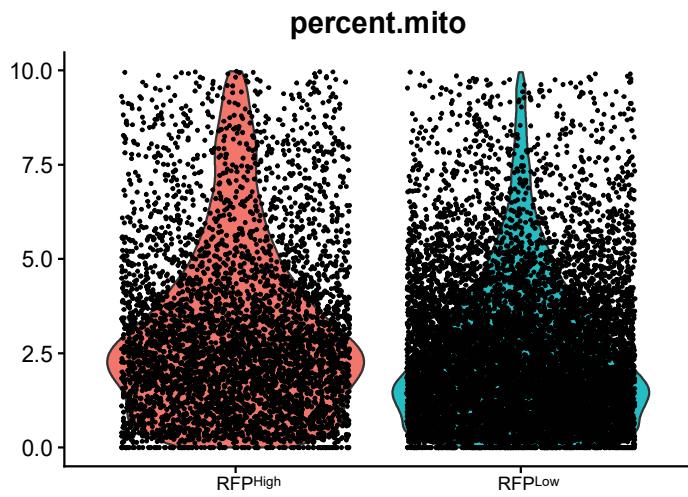


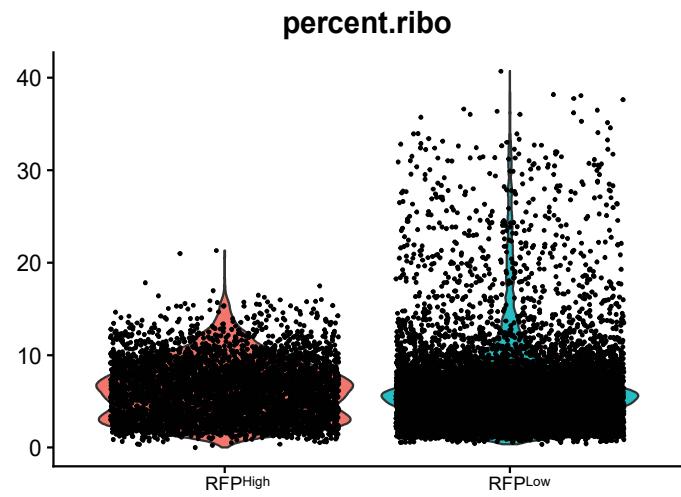
**Figure S1: WGCNA module expression between cell types.**

A: Gating strategy for sorting RFP<sup>High</sup> cells by FACS. B: p16 expression measured by qPCR in old mouse brains. C: List of significant weighted gene correlation network analysis (WGCNA) modules between RFP<sup>Low</sup> and RFP<sup>High</sup> samples. D-G: ModuleEigengene expression levels and gene ontology analysis of WGCNA modules enriched in RFP<sup>Low</sup> samples. H: UMAPs depicting the expression of WGCNA modules in all isolated cell types in single cell data set.

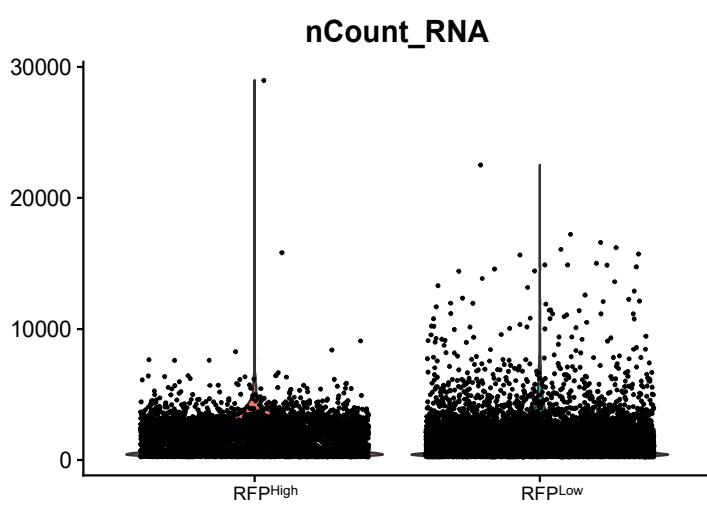
A



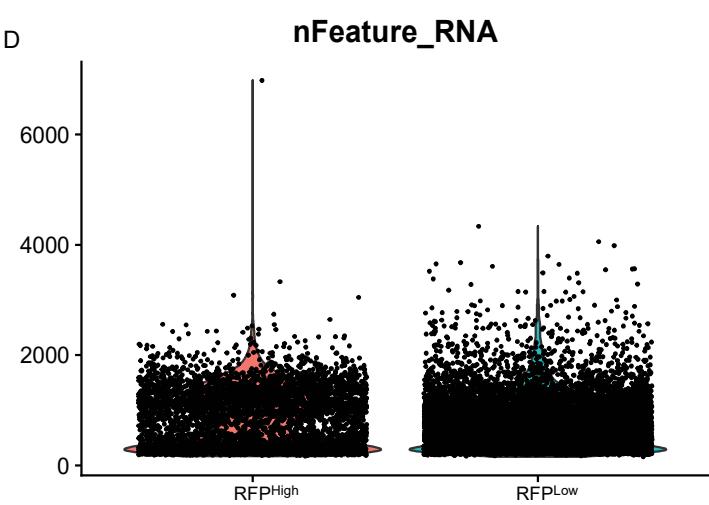
B



C



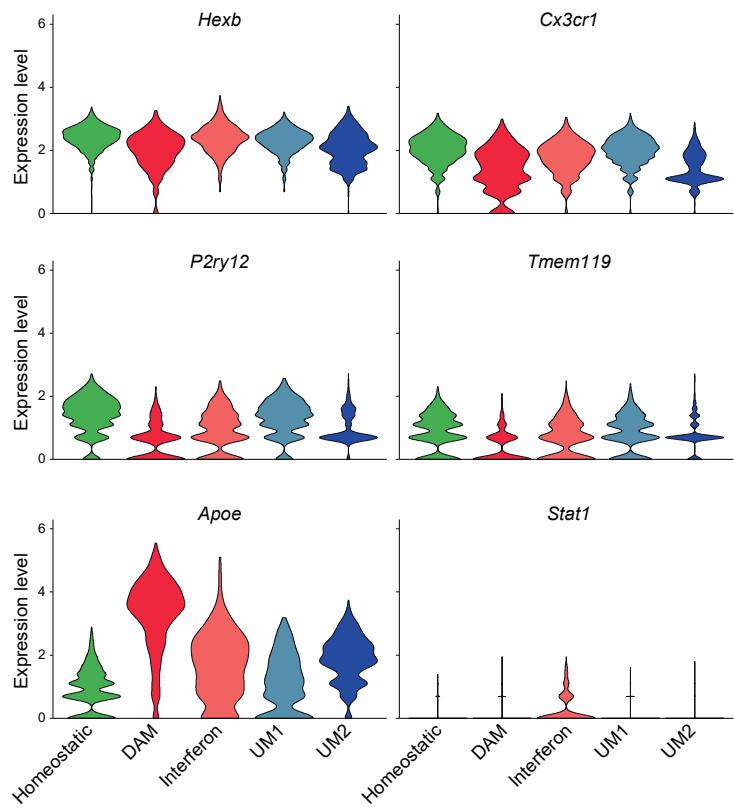
D



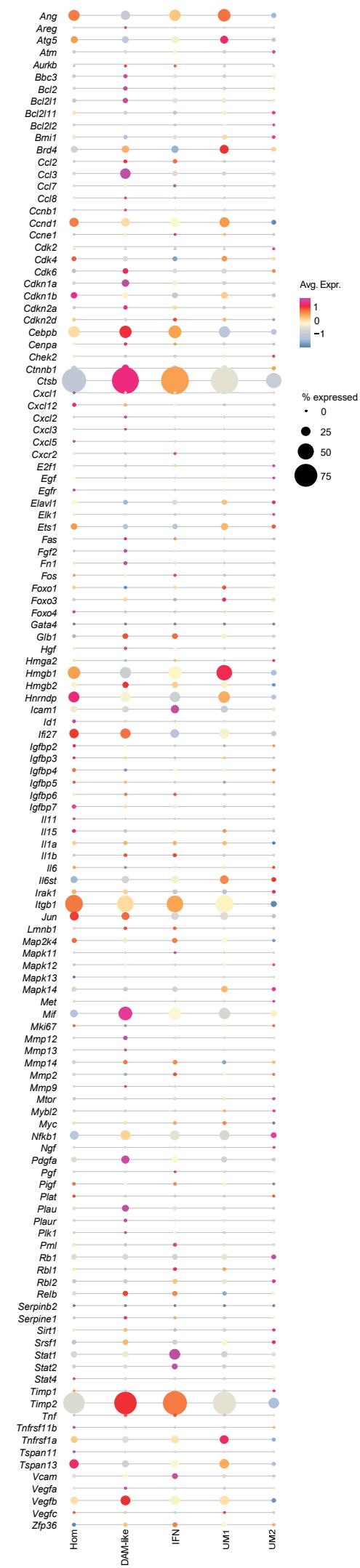
**Figure S2: Quality control of scRNaseq**

A: Percentage mitochondrial RNA, B: percentage ribosomal RNA, C: read counts and D: number of unique genes detected in RFPLow and RFPHigh samples.

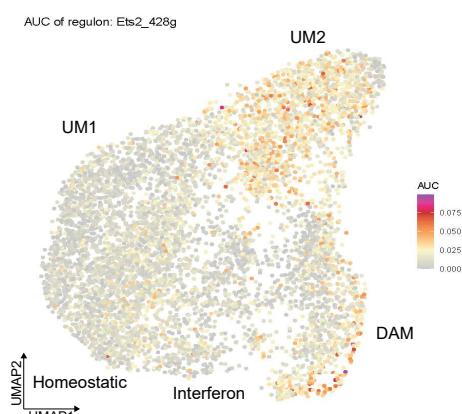
A



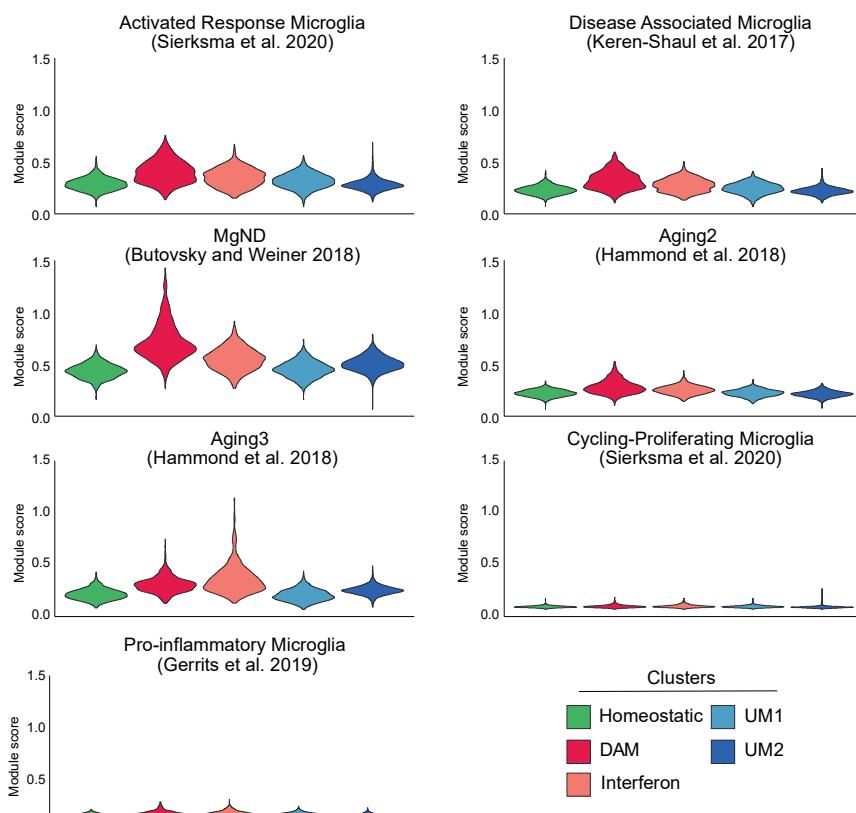
D



B



C



**Figure S3: Microglia clusters in literature do not resemble p16<sup>pos</sup> microglia.**

A: Violin plot depicting the expression of Hexb, Cxcr1, P2ry12,Tmem119, Apoe and Stat1 respectively in each cell of the microglia cluster. B: UMAP depicting Ets2 regulon expression in all microglia. C: Violin plots showing the expression of subtype signatures from the literature in each cluster. D:Dot plot showing expression of senescence genes in microglia clusters.