SUPPLEMENTAL INFORMATION

Melanocortin receptor activation alleviates amyloid pathology and glial reactivity in an Alzheimer’s disease transgenic mouse model

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**Supplemental Figure 1.** Activation of melanocortin signaling ameliorates the amyloid pathology in the cortex in APP/PS1 mice. (A–B) Chronic D-Tyr administration reduces amyloid plaque deposition in APP/PS1 mice. Representative bright-field photomicrographs (A) and quantification (B) of 6E10-stained amyloid plaques in the neocortex on coronal brain sections of APP/PS1 mice following treatment with chronic D-Tyr versus control (Veh). Scale bar = 1 mm (n = 8–9 mice per group, 4 sections per mouse at 30-µm intervals). (C–E) Chronic D-Tyr administration reduces soluble and insoluble amyloid-beta (Aβ) contents. Representative western blot (C) and quantification of soluble (D) and insoluble (E) Aβ levels in cortical homogenates from APP/PS1 mice (n = 8–9 mice per group). Full-length blots are presented at the end of Supplemental Information. As the blots were cut prior to hybridization with antibodies, membrane edges were outlined with solid black lines. (F) Quantitative assessment (i.e., ELISA) of the relative levels of Aβ₄₀ and Aβ₄₂ isomers in the soluble fraction from the cortex of APP/PS1 mice (n = 4–5 mice per group). Data are the mean ± SEM of all mice from
each group examined (*$p < 0.05$, **$p < 0.01$, ***$p < 0.001$ for chronic D-Tyr vs. Veh treatment; Student’s $t$-test).
Supplemental Figure 2. Activation of melanocortin signaling modulates inflammatory responses in the cortex in APP/PS1 mice. Quantitative real-time PCR assessment of the levels of inflammatory genes in the cortex in wild-type (WT) mice treated with control (Veh) and APP/PS1 mice treated with chronic D-Tyr versus Veh. All measurements are normalized to the level of β-actin and presented as the fold expression relative to the average of the WT-Veh group. Transcript levels of Il1β (A), Il6 (B), Icam1 (C), Aif1 (D), and Gfap (E). Quantitative assessment (i.e., ELISA) of the relative level of IL-1β (F) in the cortex. Representative western blot (G) and quantification of Iba1 (H) and GFAP (I) protein levels in cortical homogenates.
from WT and APP/PS1 mice. Full-length blots are presented at the end of Supplemental Information. As the blots were cut prior to hybridization with antibodies, membrane edges were outlined with solid black lines. Data are the mean ± SEM of all mice from each group (*n* = 4–5 mice per group; *p* < 0.05, **p** < 0.01, ***p*** < 0.001; one-way ANOVA with the Bonferroni post hoc test).
Supplemental Figure 3. Activation of melanocortin signaling reduces the reactivity of astrocytes in the cortex in APP/PS1 mice. (A–C) Chronic D-Tyr administration reduces GFAP expression in astrocytes in the cortex in APP/PS1 mice. Representative images (A), immunoreactivity in arbitrary units (A.U.) (B), and average domain areas (C) of GFAP in astrocytes in the cortical region on coronal brain sections of wild-type (WT) mice treated with control (Veh) and APP/PS1 mice treated with chronic D-Tyr versus Veh. Scale bar = 100 µm.
(\(n = 9\) mice per group; \(*p < 0.05, ***p < 0.001;\) one-way ANOVA with the Bonferroni post hoc test). (D, E) Chronic \(\text{d-Tyr}\) administration reduces the \(\text{C3}^+\) (complement component 3) A1 subtype of reactive astrocytes. Representative images (D) and quantification (E) of \(\text{C3}^+\text{GFAP}^+\) co-labeled astrocytes (white arrows) in the cortex in APP/PS1 mice treated with chronic \(\text{d-Tyr}\) versus Veh. Scale bar = 20 \(\mu\)m (\(n = 9\) mice per group; \(*p < 0.05;\) Student’s \(t\)-test).
Supplemental Figure 4. Activation of melanocortin receptor signaling mediated by D-Tyr administration does not affect microglial reactivity in the cortex in APP/PS1 mice. Representative immunostaining (A) and quantification of the density (B) of microglia (labeled with Iba1) in the cortical region on coronal brain sections of wild-type (WT) mice treated with control (Veh) and APP/PS1 mice treated with chronic D-Tyr versus Veh. Scale bar = 100 µm (n = 9 mice per group; ***p < 0.001 for WT vs. APP/PS1 mice receiving Veh treatment; one-way ANOVA with the Bonferroni post hoc test). (C, D) Chronic D-Tyr administration did not affect the association of microglia with amyloid plaques in the cortex in APP/PS1 mice. Representative images (C) and quantification (D) of amyloid plaque-associated Iba+ microglia in the cortical region in APP/PS1 mice treated with chronic D-Tyr versus Veh. Scale bar = 20 µm (n = 9 mice per group; Student’s t-test).
Supplemental Figure 5. Quantitative validation of differentially expressed transcripts in hippocampal slices from APP/PS1 mice. (A–F) Quantitative PCR showing the regulation of certain genes in acute hippocampal slices from control WT mice (Veh) and APP/PS1 mice treated with D-Tyr versus Veh. Transcript levels of *Itgam* (A), *Aif1* (B), *Gfap* (C), *Il6* (D), *Chmp3* (E), and *Zfp579* (F). All measurements are normalized to the level of β-actin and are presented as the fold expression relative to the average of the WT-Veh group. Data are the mean ± SEM of all mice from each group (n = 3–4 mice per group; *p < 0.05, **p < 0.01, ***p < 0.001; one-way ANOVA with the Bonferroni post hoc test).
Supplemental Information

Raw images of western blots

**Figure 1E**

![Western Blot Image]

- **Soluble Aβ**
- **Insoluble Aβ**
- **β-actin**

**Supplemental Figure 1C**

![Western Blot Image]

- **Soluble Aβ**
- **Insoluble Aβ**
- **β-actin**

↑ indicates band not included in main figures
Figure 2G

Supplemental Figure 2G

↑ indicates band not included in main figures