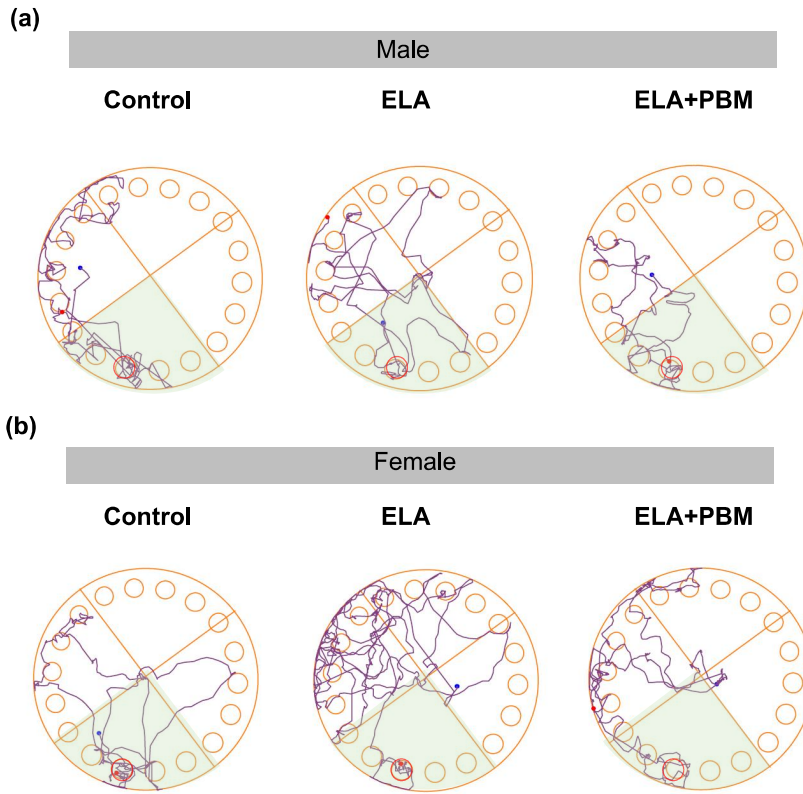
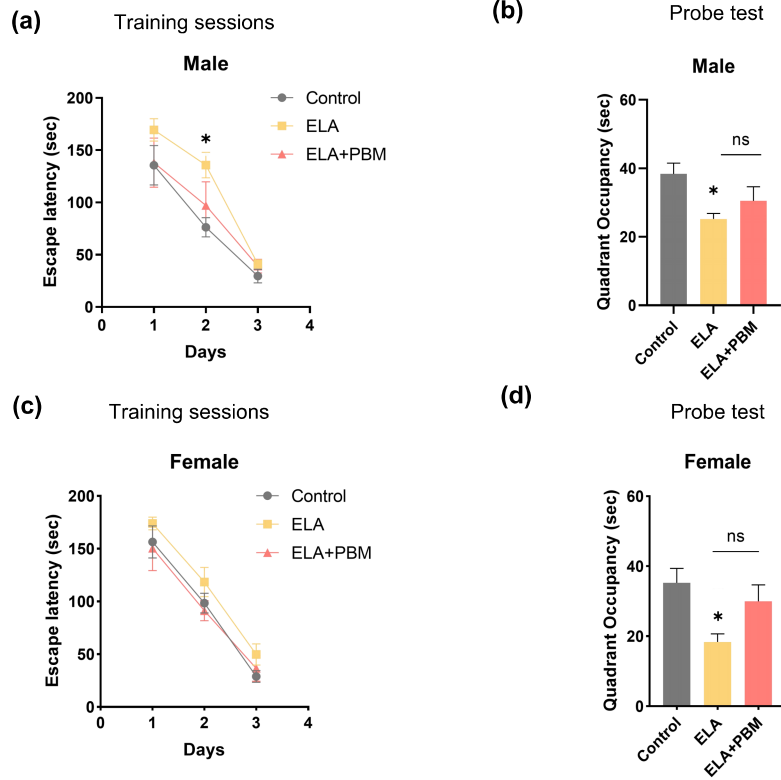


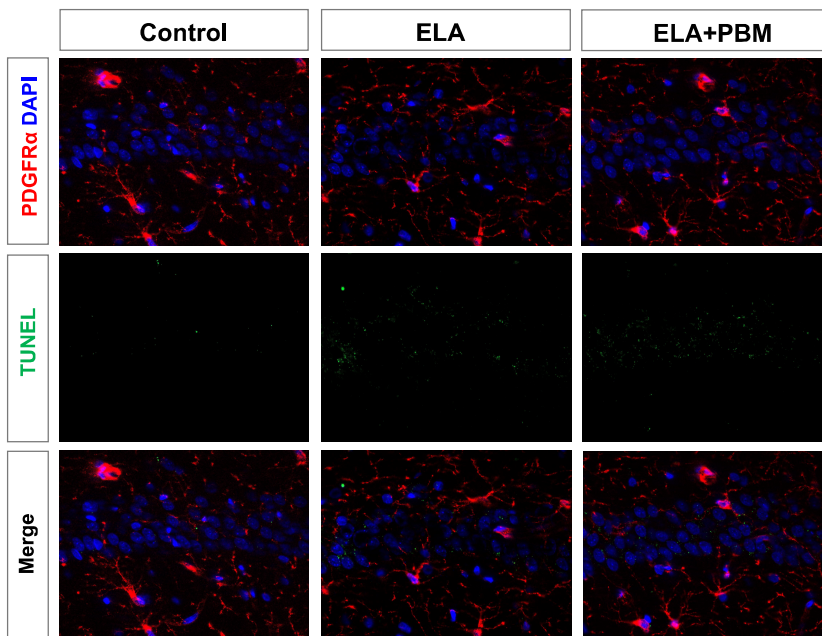
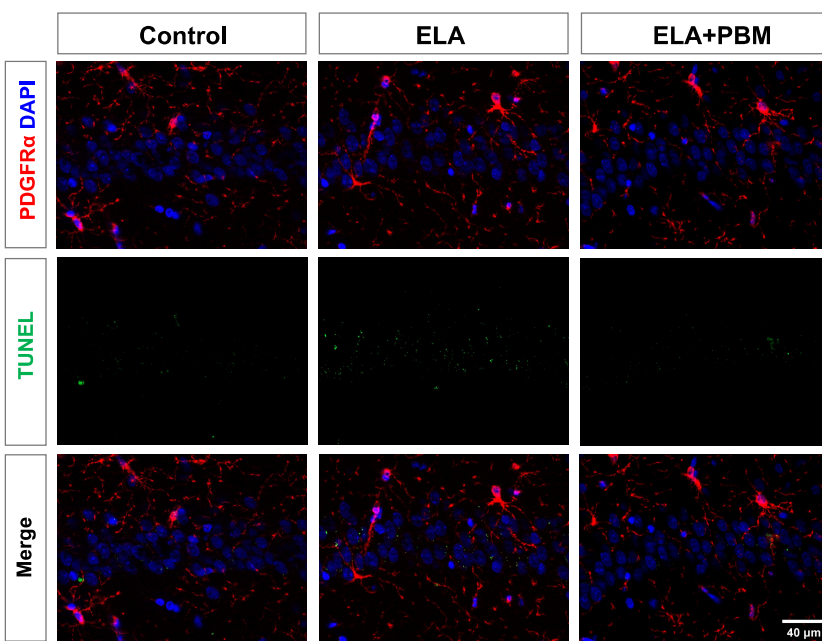
## **Supplementary Information**

**4 Supplementary Figures**

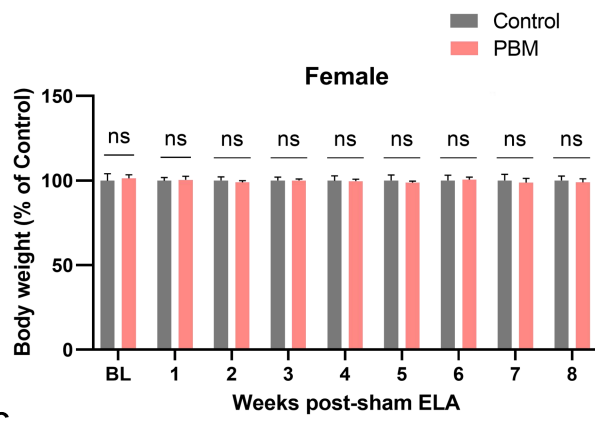
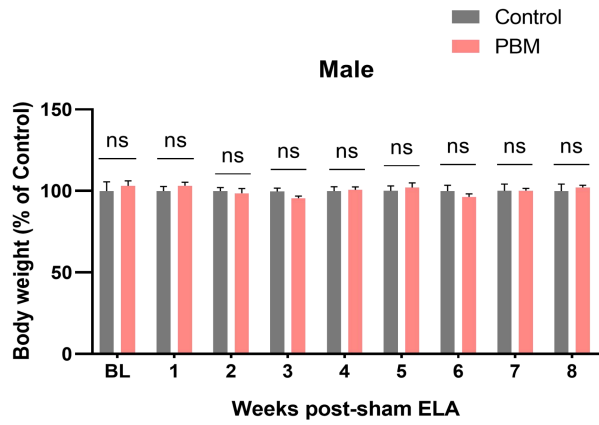
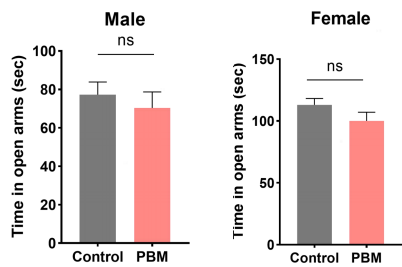
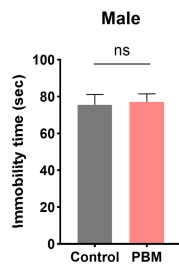
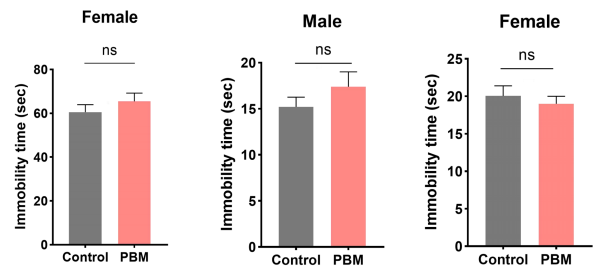
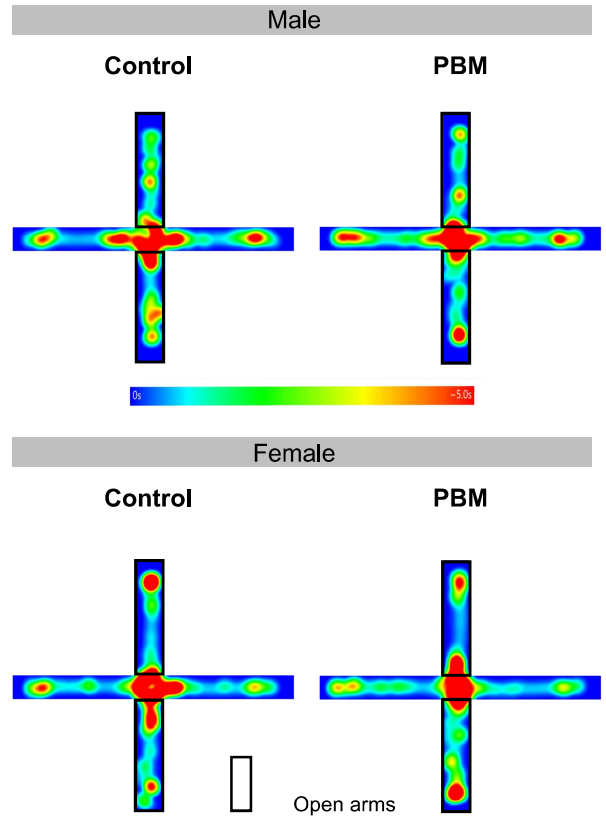
**1 Supplementary Table**

**A****B**

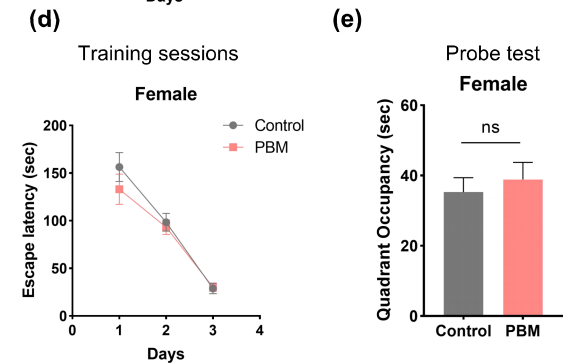
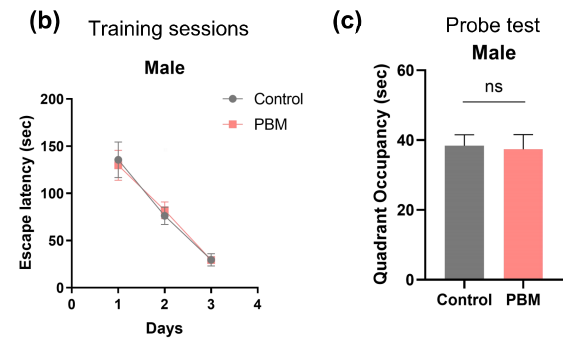
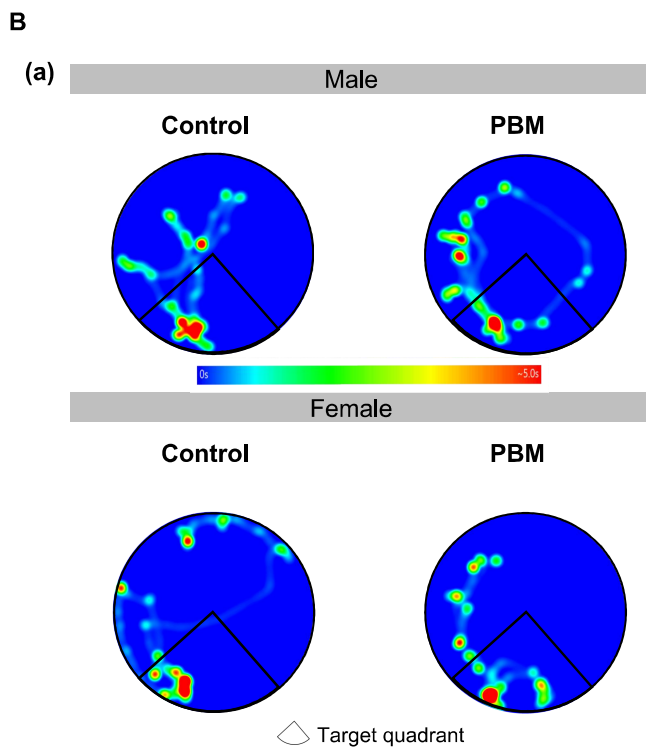
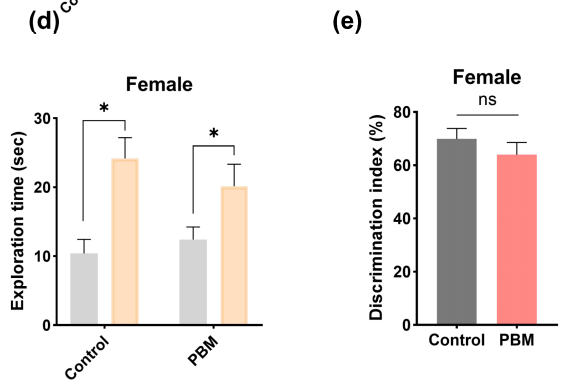
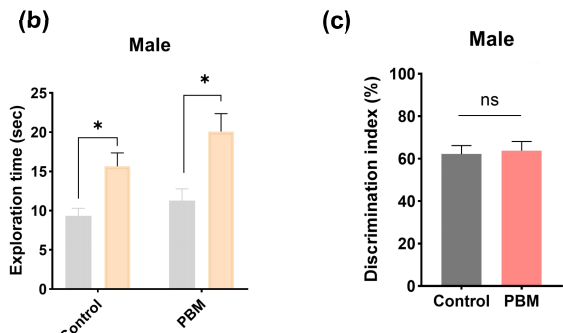
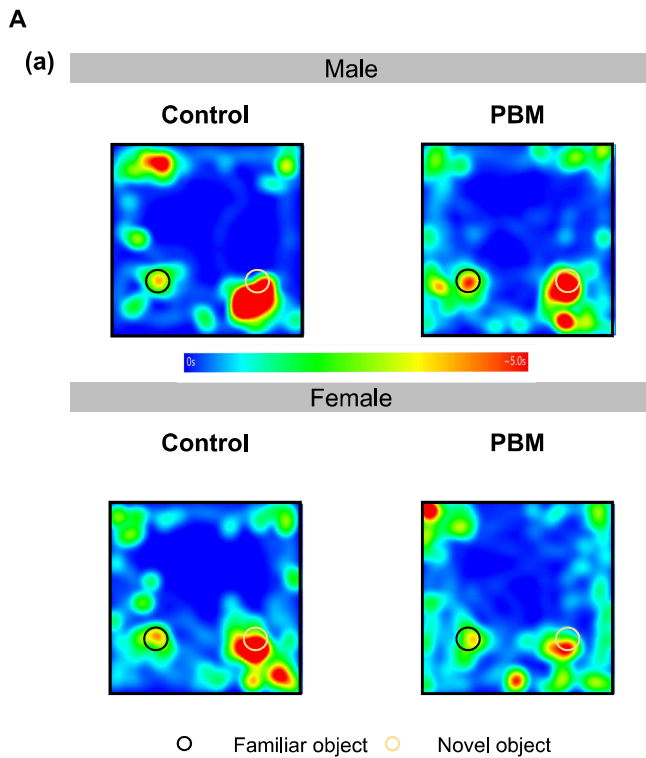
**Figure S1. ELA resulted in spatial memory impairment, which can be partially alleviated by early PBM treatment.** The Barnes maze task was used to assess spatial learning and memory abilities. **A (a, b)** The representative tracking plots of the Barnes maze task. **B** After training sessions, the probe trials were conducted, escape latency curves during the three days of training sessions (**a, c**) and quadrant occupancy time in the probe (**b, d**) were recorded and analyzed. Compared to control rats, ELA-exposed rats spent significantly less time in the target quadrant, and male rats also showed a deficit in spatial learning, early PBM treatment partially alleviated these deficits, but was not significant. All data are presented as mean  $\pm$  SE (n = 5-7). \* P < 0.05 versus Control-group; # P < 0.05 versus ELA-group.

**A****Male****B****Female**

**Figure S2. ELA does not compromise the survival of the OPCs.** TUNEL staining and immunostaining for PDGFR $\alpha$  (a marker of OPCs) with DAPI. In both male (**A**) and female animals (**B**), no appreciable TUNEL<sup>+</sup> PDGFR $\alpha$ <sup>+</sup> cells were observed. Scale bar = 40  $\mu$ m (n = 5-7).

**A****C****(a)** Elevated plus maze**(b)** Tail suspension**(c)** Forced swimming**B**

**Figure S3. Early PBM treatment does not affect the levels of anxiety and depression in normal animals.** **A** No significant difference in body weight alternation between the two groups. **B** Representative heat maps of the elevated plus maze test. **C (a)** There was no difference between groups in the time spent in open arms. **C (b)** No differences in immobility time were observed in the tail suspension test. **C (c)** No differences in immobility time were observed in the forced swimming test. All data are presented as mean  $\pm$  SE (n = 6-7).





**Figure S4. Early PBM treatment does not affect cognitive performances in normal animals.** **A (a)** Representative heat maps of the novel object recognition test. **A (b, d)** The time spent on the familiar (dark) object and novel object (orange) was calculated and statistically compared. **A (c, e)** No differences in the discrimination index were observed between the two groups. **B (a)** Representative heat maps of the Barnes maze task. **B (b, d)** There was no significant difference in escape latency between the two groups. **B (c, e)** No significant differences in the quadrant occupancy time were observed between the two groups. All data are presented as mean  $\pm$  SE (n = 6-7).

**Table S1. Primary antibodies used in this study**

Antibody	Host organism	Vendor	Cat#	Dilution	Application
Olig2	Goat	R&D Systems	AF2418-SP	1:250	IF
MBP	Mouse	Abcam	ab62631	1:300	IF
CC1	Mouse	EMD Millipore	OP80	1:500	IF
Ki67	Rabbit	Thermo Fisher Scientific	PA5-19462	1:300	IF
4-HNE	Mouse	Japan Institute For the Control of Aging	MHN-100P	1:200	IF
PDGFR $\alpha$	Rabbit	Cell Signaling Technology	#3174	1:500 for IF, 1:50 for ProteinSimple®	IF, ProteinSimple® electrophoresis syetem
MBP	Rabbit	Proteintech	10458-1-A	1:50	ProteinSimple® electrophoresis syetem
Olig2	Rabbit	Abcam	ab109186	1:50	ProteinSimple® electrophoresis syetem