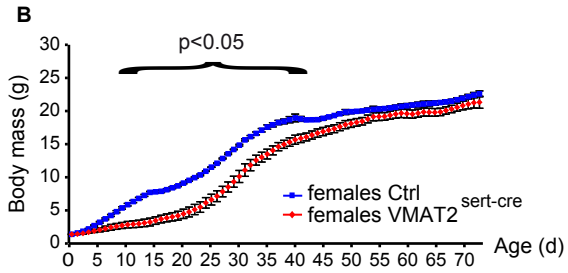
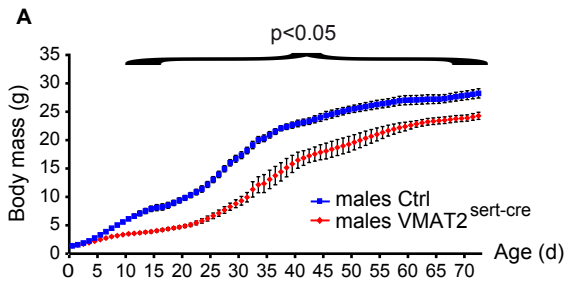
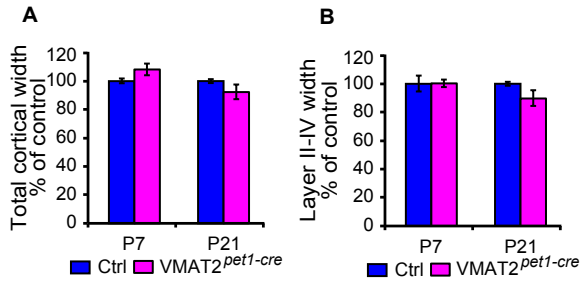


Supplementary Figure 1. Residual 5-HT in VMAT2^{sert-cre} mice.
 Control and VMAT2^{sert-cre} littermates were perfused at P7 and 50 μ m-thick coronal sections were labeled with 5-HT antibodies and revealed with DAB. A,B) In the raphe, 5-HT immunolabeling is completely absent in the axons and neuropil, while faint residual staining persists in the cell soma.
 C,D) In pinealocytes, 5-HT immunostaining remains strong in mutants, but neighboring raphe axons and neuropil are depleted.
 Scale bar = 100 μ m.



Supplementary Figure 2. Gender differences in growth rates in VMAT2^{sert-cre} mice

Mean growth curves from 11 VMAT2^{sert-cre} (red diamonds) and 7 control male mice (blue squares) (A) and 7 VMAT2^{sert-cre} (red diamonds) and 7 control female mice (blue squares) (B) from 6 different litters. The growth of the VMAT2^{sert-cre} mice was significantly different from that of control mice from P11 to P75 for males, and from P9 to P41 for females (ANOVA p<0.05).



Supplementary Figure 3: Cortical growth in VMAT2^{pet1-cre} mice

A,B) The cerebral cortex of VMAT2^{pet1-cre} mice was measured at P7 (n=4 controls and 5 VMAT2^{pet1-cre}) for total (A) or superficial layers (B) thickness in the somatosensory cortex. No difference was found between control and recombined animals (Student test). Measures from P21 brains (n=4 for each genotype) are show no difference as well. Values are expressed as % of control values.