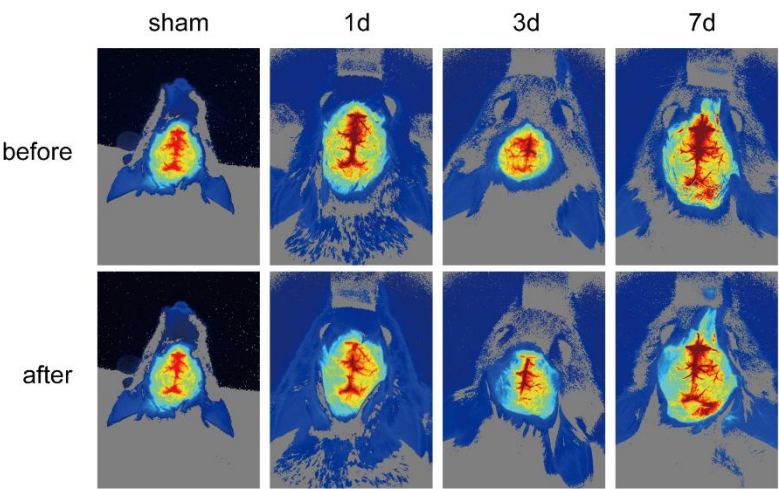
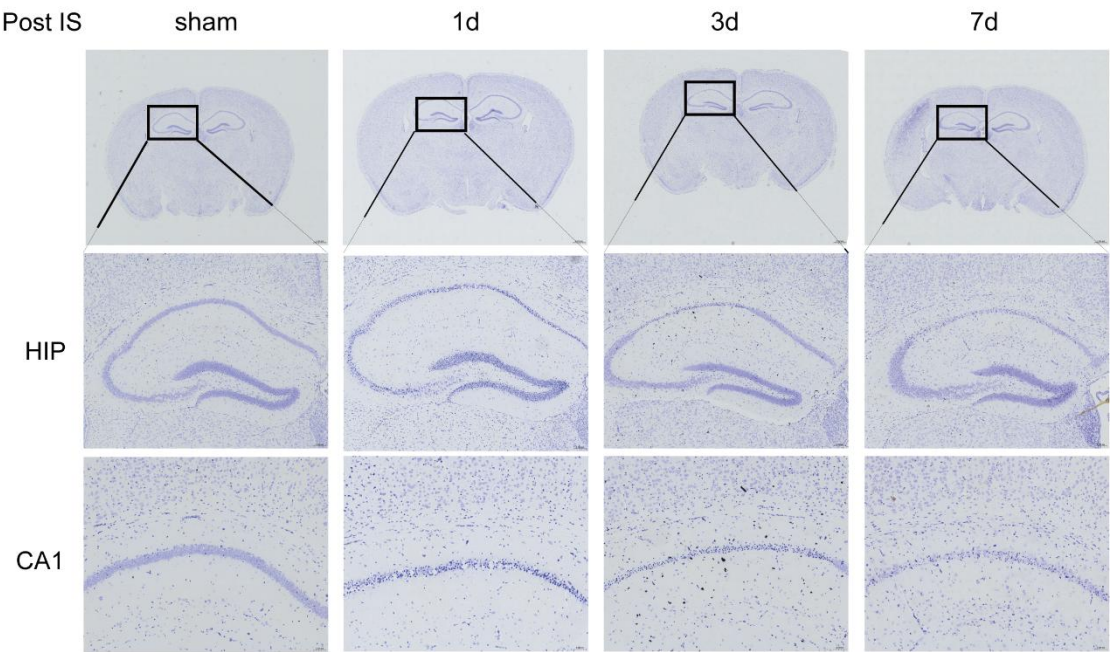


1 **Supplementary materials**

A



B

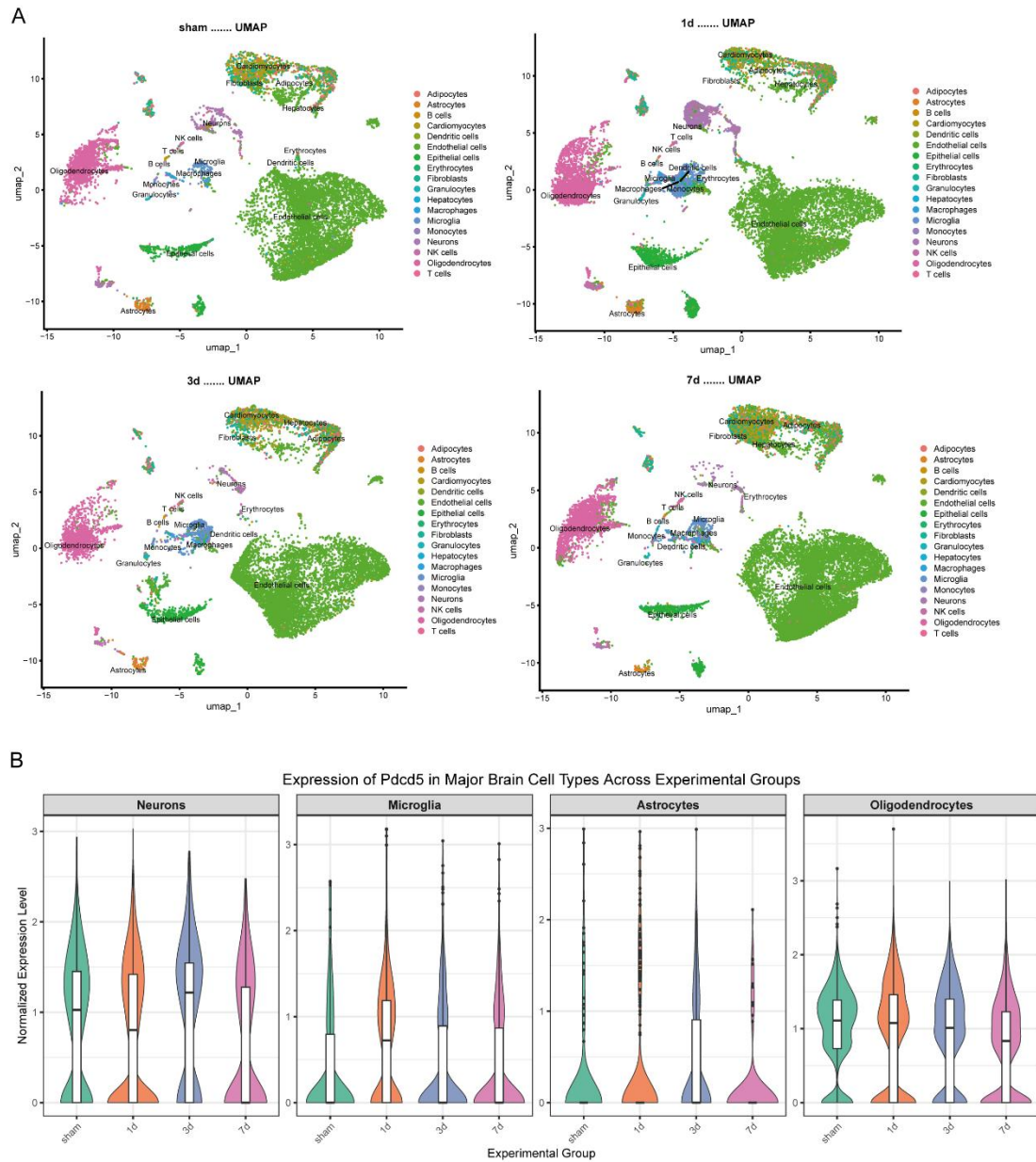


2
3 **Supplementary Figure S1. Validation of the transient MCAO mouse model and**
4 **post-ischemic histological alterations**

5 **(A)** Representative laser speckle contrast images showing cerebral blood flow before and
6 after surgery in sham mice and mice at 1, 3, and 7 days following MCAO.

7 **(B)** Representative Nissl-stained coronal brain sections from sham and ischemic mice at

8 the indicated time points post-ischemic stroke (Post IS). Upper panels display overall
 9 brain morphology; lower panels show magnified views of the hippocampus (HIP) and CA1
 10 region from the boxed areas.

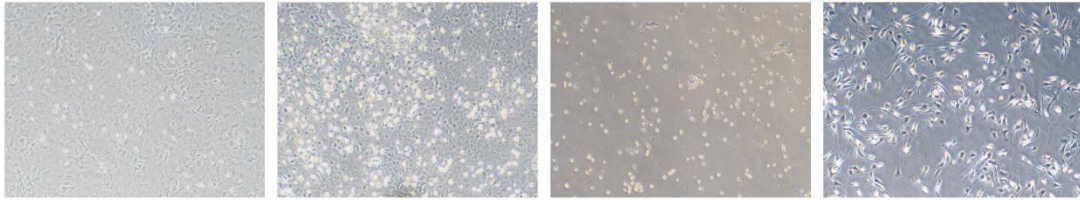


11
 12 **Supplementary Figure S2. Cell type annotation of single-cell transcriptomic data**
 13 **and cell-type-specific expression dynamics of *Pdcd5***
 14 **(A)** UMAP visualization of unsupervised cell clustering and cell type annotation in brain
 15 tissues from sham mice and mice at 1, 3, and 7 days after MCAO. Each dot represents an

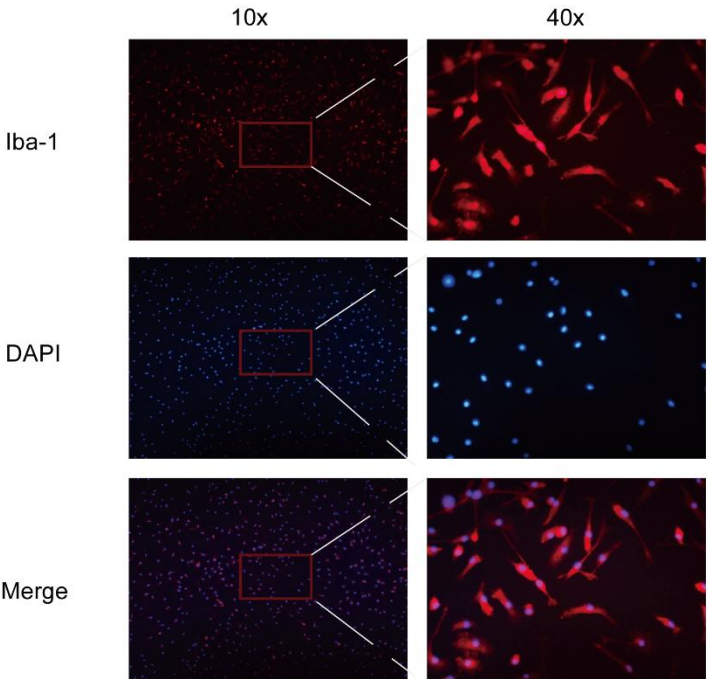
16 individual cell, and distinct cell types are color-coded.

17 **(B)** Violin plots showing normalized expression levels of *Pdcd5* in four major brain cell
18 types (neurons, microglia, astrocytes, and oligodendrocytes) across the indicated
19 experimental time points.

A



B



20

21 **Supplementary Figure S3. Isolation, purification and identification of primary**
22 **mouse microglia**

23 **(A)** Representative bright-field micrographs showing cell morphology at different stages of
24 primary microglia isolation and purification.

25 **(B)** Immunofluorescence staining for the microglial marker Iba-1 (red) to verify the purity of
26 isolated microglia. Nuclei were counterstained with DAPI (blue). Images are presented at
27 10× and 40× magnifications; the boxed areas in the 10× fields are shown at higher

28 magnification in the 40× panels.

29