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A. Liver mRNA abundance of genes involved in liver lipid synthesis, fatty acid uptake, and lipid storage was similar between hepAGT^{-/-} mice and hepAGT^{+/+} mice fed on normal laboratory diet. N=3 to 9 for each group. Comparison between genotypes by Student's t-test.

B. Hepatic mRNA of genes involved in liver fatty acid uptake was not affected by hepatocyte-specific AGT deficiency in response to western diet. N=5 to 7 for each group. Comparison between genotypes by Student's t-test.

C. Hepatic mRNA abundance of CD36 was not affected by hepatocyte-specific AGT deficiency in response to western diet. N=5 to 7 for each group. Comparison among groups by One-Way ANOVA, Holm-Sidak post hoc test.

D. Hepatic protein abundance of CD36 was not affected by hepatocyte-specific AGT deficiency in response to western diet. N=5 for each group.

E. Hepatic AGT deletion suppressed western diet-induced hepatic *Ppar γ* expression. N=5 to 9 for each group. Comparison among groups by One-Way ANOVA, Holm-Sidak post hoc test.

F. Hepatic AGT deletion suppressed western diet-induced hepatic *Cide-a* expression. N=5 to 9 for each group. Comparison among groups by One-Way ANOVA, Holm-Sidak post hoc test.

G. Liver *Cide-b* mRNA abundance was not affected by absence of hepatocyte-derived AGT. N=5 to 9 for each group. Comparison among groups by One-Way ANOVA, Holm-Sidak post hoc test.

H. Hepatic AGT deletion suppressed western diet-induced hepatic *Cide-c* expression. N=5 to 9 for each group. Comparison among groups by One-Way ANOVA, Holm-Sidak post hoc test.

Ppar γ : Peroxisome proliferator-activated receptor γ . *Cide*: Cell death-inducing DFF45-like effectors.