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In vivo Animal Models

Alzheimer's Disease



APP_{s1} Transgenic Mouse Model

The APP_{s1} mouse model overexpresses human APP751 with Swedish and London mutations under the control of the neuron specific murine Thy1 promoter.

- Early onset of brain pathology (3-4 months)
- · Progressive learning & memory impair-
- Early concomitant microgliosis & astrocytosis
- ments in the Morris water maze (MWM) · Progressive increase in amyloid plaque burden and CAA
- · Increased oxidative stress & altered cholesterol profile

Figure 1:

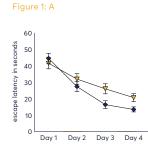
Morris water maze. Escape latencies of 6, 9 and 12 months old animals. Mean ± SEM; 6 and 9 months: n = 19 - 21; 12 months: n = 13 - 22; Two-way ANOVA with Bonferroni's post hoc test; *p<0.05, **p<0.01).

Figure 2:

Quantification of $A\beta$ levels in APPSL mice over age. Amount of Aβ1-38 (A) Aβ1-40 (B) and Aβ1-42 (C) in cortical (CTX) and hippocampal (HC) samples of 6-, 9- and 12-month old APPSL mice and 12-month old wild type (WT) littermates measured with MSD immunosorbent assay using the 4G8 antibody. n = 8 - 10 per group. Two-way ANOVA with Bonferroni's post hoc test. *p<0.05, **p<0.01; ***p<0.001.

Figure 3:

Qualitative comparison of plaque pathology of APPSL transgenic mice at 6, 9 and 12 month of age compared to non-transgenic littermates. Tissue was labeled with antibody 6E10 (green), collagen IV (red) and DAPI (blue).



Morris Water Maze

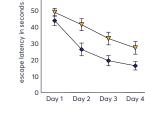




Figure 2: C

45.000

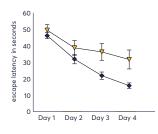
35.000

25.000

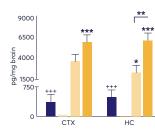
15.000

1.500

n/gq



Aß Expression Figure 2: A



стх

HC

500µm

Figure 3: B

6 months APP.

Figure 2: B

Figure 1: B

60

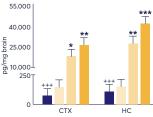
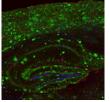


Figure 3: C

9 months APP,

Figure 3: D 12 months APP



СТХ

6E10 collagen IV DAPI

Figure 3: A

ntg

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126-54-ADAPPSL-EN (04/2024)



HC



Scantox Discovery