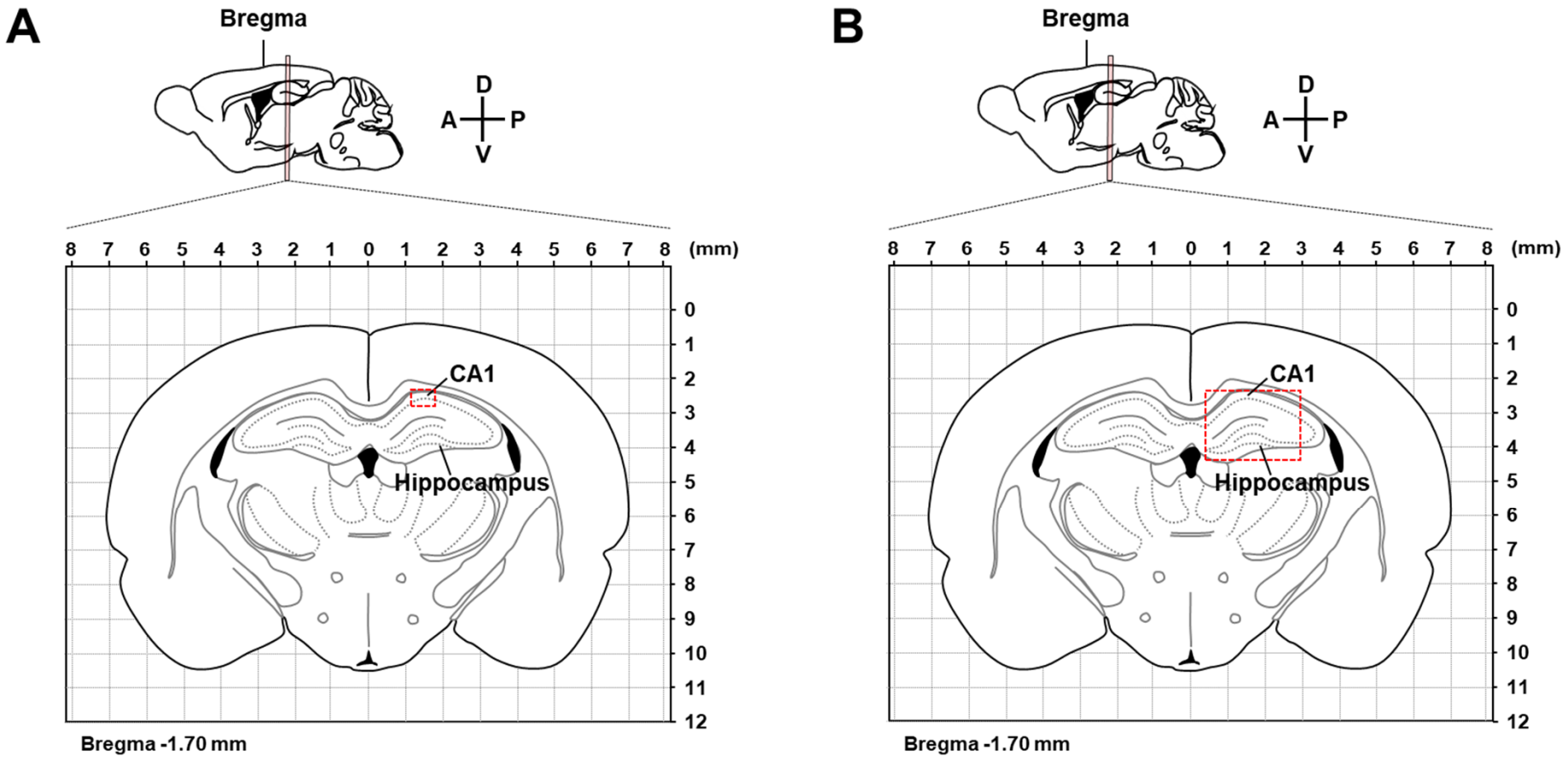

Figure S1. Schematic representation of brain sections used for immunostaining. (**A**) The coordinates of the hippocampal region of mice brain imaged under 20× objective for the study of CRAMP localization in different cell types. (**B**) The coordinates of the hippocampal region imaged under 5× objective to study the reactive gliosis in LPS-injected mice.

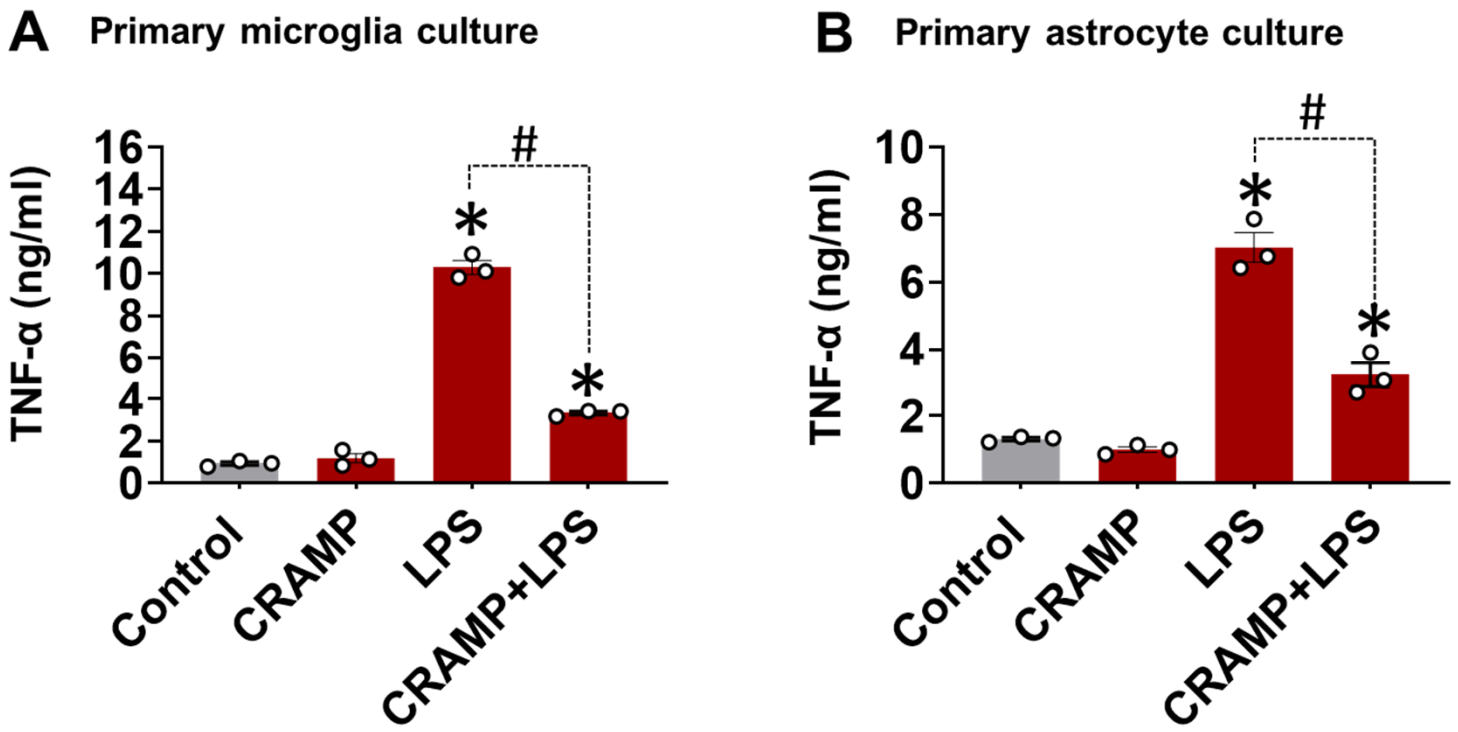

Figure S2. The effect of CRAMP peptide post-treatment on LPS-stimulated glial cells. The effect of CRAMP peptide (30 μg/mL) post-treatment on LPS-induced production of TNF-α was determined in primary microglia (**A**) and astrocyte culture (**B**). Data are mean ± SEM; n = 3; * $p < 0.05$ versus control; # $p < 0.05$ between the specified groups. One-way ANOVA with Tukey's post hoc test.

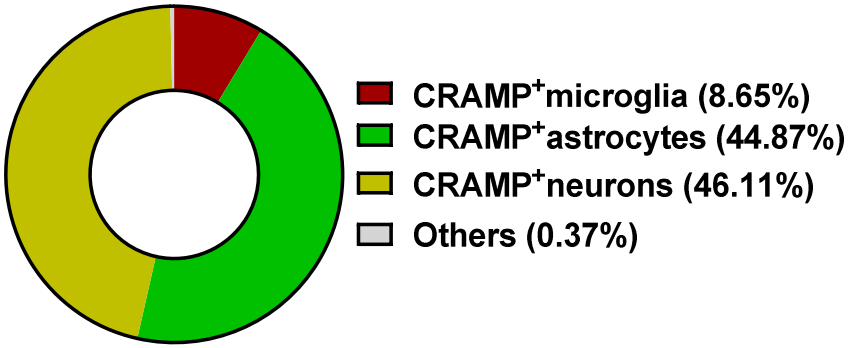

Figure S3. Cellular contribution of CRAMP expression in the hippocampus of LPS-injected mice.