be greatly strengthened by the ability to make within-animal (rather than just between-animal) comparisons of cells with different genotypes.

A long history of innovation in genetics involves elaborate ways to create mosaics in mice, *C. elegans* and other laboratory organisms in order to distinguish cell-autonomous from non-cell autonomous genetic effects. Natural chimerism in marmosets may enable many straightforward ways to pursue such kinds of studies. Natural chimerism may also make it possible to determine when microglia or macrophages, as opposed to other cell types, mediate the effect of a mutation on an animal's phenotype.

Microglia perform essential roles in the development and regulation of the central nervous system (CNS), including by sculpting or "pruning" neuronal circuits (Hammond et al., 2018; Schafer et al., 2012; Schafer & Stevens, 2015), and are implicated in or hypothesized to contribute to a wide range of brain disorders and diseases, including Alzheimer's disease, Parkinson's disease, autism spectrum disorder, and schizophrenia. Marmoset microglial chimerism will enable many new ways of studying microglia and the effects of genes and alleles upon brain biology.

Methods

Ethical compliance

Marmoset experiments were approved by and in accordance with Massachusetts Institute of Technology IACUC protocol number 051705020.

Nucleus Drop-seq library preparation and sequencing

Nucleus suspensions were prepared from frozen tissue and used for nucleus Drop-seq following the protocol we have described at https://doi.org/10.17504/protocols.io.2srged6. Drop-seq libraries were prepared as previously described (Macosko et al., 2015), with modifications, quantification and quality control as described in a previous study (Saunders et al., 2018), as well as the following modifications optimized for nuclei: in the Drop-seq lysis buffer, 8 M guanidine hydrochloride (pH 8.5) was substituted for water, nuclei were loaded into the syringe at a concentration of 176 nuclei/μl, and cDNA amplification was performed using around 6,000 beads per reaction, 15 PCR cycles. Raw sequencing reads were aligned to the calJac3 marmoset reference genome assembly and reads that mapped to exons or introns of each

assembly were assigned to annotated genes (https://github.com/broadinstitute/Drop-seq). Drop-seq libraries are indicated in Supplementary Table 1.

Nucleus 10X Chromium library preparation and sequencing

Single-nucleus suspensions from frozen tissue were generated as for Drop-seq; GEM generation and library preparation followed the manufacturer's protocol (protocol versions #CG00052 Chromium Single Cell 3' v2 and #CG000183 Chromium Single Cell3′ v3 UG_Rev-A). Raw sequencing reads were processed and aligned using the same method for aligning Drop-seq reads. 10X Chromium libraries are indicated in Supplementary Table 1.

Clustering of cells using Independent Component Analysis

Nuclei from intact cells were identified and clustered into cell types using a method that we have previously described (Krienen et al., 2020; Saunders et al., 2018). Briefly, nuclei with less than 400 detected genes were not used in the analysis. A digital gene expression matrix was created for a set of libraries from the same animal that were to be co-analyzed (**Supplementary Table 5**), and independent component analysis using the `fastICA` package in R was used after normalization and variable gene selection as previously described (Krienen et al., 2020; Saunders et al., 2018). A Louvaine-based clustering algorithm was performed on the top 60 independent components. Due to the large number of nuclei profiled in some marmosets (CJ027, CJ028, CJ029), memory requirements exceeded machine limits and for these marmosets, we divided the clustering analysis into two or three batches (**Supplementary Table 5**). The brain of marmoset CJ022 was profiled using both Drop-seq and 10X and a separate clustering was done for each snRNA-seq method (**Supplementary Table 5**). We ran the clustering algorithm 12 times using 3 nearest neighbor parameters (10,20,30) and 4 resolution parameters (0.3, 0.5, 0.1, 1.0). Markers for each cluster were identified using differential gene expression analysis (Krienen et al., 2020; Saunders et al., 2018). We inspected each clustering result and chose the one which yielded separate clusters for microglia and macrophages (**Supplementary Table 5**).

Identification of cell types

For the brain datasets, the microglia and macrophage clusters were identified by the markers *TREM2, C3, LAPTM5* for microglia and *F13A1*, *LYVE1* for macrophages. The other brain cell types were identified using cell type markers for neurons, astrocytes, oligodendrocytes, polydendrocytes and endothelial cells that we used as before (Krienen et al., 2020). Cell types in blood, liver and kidney were identified using the `ScType` method (Ianevski et al., 2022).

Donor-of-origin analysis and detection of host-sibling doublets (Dropulation)

We used the Dropulation suite to calculate a donor likelihood for each cell (Wells et al., 2023) (software available at https://github.com/broadinstitute/Drop-seq). The host and birth sibling genotypes were provided as input to Dropulation's `AssignCellsToSamples` tool, together with the snRNA-seq BAM file and a list of cell barcodes that were identified to be intact cells. To generate chimerism-free reference genotypes, we cultured fibroblasts and performed whole genome sequencing (WGS) on the resulting DNA. We found that the difference in likelihoods between host and sibling increase with the number of UMI of the cell, and hence we imposed a minimum number of UMI for each marmoset's cells (**Supplementary Table 5**). We also performed doublet detection using Dropulation's `DetectDoublets` to obtain a likelihood of a cell having a mix of transcripts from the host and sibling(s). Doublets lie between the host and sibling curves (**Supplementary Fig. 6**) and for each marmoset we empirically obtained a threshold for the Dropulation test statistic to identify them and were discarded in all analyses

399 (**Supplementary Fig. 6**, **Supplementary Table 5**). The likelihoods plotted in Figures 1C,1E,2C
400 and Supplementary Fig. 2 are from cells that have been filtered for minimum UMI and doublets.
401
402 Marmosets CJ006 and CJ007 were born in a triplet litter (tri-zygotic) that all died shortly after
403 birth. We did not have access to any tissue from the third sibling and were not able to perform
404 whole genome sequencing on it. For Dropulation analysis of CJ006 and CJ007's brains, we
405 provided only the genotypes of marmosets CJ006 and CJ007. Nuclei that contain the genome
406 of the third unknown sibling will mostly be identified as doublets and were discarded in our
407 analysis.
408
409 **Additional filtering for microglia and macrophage clusters**
410 We performed additional filtering of microglia and macrophage cells. When we compared the
411 gene expression of host microglia and sibling microglia using cell-types from first-round
412 clustering (and with UMI and doublet filtering), we found an abundance of genes that have
413 higher expression in host than in the sibling (**Supplementary Fig. 7**, see panels B, C, G, I and
414 K). The asymmetry could arise from neuronal cells mis-classified as microglia or macrophages.
415 To filter out these mis-classified cells, we sub-clustered the microglia and macrophage cell
416 types of each marmoset using the same fastICA and Louvaine-based clustering used in the first
417 round of clustering. We found that some sub-clusters were not chimeric, indicating that they
418 were not cells of hematopoietic origin, and that discarding these cells improved the symmetry
419 between host and sibling gene expression (**Supplementary Fig. 7**).
420
421 **Whole genome sequencing**
422 Illumina libraries from fibroblast, blood, brain and buccal cells (**Supplementary Table 6**) were
423 created as follows. An aliquot of genomic DNA (150ng in 50µL) is used as the input into DNA
424 fragmentation (aka shearing). Shearing is performed acoustically using a Covaris focused-
425 ultrasonicator, targeting 385bp fragments. Following fragmentation, additional size selection is
426 performed using a SPRI cleanup. Library preparation is performed using a commercially
427 available kit provided by KAPA Biosystems (KAPA Hyper Prep with Library Amplification Primer
428 Mix, product KK8504), and with palindromic forked adapters using unique 8-base index
429 sequences embedded within the adapter (purchased from Roche). The libraries are then
430 amplified by 10 cycles of PCR. Following sample preparation, libraries are quantified using
431 quantitative PCR (kit purchased from KAPA Biosystems) with probes specific to the ends of the
432 adapters. This assay is automated using Agilent's Bravo liquid handling platform. Based on
433 qPCR quantification, libraries are normalized to 2.2nM and pooled into 24-plexes. Sample pools
434 are combined with NovaSeq Cluster Amp Reagents DPX1, DPX2 and DPX3 and loaded into
435 single lanes of a NovaSeq 6000 S4 flowcell cell using the Hamilton Starlet Liquid Handling
436 system. Cluster amplification and sequencing occur on NovaSeq 6000 Instruments utilizing
437 sequencing-by-synthesis kits to produce 151bp paired-end reads. Output from Illumina software
438 is processed by the Picard data-processing pipeline to yield CRAM or BAM files containing
439 demultiplexed, aggregated aligned reads. All sample information tracking is performed by
440 automated LIMS messaging. All samples were sequenced to 30X coverage.
441
442 **Variant site detection and genotyping from whole genome sequencing**
443 Illumina paired-end reads were aligned to the calJac3 reference marmoset genome assembly
444 using bwa (Li & Durbin, 2010) with command "`bwa mem`". Duplicate reads were marked using
445 Picard Markduplicates and for each chromosome, the GATK Haplotype Caller (McKenna et al.,
446 2010) was run in genotype discovery GVCF mode. For each chromosome, the GVCFs of all
447 samples analyzed in this study (from fibroblasts, blood, buccal cells, skin, brain and hair), were
448 combined into a single GVCF file using GATK CombineGVCFs. To obtain the highest sensitivity
449 in calling SNPs, we included in the GVCF additional fibroblasts whole genome sequences from

the colony, yielding a total of 113 marmosets for multi-sample variant calling. The GVCF of each chromosome was genotyped using GATK GenotypeGVCFs. Only bi-allelic SNPs were used in the analysis and the following filters were used: QD<4.0 | FS>60.0 | MQ<40.0 | MQRankSum<-12.5 | ReadPosRankSum<-8.0 | MAF<0.01 | QUAL<500. SNP calls from all chromosomes were combined into one VCF file and additional filtering was performed to discard heterozygous sites that exhibited extreme allelic imbalance, i.e., the fraction of non-reference allele (from all samples) is less than 0.2 or greater than 0.8, and furthermore, sites in copy number variant regions were discarded (copy number variant regions were obtained by running Genome STRiP (Handsaker et al., 2015) on whole genome sequencing data from 113 fibroblast samples).

Dropulation analysis using sibling genotypes from whole genome sequencing of buccal cells

For four marmosets in our dataset (CJ022, CJ025, CJ026, CJ102; all born with one sibling), only the buccal cells (from cheek swabs) of their siblings were available for whole genome sequencing (the siblings are, CJ106, CJ104, CJ105 and CJ103, respectively). Using a method that quantifies chimerism from whole genome sequencing data (Census-seq; software available at https://github.com/broadinstitute/Drop-seq) (Mitchell et al., 2020), we estimated the chimerism fraction in buccal cells as follows: CJ106: 10%, CJ104: 24%, CJ105: 24% and CJ103: 9%. Thus, the genotypes we obtained for these marmosets will include errors and those genotyping errors could subsequently affect Dropulation (donor-of-origin) analysis that was used to estimate chimerism. To empirically estimate how sibling genotypes obtained from a chimeric tissue affect Dropulation analysis, we selected a host-sibling pair whose genome sequencing were both obtained from fibroblast cultures: CJ027 and its birth sibling CJ140. To simulate DNA contamination, we fixed the sequencing coverage of CJ140 to 40X, and replaced between 1% to 60% of the reads from CJ027's sequencing reads (random subsampling using '`samtools view -s`'). We genotyped CJ140's "chimeric" bam files using GATK's "genotype given alleles" mode and compared the genotypes with CJ140's true genotypes from its pure fibroblast WGS. We found that the sensitivity at heterozygous sites remains constant with different contamination levels, while the false positive rate increases. The false positive calls at heterozygous sites come from homozygous sites incorrectly genotyped as heterozygous (**Supplementary Fig. 8A-F**). Next, we re-analyzed donor-of-origin on brain snRNA-seq of CJ027 sibling (CJ140) genotypes from simulated contaminated DNA (300,000 nuclei; CJ027 genotypes from pure fibroblast WGS, CJ140 genotypes from WGS with various contamination levels). We found that chimerism in CJ140's WGS resulted in doublets being assigned to the twin (**Supplementary Fig. 8G-L**), which subsequently causes a slight increase in chimerism estimates (**Supplementary Fig. 8M-U**). The contamination levels in buccal cells were from 9% to 24%, which we estimate will result in an overestimation in microglia chimerism of up to 3.5 percentage points and 4.5 percentage points in macrophage chimerism. Our results will not be affected by this over-estimation of chimerism in 4 marmosets since the conclusions were made from the analysis of all 11 marmosets (including 7 marmosets whose siblings were genotyped from fibroblast cultures).

Gene expression analysis of host and sibling meta cells

For each cell type, the host and sibling "meta cells" were calculated from the sum of UMI counts per gene across cells and were scaled to counts per 100,000 transcripts. The fold-changes and *P*-values of differentially expressed genes were identified using the `binomTest` method from the *edgeR* package (Robinson et al., 2010).

Software availability

All software used in the analysis are publicly available. Drop-seq (analysis of snRNA-seq data, clustering, marker genes), Census-seq (estimation of chimerism in whole-genome sequencing data) and Dropulation analysis (estimation of chimerism in snRNA-seq data): https://github.com/broadinstitute/Drop-seq; alignment and variant detection of Illumina whole genome sequencing data: bwa (https://github.com/lh3/bwa), GATK (https://gatk.broadinstitute.org), BCFtools (https://github.com/samtools/bcftools), samtools (http://www.htslib.org/download), Picard Tools (https://broadinstitute.github.io/picard); R environment (https://www.rstudio.com/products/rstudio/download/, https://www.r-project.org); cell type identification scType (https://github.com/IanevskiAleksandr/sc-type).

Data availability

Brain snRNA-seq of 6 marmosets (CJ022, CJ023, CJ025, CJ026, CJ027, CJ028) were generated as part of the NIH's Brain Initiative Cell Census Network (BICCN) project, while brain snRNA-seq of 5 marmosets (CJ001, CJ006, CJ007, CJ023, CJ102), and all blood, liver and kidney snRNA-seq were generated for this project. All snRNA-seq datasets are available in the BICCN NeMO portal (https://assets.nemoarchive.org/dat-hsgdsgu and https://assets.nemoarchive.org/dat-1je0mn3). All whole genome sequencing datasets will be provided in a manuscript (in preparation) that will describe naturally occurring genome variation in captive marmosets.

Acknowledgements

This work was supported by the NIH BRAIN Initiative U01MH114819, the Stanley Center for Psychiatric Research at the Broad Institute of MIT and Harvard, the James and Patricia Poitras Center for Psychiatric Disorders Research at MIT and the Hock E. Tan and K. Lisa Yang Center for Autism Research at MIT.

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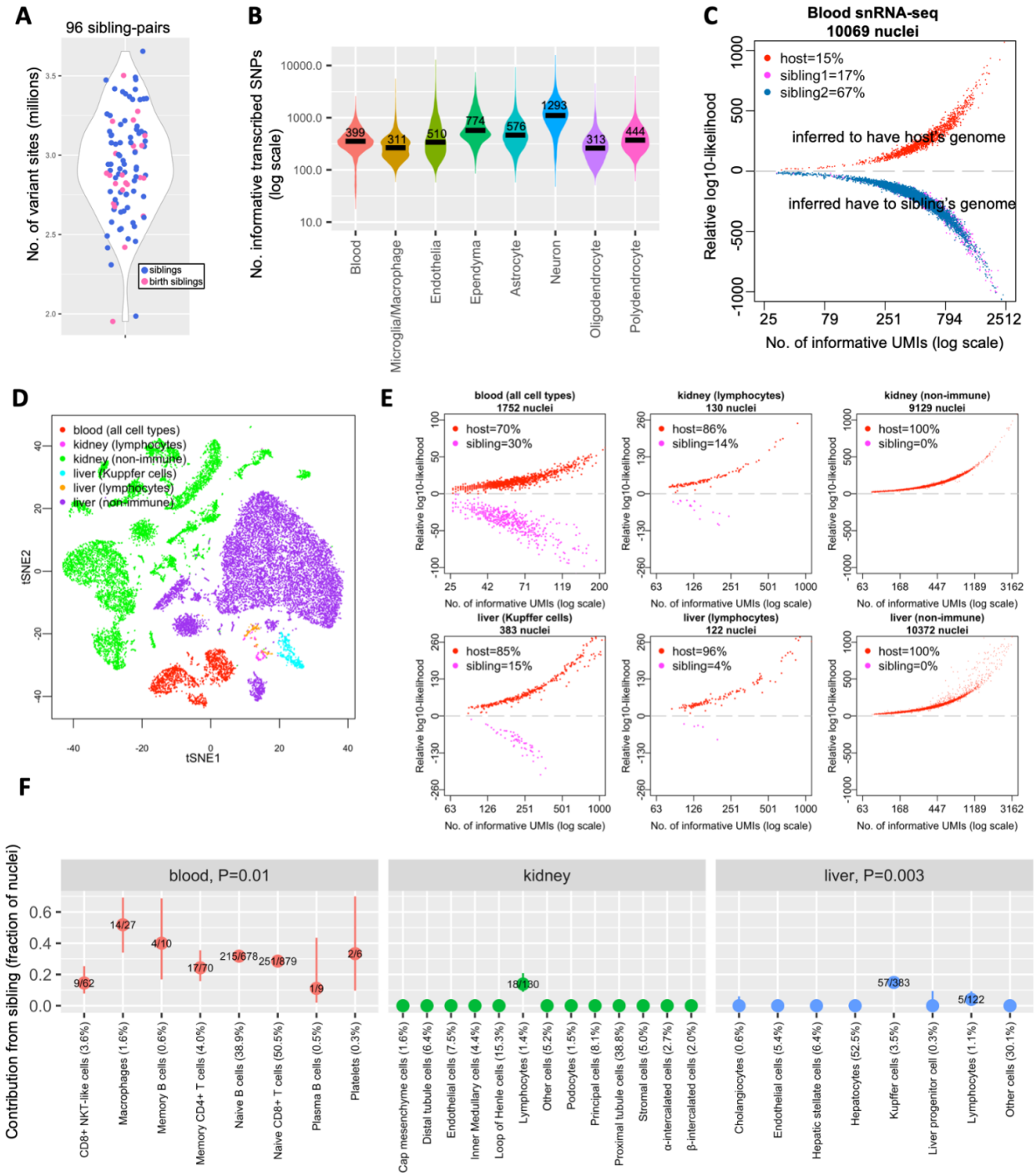

Figure 1. Sibling chimerism analysis at single-cell resolution. (A) Numbers of variant sites at which marmoset sibling genomes differ, for 96 sibling-pairs. Each dot is a sibling-pair; birth siblings are colored in pink. **(B)** Numbers of transcribed SNPs (per nucleus) visible in various cell types from blood and brain snRNA-seq datasets of marmoset CJ028. x-axis: snRNA-seq library; y-axis: number of ascertained, transcribed SNPs for which host and sibling have different genotypes; black horizontal lines: median values per cell type. **(C)** Donor-of-origin assignment of each nucleus in blood snRNA-seq of a marmoset. The host marmoset (CJ028) was born with two siblings; each nucleus was assigned to either the host or to one of the two

621 birth siblings. x-axis: number of unique molecular identifiers (UMI; a measure of transcript
622 abundance) that contains SNPs for which the host and sibling's genomes differ, in log scale; y-
623 axis: inferred likelihood that the cell has host genome minus likelihood that the cell has sibling
624 genome (log10). (**D**) Two-dimensional visualization (tSNE plot) of snRNA-seq data from a
625 marmoset's (CJ026) blood, kidney and liver. **(E)** Donor-of-origin assignment in marmoset
626 CJ026's blood, kidney and liver. axes: same as in (C). **(F)** Levels of chimerism in each cell type
627 ascertained in blood, kidney and liver snRNA-seq of marmoset CJ026. y-axis: fraction of sibling
628 cells in each cell type; numbers in fraction: number of sibling nuclei over total nuclei in the cell
629 type; percentage in x-axis labels: cell type representation in the tissue; vertical bars: binomial
630 confidence interval (95%); *P*-values: test of heterogeneity (Chi-square) across immune cell
631 types of a tissue (to test for differences in contribution of sibling across immune cell types).
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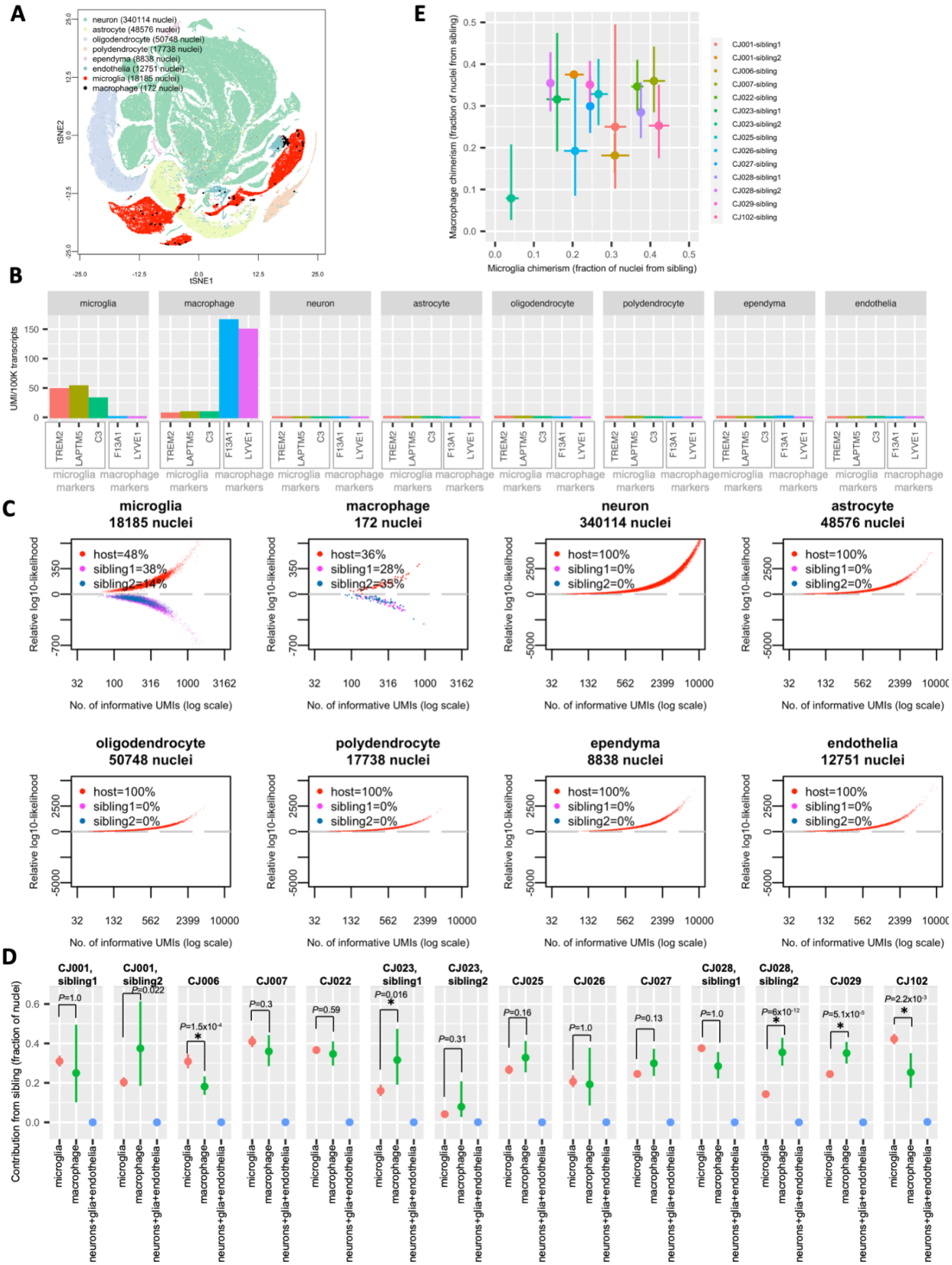


Figure 2. Microglia and macrophages, but not neurons and glia, are chimeric in the

marmoset brain. (A) Two-dimensional visualization (tSNE plot) of marmoset CJ028's snRNA-seq profiles from 8 brain regions. **(B)** Expression of markers of microglia and macrophages in microglia, macrophages, neurons, glia (astrocytes, oligodendrocytes, polydendrocytes, ependymal cells) and endothelial cells. y-axis: expression levels, measured as numbers of detected transcripts (unique molecular identifiers (UMIs), a measure of transcript abundance) from that gene per 100,000 total. Microglia markers: *TREM2, LAPTM5, C3*; macrophage markers: *F13A1, LYVE1*. Data from all CJ028's brain regions. **(C)** Donor assignment of each nucleus to one of three possible donors (host, sibling1 or sibling2) of marmoset CJ028's brain snRNA-seq data. x-axis: number of UMI that contains SNPs for which the host and sibling's genomes differ, in log10 scale; y-axis: inferred likelihood that the cell has host genome minus likelihood that the cell has sibling genome (log10). Data from all of CJ028's brain regions. **(D)** Fractions of cells with a sibling's genome among microglia, macrophages, and other brain cell types (neurons, astrocytes, oligodendrocytes, polydendrocytes, ependyma, endothelial) from 11 marmosets. CJ001, CJ023 and CJ028 were part of a triplet litter and the chimerism fraction of each birth sibling is shown in separate panels. y-axis: fraction of sibling-cells in the cell type. vertical bars: binomial confidence interval (95%). *P*-values: Chi-square test from comparison of chimerism fractions in microglia and macrophage; * (*P*-value<0.05). (**E**) Sibling contributions to microglial (x-axis) and macrophage (y-axis) populations in the same animals. Dots and bars are from (D). Pearson correlation *R*=0.32, 95% confidence interval (-0.26 to 0.73), *P*-value=0.27.

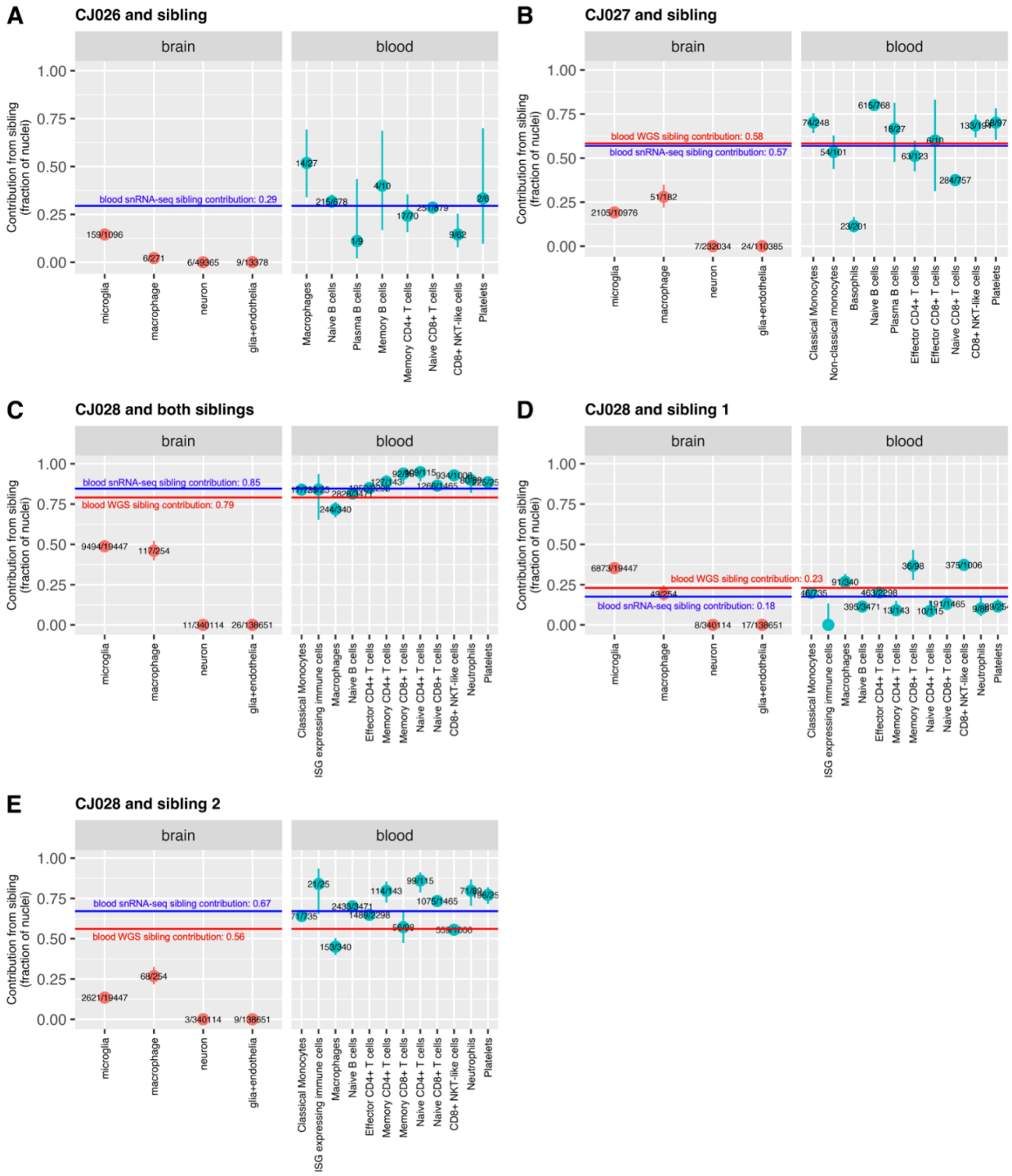


Figure 3. Sibling contributions to hematopoiesis-derived cells diverge between blood and brain. (A)-(E) Chimerism fractions in brain and blood of three animals. CJ028 was part of a triplet litter, and the contribution of each sibling is shown in a separate panel (D,E). Red horizontal lines: twin contribution to blood cells as ascertained from whole-genome sequencing of whole-blood-derived genomic DNA (Census-seq). Blue horizontal lines: total twin contribution to blood cells as estimated from PBMC snRNA-seq (all cells). Vertical bars: binomial confidence interval (95%). Numbers in fraction: number of sibling cells over total cells in the cell type.

glia+endothelia: astrocytes, oligodendrocytes, polydendrocytes, ependymal cells, endothelial cells.

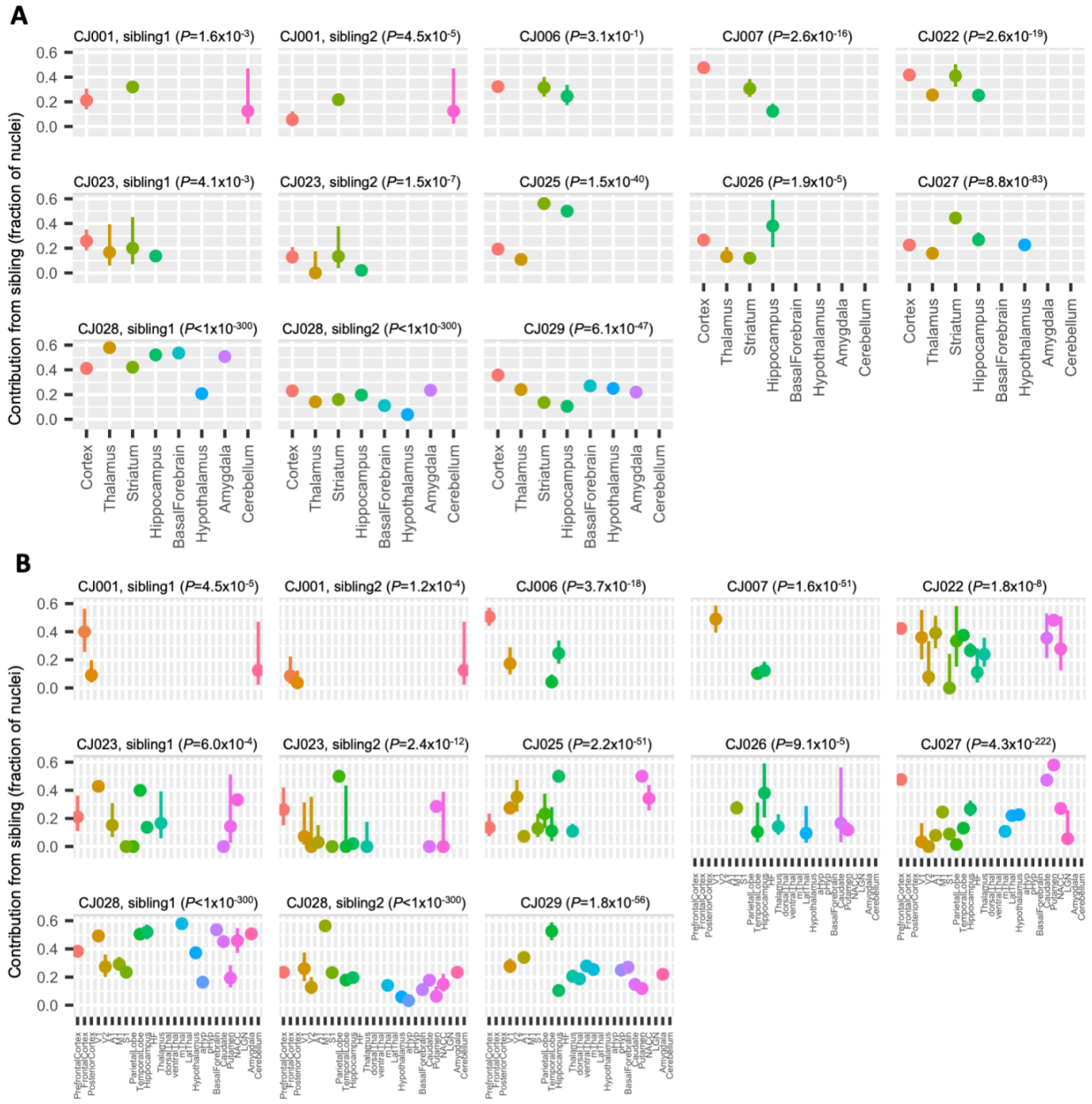
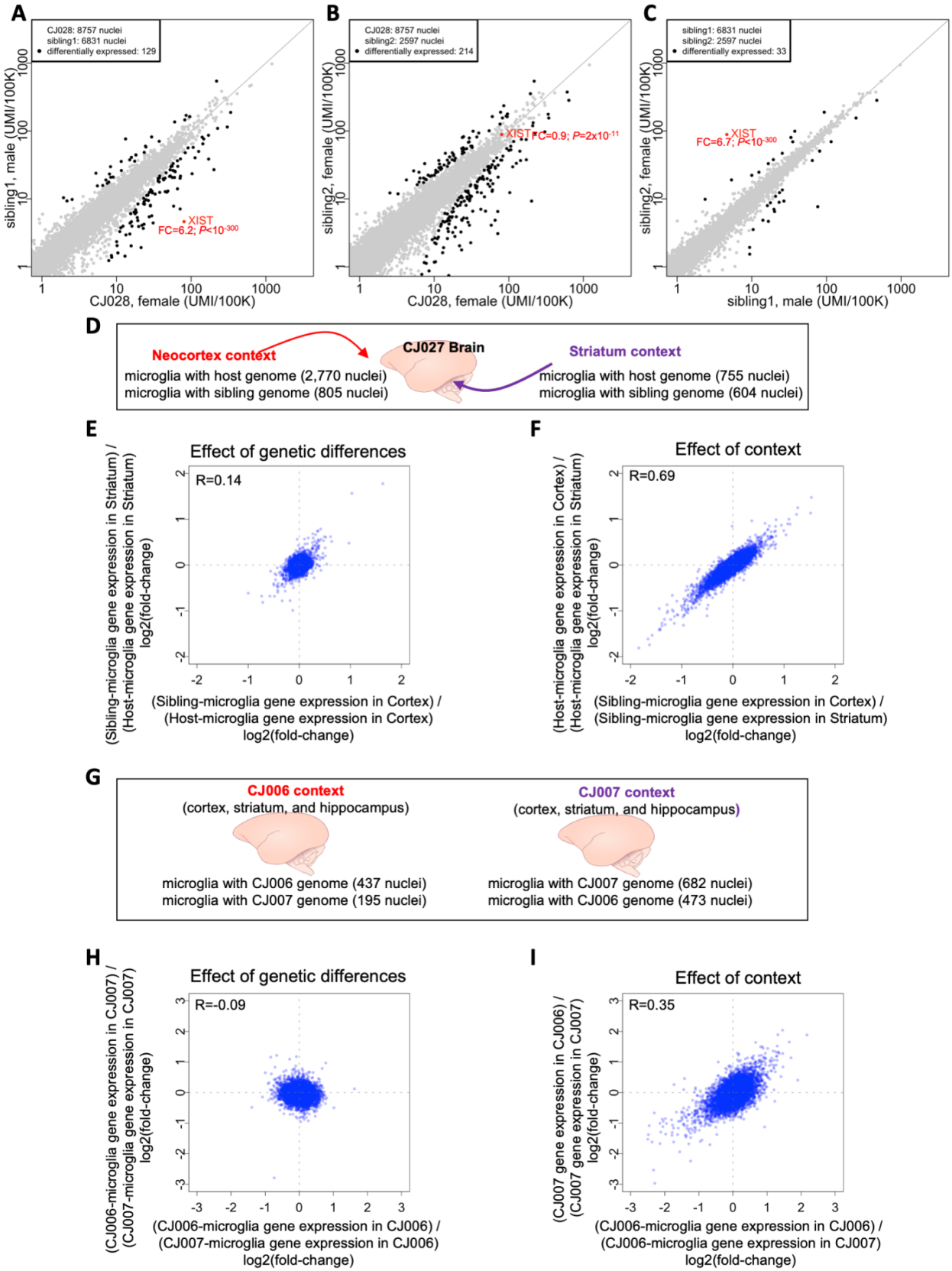


Figure 4. **Sibling contributions to brain microglial populations vary across an animal's brain areas.** Contributions of sibling(s) to the microglial populations ascertained in principal brain areas **(A)**, and finer-scale brain substructures **(B)**. CJ001, CJ023 and CJ028 were part of a triplet litter and the chimerism contribution of each twin is shown in a separate panel. CJ102 was profiled in only one brain region and hence was not included in the analysis. y-axis: fraction of twin cells. Vertical bars: binomial confidence interval (95%); *P*-values: test of heterogeneity across an animal's brain regions.



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684 **Figure 5. Utilizing natural chimerism to distinguish cell-autonomous from non-cell-**

autonomous effects on gene expression, and to compare the effects context and genetic variation in shaping gene expression. (A-C) Comparisons of RNA expression between microglial populations within host animal CJ028, who had two birth siblings. Comparisons of gene expression between microglia with the genomes of **(A)** the female host and male sibling, **(B)** the female host and female sibling, and **(C)** the two siblings (male and female). Each point represents a gene; its location on the plot represents the level of expression of that gene among microglia with two different genomes in the same animal. x- and y-axes: normalized gene expression levels (number of transcripts per 100,000 transcripts). FC: fold-change of gene expression, female/male for *XIST*. Fold-change and *P*-values were calculated using edgeR. Differentially expressed genes (black dots) were defined as: FDR *Q*-value<0.05 and fold-change>1.5 or less than 1/1.5 and the gene must be expressed in at least 10% of one of the microglia sets. **(D-I)** Higher effect of context than genetic differences in shaping gene expression. **(D)** In the brain of marmoset CJ027, the neocortex and striatum are two contexts where two sets of microglia with different genomes reside. **(E)** Effect of genetic differences. x-axis: log2-fold-change of cortical microglia gene expression between host and sibling cells; y-axis: log2-fold-change of striatal microglia gene expression between host and sibling cells. **(F)** Effect of context. x-axis: log2-fold-change of the sibling's gene expression between the two brain regions; y-axis: log2-fold-change of the host's gene expression between the two brain regions. **(G-I)** The brains (cortex, striatum and hippocampus) of two birth siblings provide biological contexts in which populations of microglia with two sibling genomes reside. The effect of genetic differences (H) and effect of animal context (I) are compared, for the same brain areas (combined data from cortex, striatum and hippocampus). x- and y-axes: log2-fold-changes of the gene expression between two sets of microglia (the sets being compared are indicated in the axis labels). *R*: Spearman correlation.

712 **Supplementary Table 1.** Marmosets analyzed with snRNA-seq in this study. Colonies: NEPRC
713 - New England Primate Research Colony; CLEA - Central Institute for Experimental Animals,
714 Japan; Company A: marmosets obtained from a non-clinical contract research organization.

ID	Tissue	Sex	Age	No. birth siblings (sex of sibling(s))	Colony	Single-cell platform
CJ001	Brain	F	Neonate	2 (M, M)	NEPRC	Drop-seq
CJ006	Brain	M	Neonate	2 (F, unknown)	CLEA	10X
CJ007	Brain	F	Neonate	2 (M, unknown)	CLEA	10X
CJ022	Brain	F	2 yrs 8 mos	1 (M)	NEPRC	Drop-seq, 10X
CJ023	Brain	F	2 yrs	2 (F, M)	NEPRC	Drop-seq
CJ025	Brain	M	2 yrs	1 (M)	NEPRC	Drop-seq
CJ026	Brain	M	2 yrs 8 mos	1 (F)	NEPRC	Drop-seq
CJ026	Blood					10X
CJ026	Liver					10X
CJ026	Kidney					10X
CJ027	Brain	M	2 years	1 (M)	CompanyA	10X
CJ027	Blood					10X
CJ028	Brain	F	2 years	2 (M, F)	NEPRC	10X
CJ028	Blood					10X
CJ029	Brain	M	2 years	1 (M)	CLEA	10X
CJ102	Brain	F	6 mos	1 (M)	CompanyA	Drop-seq, 10X

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717 **Supplementary Table 2.** Number of microglia and macrophage cells identified in brain datasets
718 and number of nuclei profiled in blood, liver and kidney.

ID	Tissue	Microglia	Macrophage	Other Cell Types	Total nuclei	Percent Microglia	Percent Macrophage
CJ001	Brain	1,063	16	89,599	90,678	1.17%	0.02%
CJ006	Brain	632	259	165,272	166,163	0.38%	0.16%
CJ007	Brain	1,155	139	105,407	106,701	1.08%	0.13%
CJ022	Brain	3,659	228	217,312	221,199	1.65%	0.10%
CJ023	Brain	581	38	79692	80,311	0.72%	0.05%
CJ025	Brain	1,366	131	210628	212,125	0.64%	0.06%
CJ026	Brain	693	26	62743	63,462	1.09%	0.04%
CJ027	Brain	8,398	167	342,419	350,984	2.39%	0.05%
CJ028	Brain	18,185	172	478,765	497,333	3.66%	0.03%
CJ029	Brain	11,874	285	320,000	332,159	3.57%	0.09%
CJ102	Brain	1,124	43	56,602	57,769	1.95%	0.07%
CJ026	Blood	NA	NA	1,741	1,741	NA	NA
CJ026	Liver	NA	NA	10,877	10,877	NA	NA
CJ026	Kidney	NA	NA	9,262	9,262	NA	NA
CJ027	Blood	NA	NA	2,529	2,529	NA	NA
CJ028	Blood	NA	NA	10,042	10,042	NA	NA

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723 **Supplementary Table 3**. Comparison of chimerism between CJ028's two birth siblings, in blood
724 and in brain myeloid cells (microglia and macrophage). *P*-values are from a two-sided test of
725 proportions between chimerism fractions of sibling 1 and sibling 2 using the prop.test function in
726 R.

Cell type	Total nuclei	Sibling 1 nuclei	Sibling 2 nuclei	Sibling 1 Fraction	Sibling 2 Fraction	Sibling 1 + Sibling 2 Fraction	*P*-value
microglia and macrophage	19701	6922	2689	0.35	0.14	0.49	$<2.2\times10^{-16}$
microglia only	19447	6873	2621	0.35	0.13	0.49	$<2.2\times10^{-16}$
macrophage only	254	49	68	0.20	0.27	0.46	5.8×10^{-2}
blood	10042	1758	6739	0.18	0.67	0.85	$<2.2\times10^{-16}$

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730 **Supplementary Table 4.** Summary of context versus genetic effects analysis, with two brain
731 regions of an animal as two contexts. The analysis described in Fig. 5D-F was repeated across
732 all animals and brain regions with at least 60 cells that are available for analysis in each context,
733 and the summary of the correlations are tabulated here. The correlations are plotted in
734 Supplementary Fig. 5. Abbreviations; STR: striatum; Thal: thalamus; Hippo: hippocampus; Hyp:
735 hypothalamus; BF: basal forebrain.

Host ID	Sibling ID	Brain Region1	Brain Region2	Genetic Effects Spearman Correlation	Context Effects Spearman Correlation
CJ022	CJ106	Cortex	Thal	0.1	0.73
CJ025	CJ104	Cortex	STR	0.03	0.2
CJ027	CJ140	Cortex	Thal	0.13	0.64
CJ027	CJ140	Cortex	STR	0.14	0.69
CJ027	CJ140	Cortex	Hippo	0.08	0.46
CJ027	CJ140	Cortex	Hyp	0.03	0.68
CJ027	CJ140	Thal	Hippo	0.04	0.56
CJ027	CJ140	Thal	Hyp	0.09	0.66
CJ027	CJ140	STR	Hippo	0.08	0.43
CJ027	CJ140	STR	Hyp	0.08	0.59
CJ027	CJ140	Hippo	Hyp	0.04	0.51
CJ028	CJ141	Cortex	Thal	0.1	0.66
CJ028	CJ142	Cortex	Thal	0.03	0.55
CJ141	CJ142	Cortex	Thal	0.07	0.59
CJ028	CJ141	Cortex	STR	0.13	0.68
CJ028	CJ142	Cortex	STR	0.1	0.58
CJ141	CJ142	Cortex	STR	0.09	0.61
CJ028	CJ141	Cortex	Hippo	0.06	0.39

CJ028	CJ142	Cortex	Hippo	0	0.33
CJ141	CJ142	Cortex	Hippo	0.04	0.38
CJ028	CJ141	Cortex	BF	0.13	0.73
CJ028	CJ142	Cortex	BF	0.07	0.6
CJ141	CJ142	Cortex	BF	0.09	0.6
CJ028	CJ141	Cortex	Hyp	0.06	0.78
CJ028	CJ142	Cortex	Hyp	0.07	0.72
CJ141	CJ142	Cortex	Hyp	0.08	0.75
CJ028	CJ141	Cortex	Amygdala	0.14	0.65
CJ028	CJ142	Cortex	Amygdala	0.1	0.53
CJ141	CJ142	Cortex	Amygdala	0.12	0.57
CJ028	CJ141	STR	Hippo	0.06	0.63
CJ028	CJ142	STR	Hippo	0.01	0.56
CJ141	CJ142	STR	Hippo	0.03	0.61
CJ028	CJ141	STR	BF	0.14	0.63
CJ028	CJ142	STR	BF	0.06	0.5
CJ141	CJ142	STR	BF	0.05	0.52
CJ028	CJ141	Hyp	Amygdala	0.14	0.68
CJ028	CJ142	Hyp	Amygdala	0.21	0.65
CJ141	CJ142	Hyp	Amygdala	0.07	0.68
CJ029	CJ143	Cortex	Thal	0.09	0.61
CJ029	CJ143	Cortex	STR	0.06	0.5
CJ029	CJ143	Cortex	Hippo	0.05	0.62
CJ029	CJ143	Cortex	BF	0.09	0.79
CJ029	CJ143	Cortex	Hyp	0.07	0.78
CJ029	CJ143	Cortex	Amygdala	0.07	0.32
CJ029	CJ143	Thal	STR	0.09	0.48
CJ029	CJ143	Thal	Hippo	0.06	0.71
CJ029	CJ143	Thal	BF	0.14	0.75
CJ029	CJ143	Thal	Hyp	0.13	0.8
CJ029	CJ143	Thal	Amygdala	0.05	0.52
CJ029	CJ143	Hippo	BF	0.06	0.77
CJ029	CJ143	Hippo	Hyp	0.03	0.78
CJ102	CJ103	Cortex	STR	0.07	0.32

740 **Supplementary Table 5**. Clustering parameters used to identify microglia and macrophage cell
741 types, and thresholds for identifying host-sibling doublets. The final number of microglia and
742 macrophages after the second round of clustering are in Supplementary Table 2. A cell is
743 assigned as a doublet if the Dropulation tool `DetectDoublets` assigned the highest likelihood
744 for the cell as a doublet and if the log10 of the best likelihood minus the log10 of the second-
745 best likelihood (lrt_test_stat) is greater than the doublet detection threshold (last column).

Marmo set ID	Tissue	Resolution parameter	Nearest neighbor parameter	UMI threshold (min log-UMI per cell)	Doublet Detection Threshold (lrt_test_stat)
CJ001	Brain	0.8	10	1.2	0.0
CJ006	Brain	0.5	10	1.2	5.0
CJ007	Brain	0.5	20	1.2	10.0
CJ022	Brain Drop-seq	0.5	20	1.2	2.0
CJ022	Brain 10X	0.5	20	1.5	2.0
CJ023	Brain	0.5	20	1.2	3.0
CJ025	Brain	0.5	20	1.0	0.5
CJ026	Brain	0.5	20	1.0	2.0
CJ026	Blood	0.5	20	1.4	4.0
CJ026	Liver	0.5	20	1.9	8.0
CJ026	Kidney	0.5	20	1.9	8.0
CJ027	Brain Cortex A	0.5	20	1.5	20.0
CJ027	Brain Cortex B	0.5	20	1.5	20.0
CJ027	Brain others	0.5	20	1.5	20.0
CJ027	Blood	0.5	20	1.5	1.0
CJ028	Brain Cortex A	0.5	20	1.5	25.0
CJ028	Brain Cortex B	0.8	10	1.5	25.0
CJ028	Brain others	0.5	20	1.5	25.0
CJ028	Blood	0.5	20	1.5	1.0
CJ029	Brain Cortex	0.5	20	1.5	20.0
CJ029	Brain others	0.5	20	1.5	20.0
CJ102	Brain, Drop-seq	0.8	10	1.2	3.0
CJ102	Brain, 10X	0.5	20	1.2	5.0

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749 **Supplementary Table 6**. Whole genome sequencing datasets used in (1) donor-of-origin
750 assignment from snRNA-seq (Dropulation), and (2) estimating chimerism from blood whole
751 genome sequencing (Census-seq).

ID	Tissue/DNA source	Sequencing coverage
CJ001	Fibroblast culture	36.4X
CJ119 (CJ001's sibling1)	Fibroblast culture	39.7X
CJ120 (CJ001's sibling2)	Fibroblast culture	36.7X
CJ006 (CJ007's sibling)	Fibroblast culture	45.9X
CJ007 (CJ006's sibling)	Fibroblast culture	55.8X
CJ022	Fibroblast culture	38.4X
CJ106 (CJ022's sibling)	Buccal swab	36.0X
CJ023	Fibroblast culture	34.9X
CJ131 (CJ023's sibling1)	Fibroblast culture	39.5X
CJ116 (CJ023's sibling2)	Fibroblast culture	28.3X
CJ025	Fibroblast culture	72.7X
CJ104 (CJ025's sibling)	Buccal swab	58.0X
CJ026	Fibroblast culture	74.1X
CJ105 (CJ026's sibling)	Buccal swab	43.7X
CJ027	Fibroblast culture	47.5X
CJ027	Blood	46.9X
CJ140 (CJ027's sibling)	Fibroblast culture	40.2X
CJ028	Fibroblast culture	44.6X
CJ028	Blood	52.6X
CJ141 (CJ028's sibling1)	Fibroblast culture	43.6X
CJ142 (CJ028's sibling2)	Fibroblast culture	41.2X
CJ029	Fibroblast culture	49.2X
CJ143 (CJ029's sibling)	Fibroblast culture	40.8X
CJ102	Brain	32.4X
CJ103 (CJ102's sibling)	Buccal swab	38.5X

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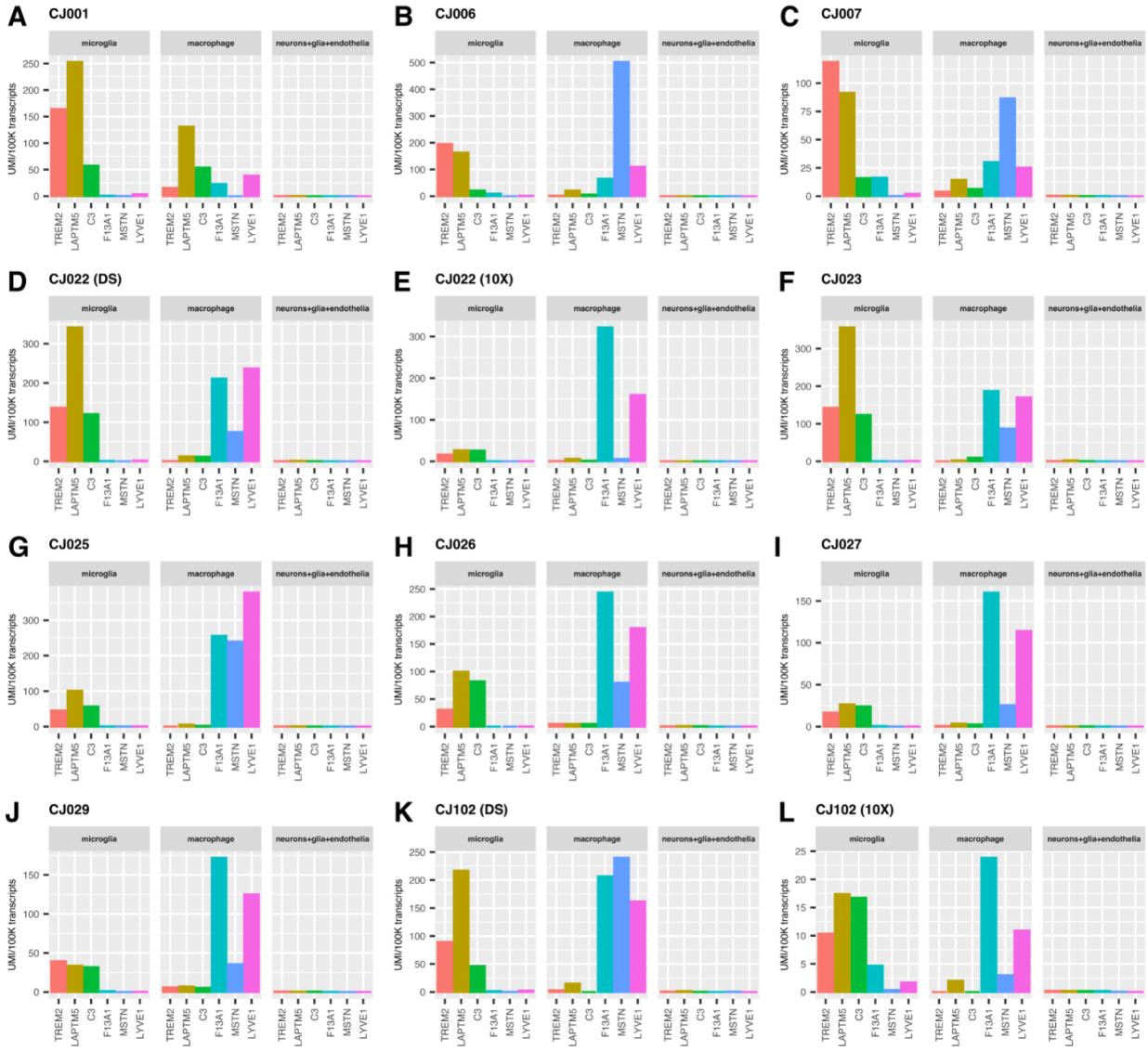
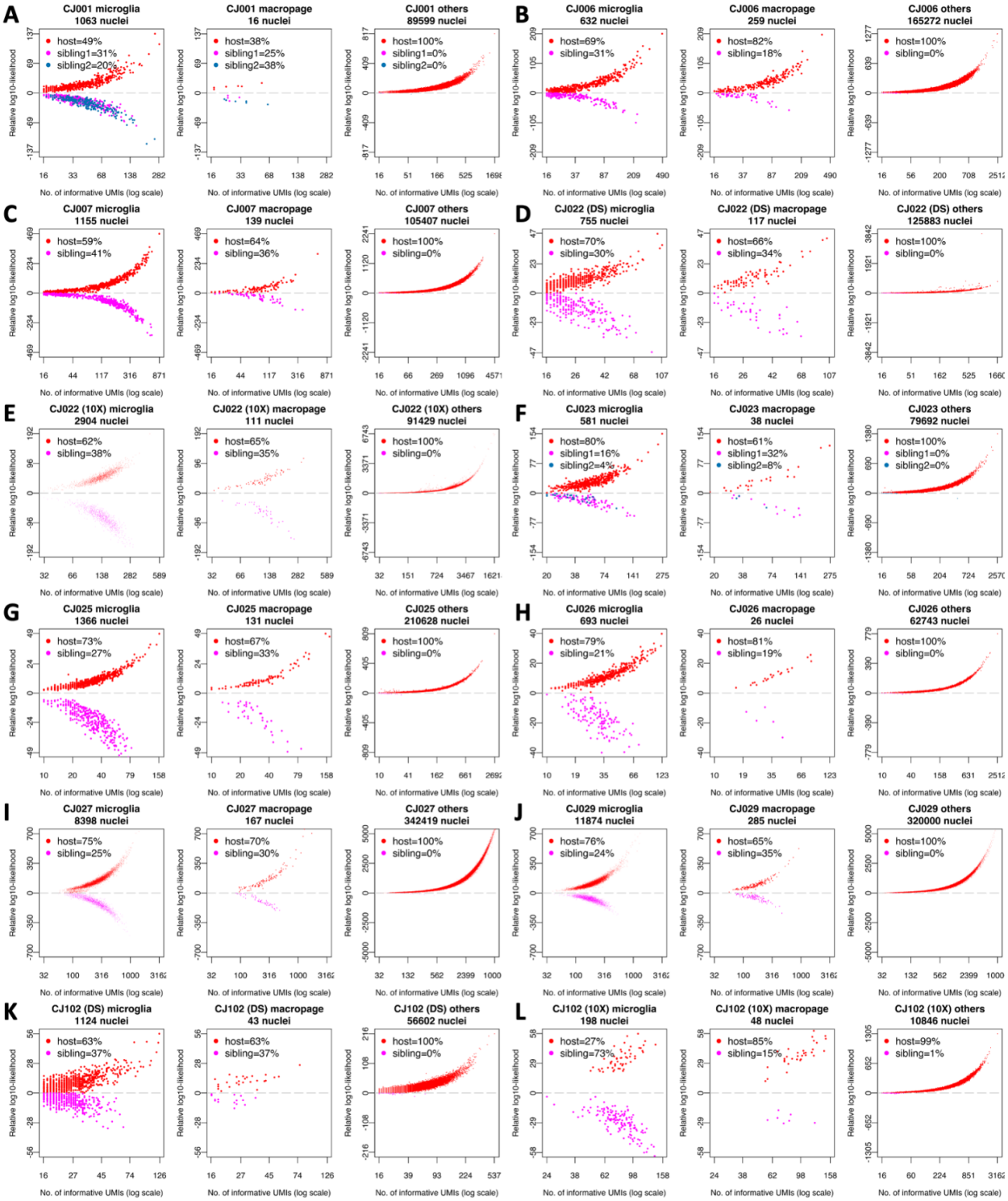


Supplementary Figure 1. Microglia and brain macrophages can be identified in all animals. Expression of microglia and macrophage markers in microglia (left sub-panels), macrophages (middle sub-panels) and all other cell types in the brain (right sub-panels; neurons, glia (astrocytes, oligodendrocytes, polydendrocytes, ependymal cells), and endothelial cells of each animal. Marmosets CJ022 and CJ102 were profiled using two technologies (DS: Drop-seq, 10X: 10X Chromium). y-axis: unique molecular identifier (UMI, a measure of transcript abundance) of each gene across cells, summed and normalized to 100,000 transcripts. Microglia markers: *TREM2, LAPTM5, C3*; macrophage markers: *F13A1, MSTN, LYVE1*.



Supplementary Figure 2. Donor-of-origin assignments from brain snRNA-seq reveals only microglia and macrophages are chimeric. (A-L) Donor of origin (Dropulation) assignments of each nucleus from brain snRNA-seq of 10 animals. Marmosets CJ022 and CJ102 were profiled using two technologies (DS: Drop-seq, 10X: 10X Chromium). For each marmoset, the snRNA-seq data are grouped into microglia, macrophage, and all other cell types (others: neurons, astrocytes, oligodendrocytes, polydendrocytes, ependymal cells, endothelial

772 cells). x-axis: number of UMI that contains SNPs for which the host and sibling's genomes differ,
773 in log scale; y-axis: inferred likelihood that the cell has host genome minus likelihood that the
774 cell has sibling genome (log10). Nuclei with positive y-values are assigned to the host and
775 those on the negative y-axes are assigned to the sibling.
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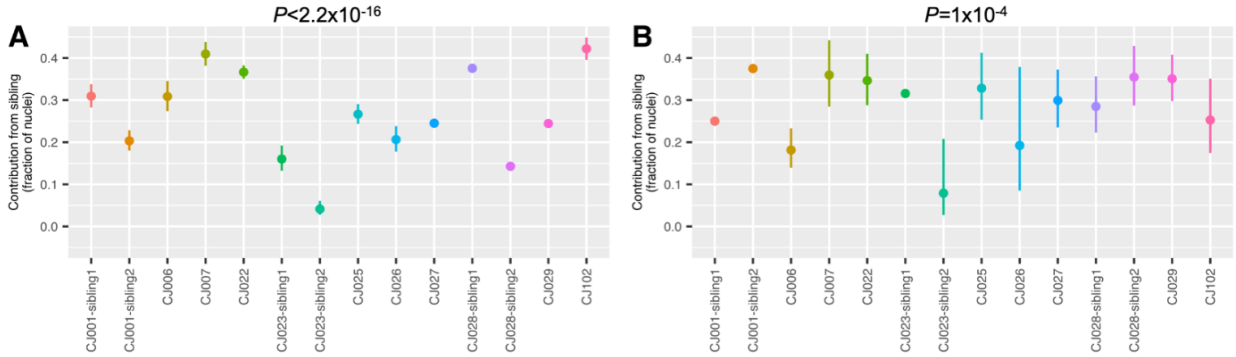
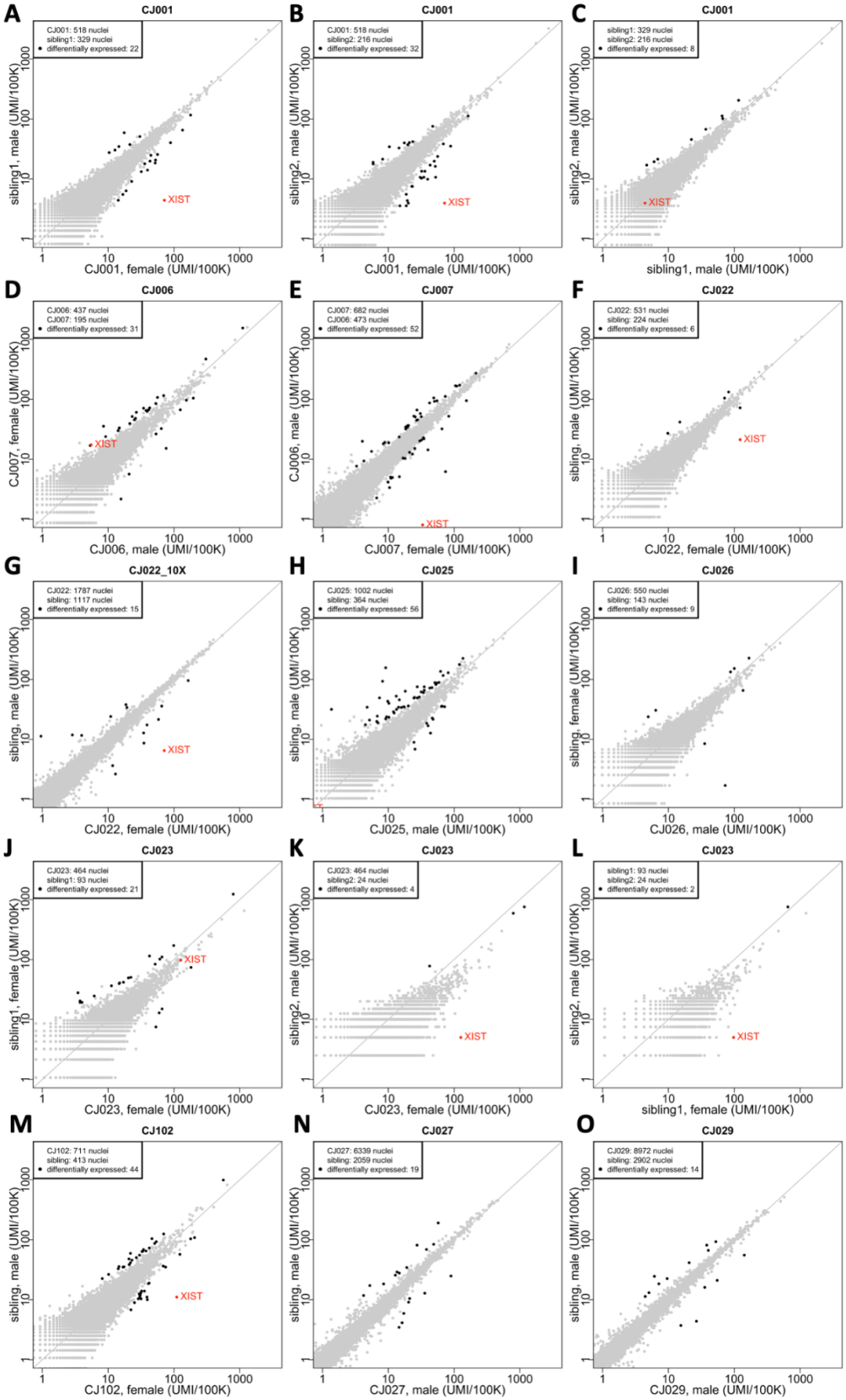
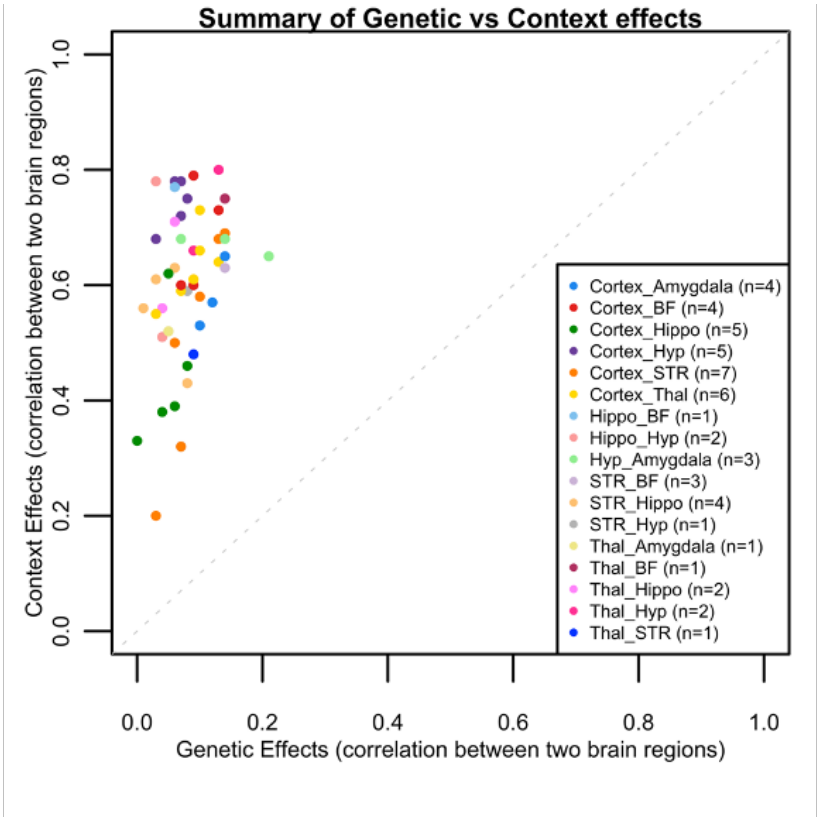


Supplementary Figure 3. Summary of microglia (A) and macrophage (B) chimerism across animals. y-axis: fraction of twin cells. Vertical bars: binomial confidence interval (95%). *P*-values: test of heterogeneity across animals.

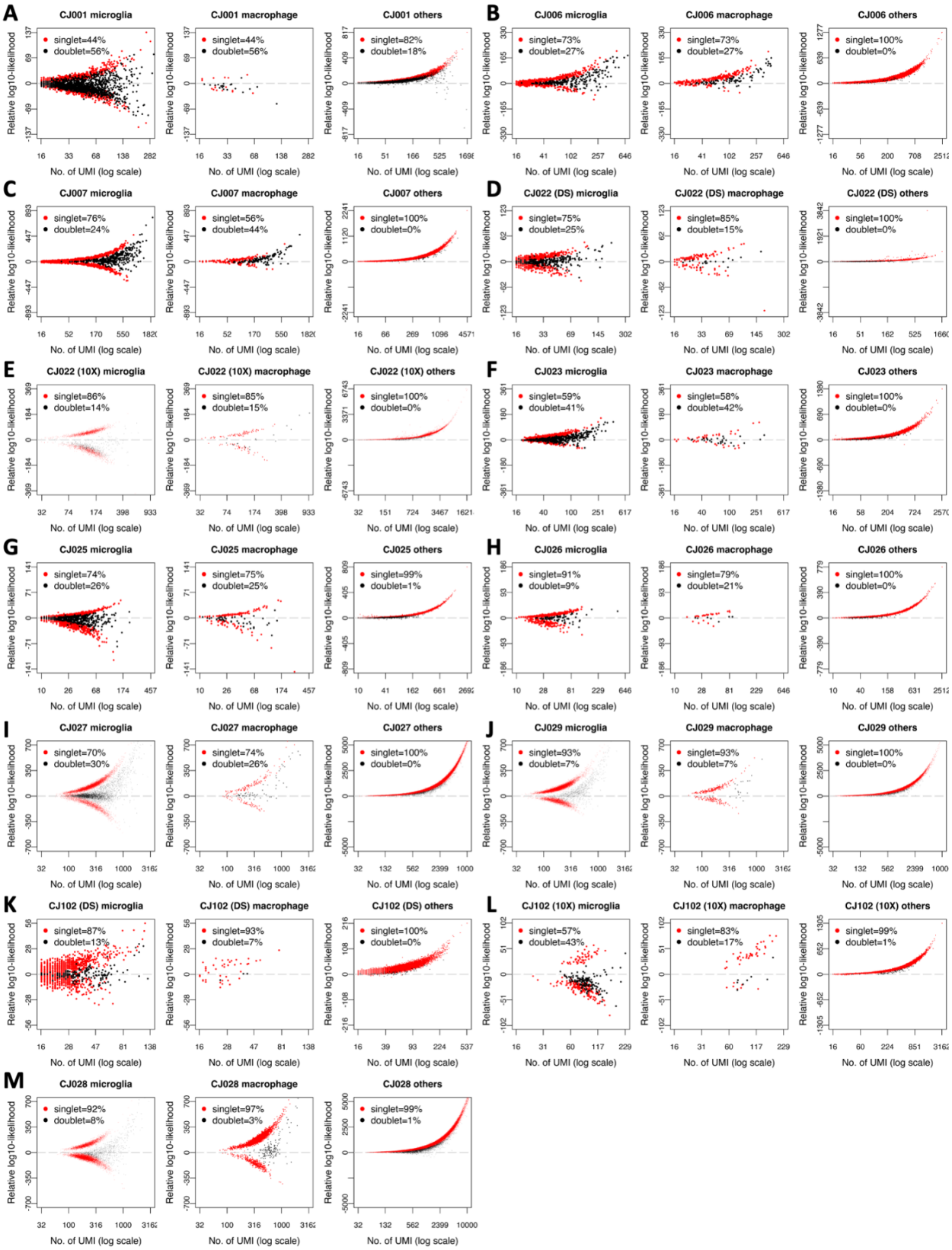


Supplementary Figure 4. Comparison of gene expression between microglia with

different genomes in each host animal's brain. Each point represents a gene; its location on the plot represents the level of expression of that gene among microglia with two different genomes in the same animal. x- and y-axes: normalized gene expression levels (number of transcripts per 100,000 transcripts). Fold-change and *P*-values were calculated using edgeR and differentially expressed genes (black dots) were defined as: FDR *Q*-value<0.05 and fold-change>1.5 or less than 1/1.5 and the gene must be expressed in at least 10% of one of the microglia sets.

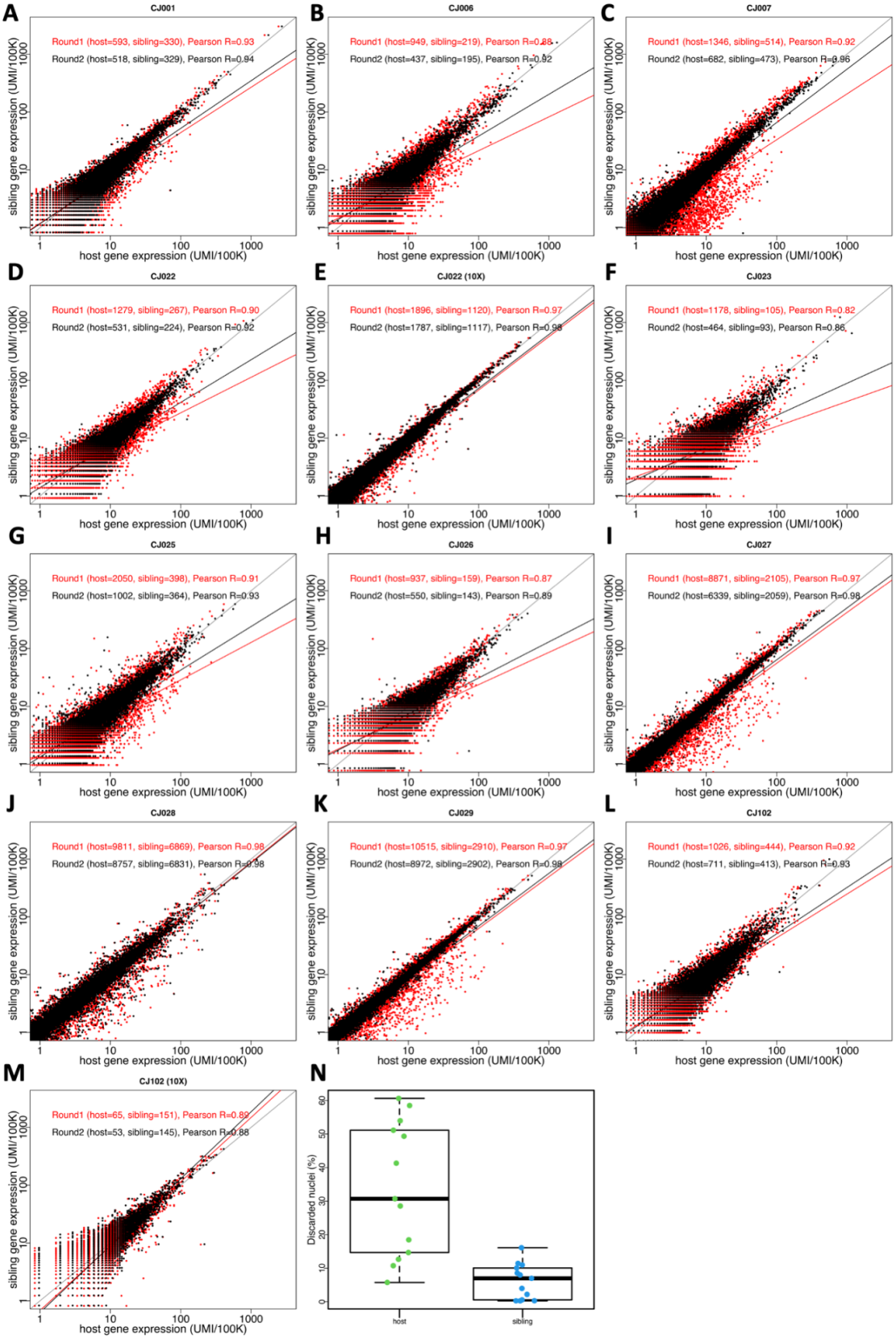


Supplementary Figure 5. Summary of genetic versus context effects. This is a plot of the correlation values from Supplementary Table 5. Abbreviations; STR: striatum; Thal: thalamus; Hippo: hippocampus; Hyp: hypothalamus; BF: basal forebrain



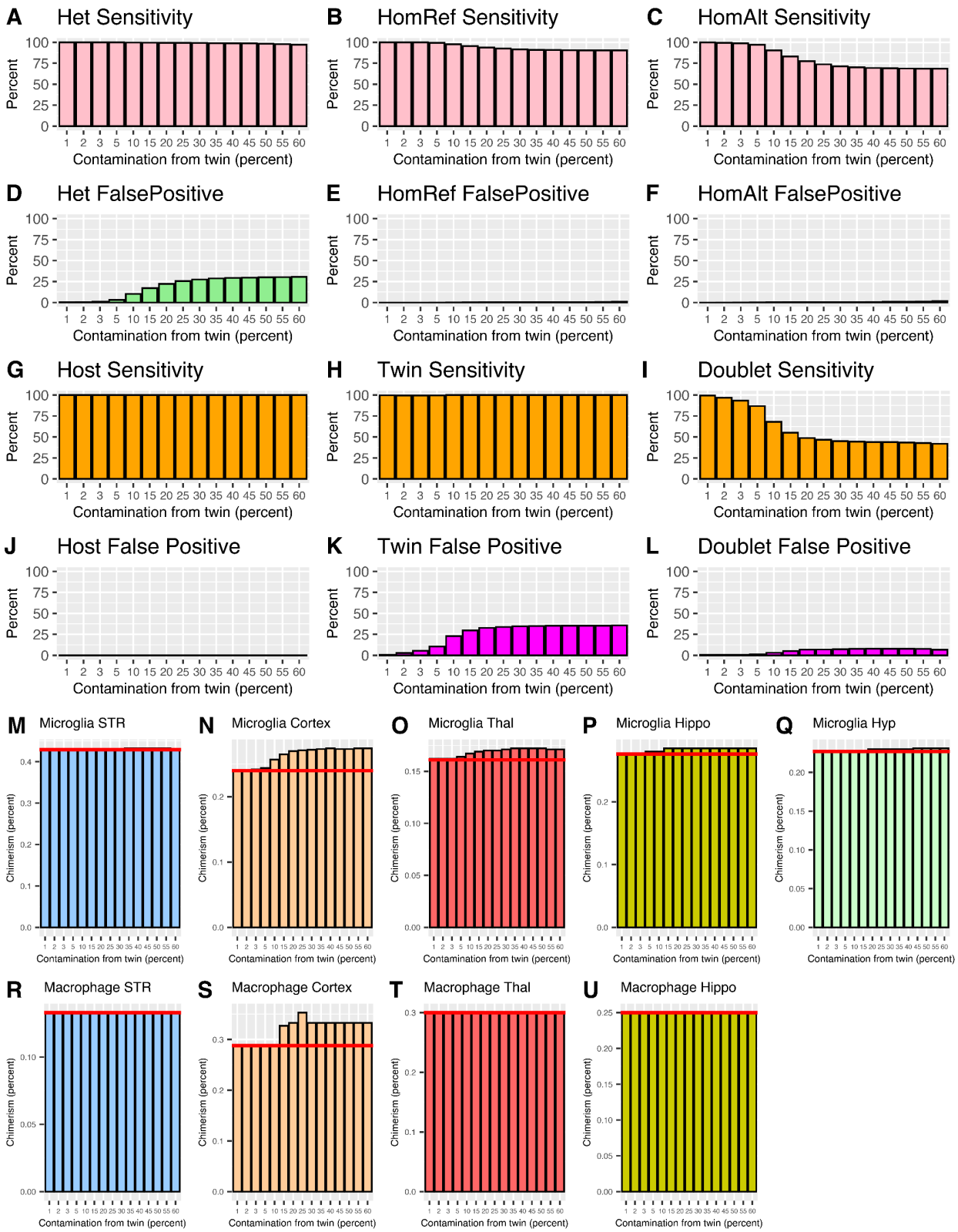
Supplementary Figure 6. Doublet detection using host and sibling genotypes. The axes are the same as in Supplementary Fig. 2, and each dot is a nucleus. Here, nuclei that were

identified as doublets and discarded in analyses were indicated (black dots). Marmosets CJ022 and CJ102 were profiled using two technologies (DS: Drop-seq, 10X: 10X Chromium).



Supplementary Figure 7. Second round clustering of microglia to discard mis-classified

cells. (A-M) Gene expression comparison between host and sibling cells. Red dots: nuclei identified as microglia from first-round of clustering, black dots: nuclei that were retained after second-round clustering. For triplets, only the first sibling is included in the plots. Pearson correlation as calculated for each set (before and after second round clustering) and shows an improvement in correlation after discarding mis-classified cells. Marmosets CJ022 and CJ102 were profiled using two technologies (Drop-seq and 10X Chromium). (**N**) Summary (box plot) of fraction of microglia cells discarded during second round of clustering, for host and birth sibling.



Supplementary Figure 8. Analysis of genotyping and chimerism if the genotypes of the sibling are contaminated by the hosts' DNA. (A)-(F) Sensitivity and false positives in

genotyping; HomRef: homozygous reference, HomAlt: homozygous alternate allele, Het: heterozygous. **(G)-(L)** Sensitivity and false positives in donor-of-origin assignment. **(M)-(Q)** Microglia chimerism estimates when sibling WGS are contaminated by the hosts' DNA, for 5 brain regions; red horizontal line: chimerism estimates when there's no error in sibling genotypes. **(R)-(U)** Macrophage chimerism estimates when sibling WGS are contaminated by hosts' DNA; red horizontal line: chimerism estimates when there's no error in sibling genotypes.