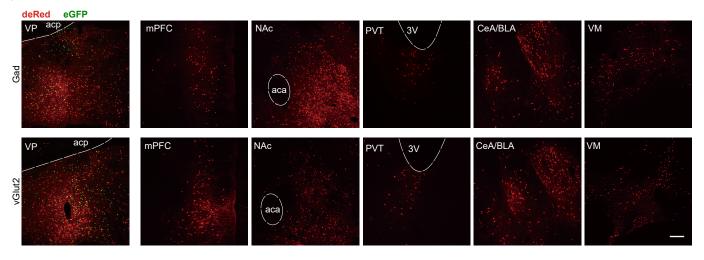
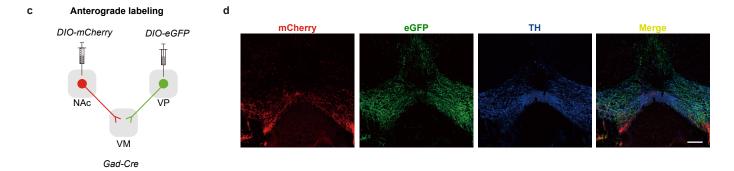


b





Supplementary information, Fig. 10 Rabies virus-based retrograde labeling of GABAergic or glutamatergic neurons in the VP.

a, Schematic of experimental design. AAV9-EF1α-DIO-his-eGFP-2α-TVA and AAV9-EF1α-DIO-RVG were unilaterally injected into the VP of Gad-Cre or vGlut2-Cre mice. Two weeks later, RV-ENVA-deltaG-dsRed (RVdG) was injected into the same VP site. Eight days were allowed for rabies to transduce and label synaptic connected input neurons. b, Representative confocal images of rabies labeling of given regions inputs to VP GABAergic or glutamatergic neurons from Gad-Cre or vGlut2-Cre mice, respectively. Scale bar: 100 μm. c, Schematic of anterograde labeling of GABA NAc-VM and GABAVP-VM projections. AAV9-EF1α-DIO-mCherry and AAV9-EF1α-DIO-eGFP were bilaterally injected into the NAc and VP of Gad-Cre mice. d, Representative images of mCherry⁺ and eGFP⁺ projectors in the VM with immunostaining of TH⁺ neurons. Scale bar: 100 μm. Related to Figure 5.