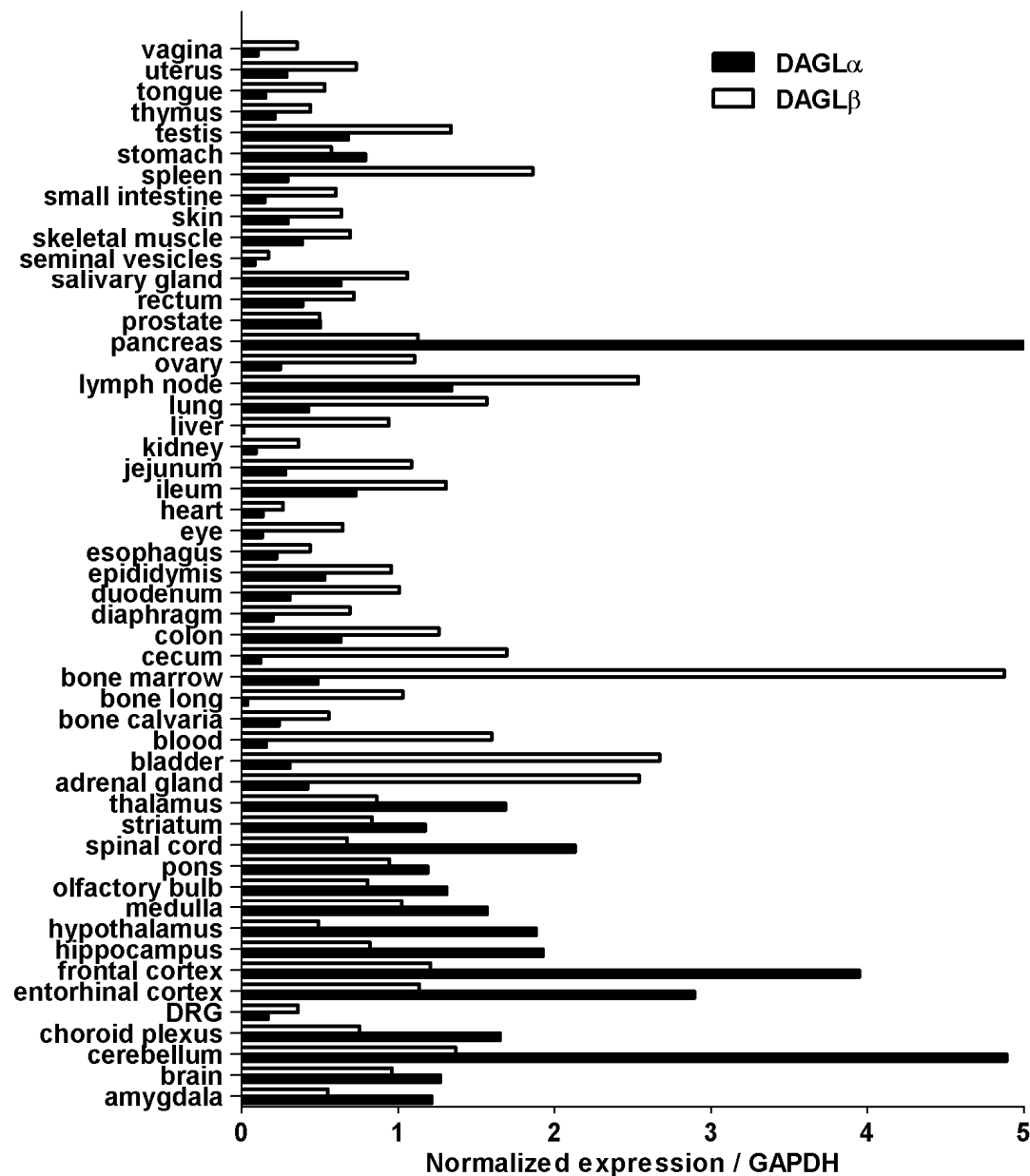


Supplementary Fig 1. Loss of either DAGL α or DAGL β has no impact on expression of other eCB pathway transcripts. Taqman analysis for mRNA levels of DAGL α , DAGL β , MAGL, FAAH, CB1, and CB2 in the cortex (CTX), cerebellum (Cb), spinal cord (SC), and liver (LI) from 3-5 wt (+/+), DAGL $\alpha^{+/-}$ ($\alpha^{+/-}$), DAGL $\alpha^{-/-}$ ($\alpha^{-/-}$), DAGL $\beta^{+/-}$ ($\beta^{+/-}$), and DAGL $\beta^{-/-}$ ($\beta^{-/-}$) mice. mRNA expression is normalised as the percentage of GAPDH levels.



Supplementary Fig 2. Tissue distribution of DAGL α/β transcripts. TaqMan qRT-PCR was performed on RNA samples extracted from multiple mouse tissue samples to detect DAGL α/β mRNA. Data were fitted to a standard curve generated with varying amounts of cDNA. Expression levels of DAGL α/β were normalized to GAPDH.