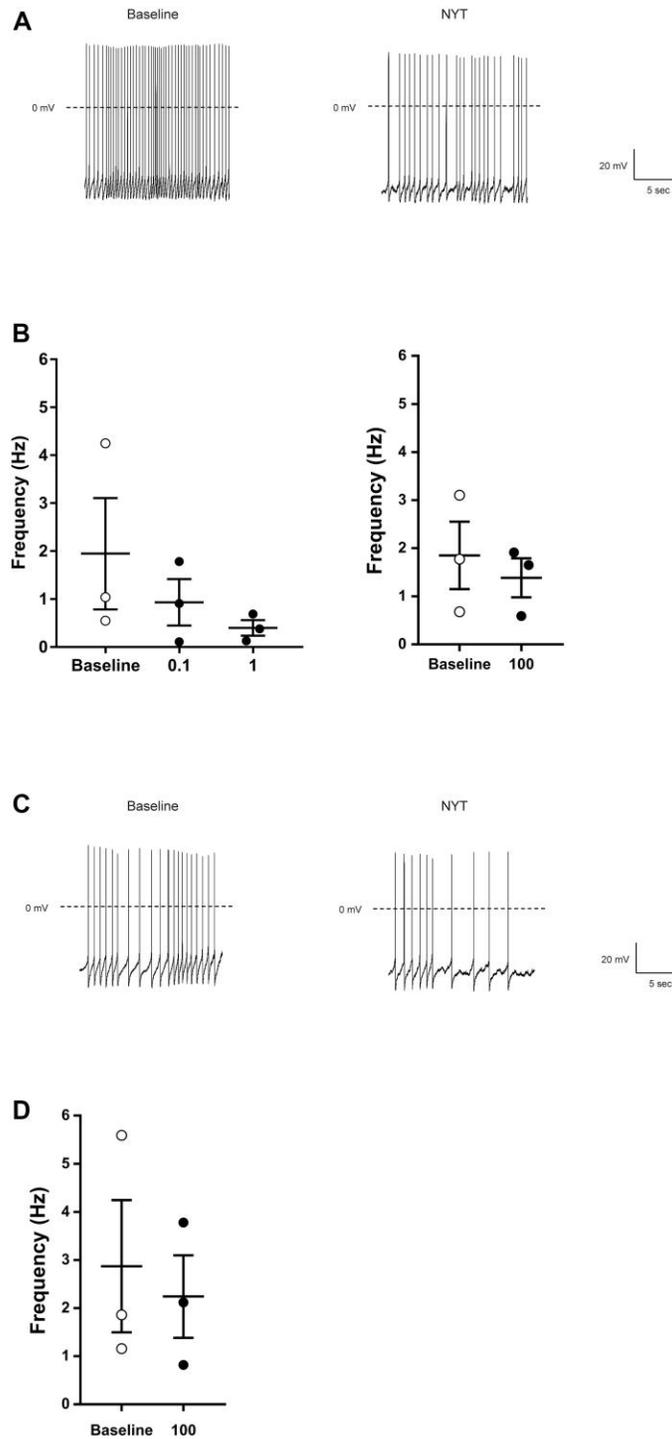
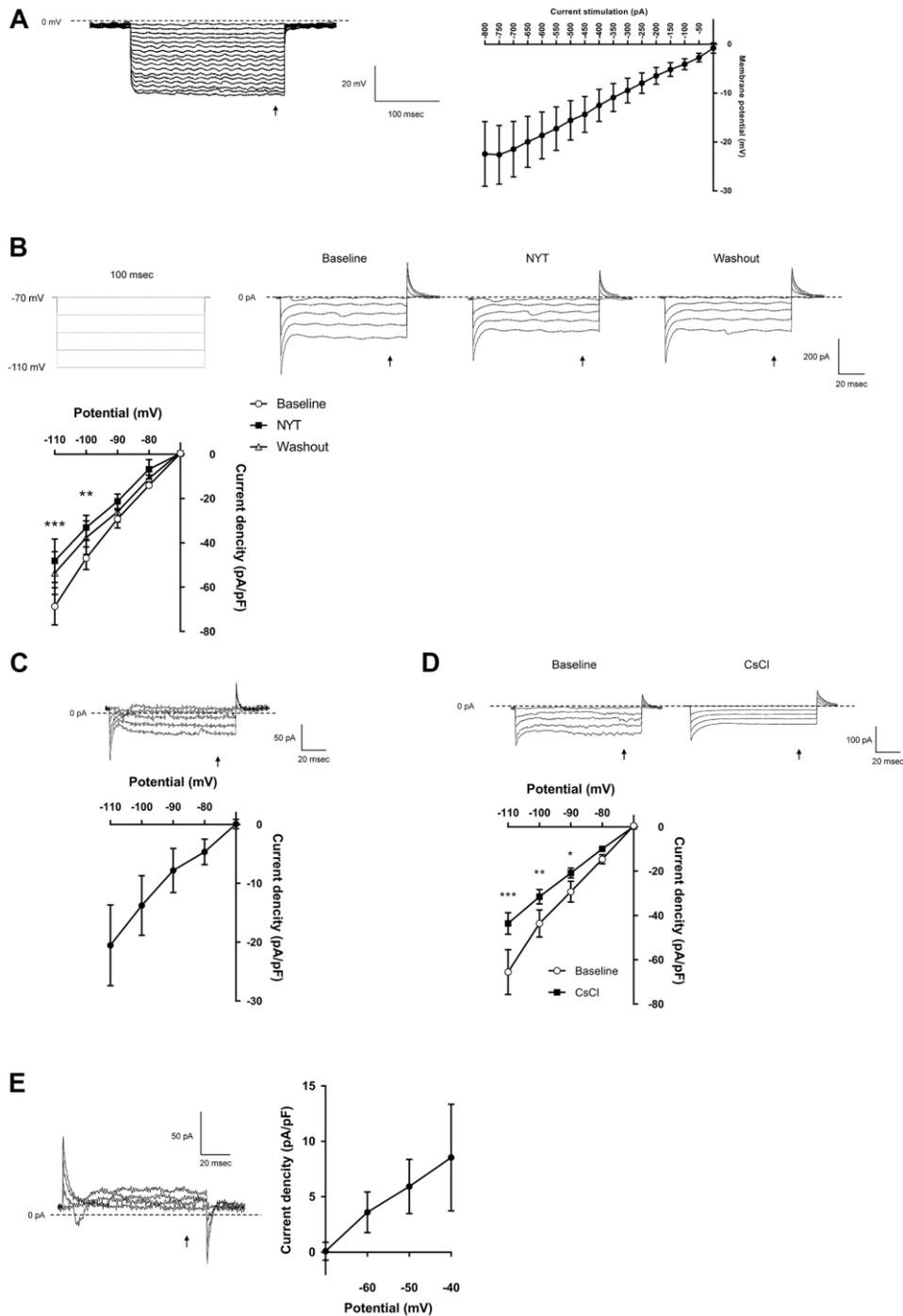


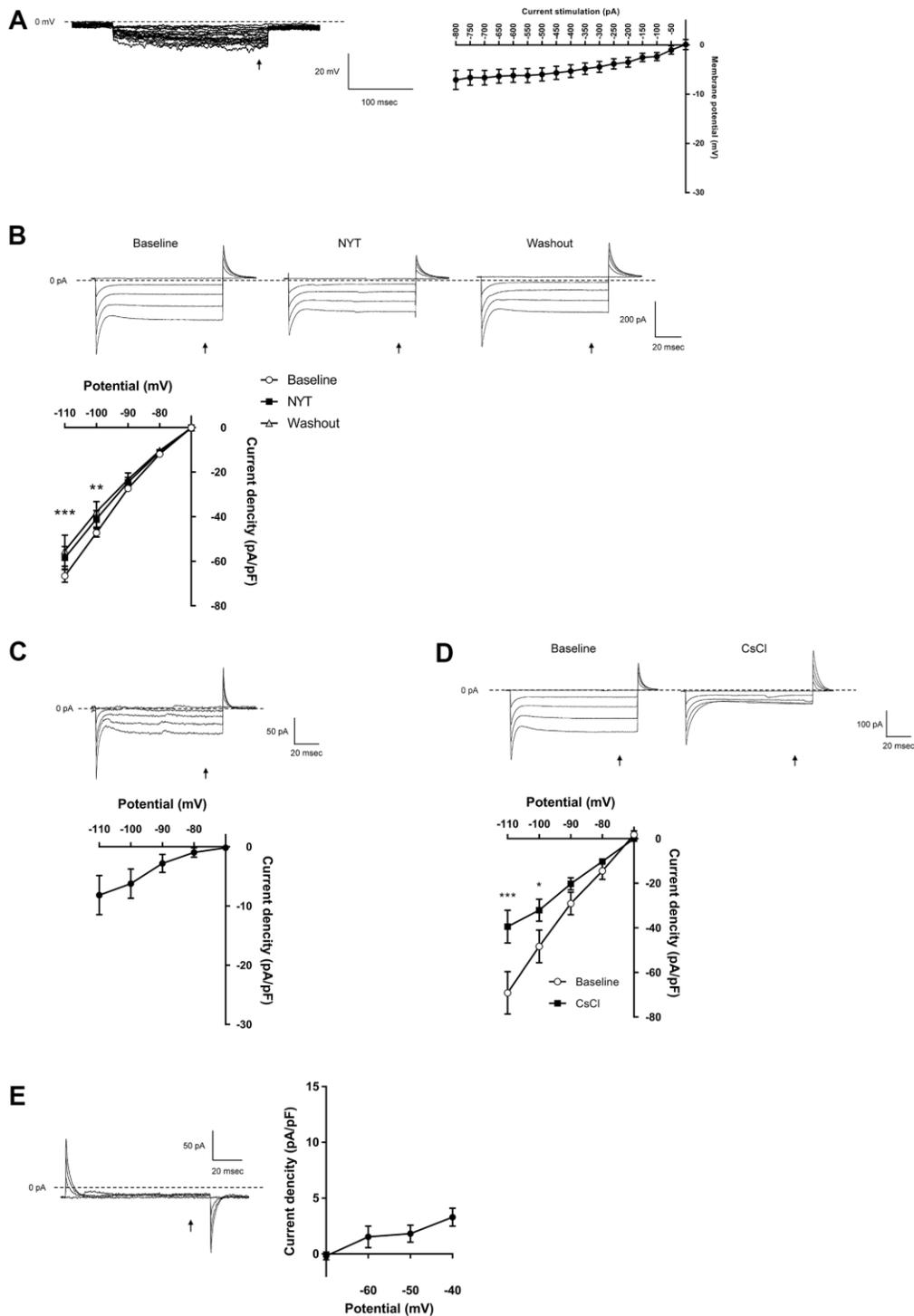
SUPPLEMENTARY FIGURES



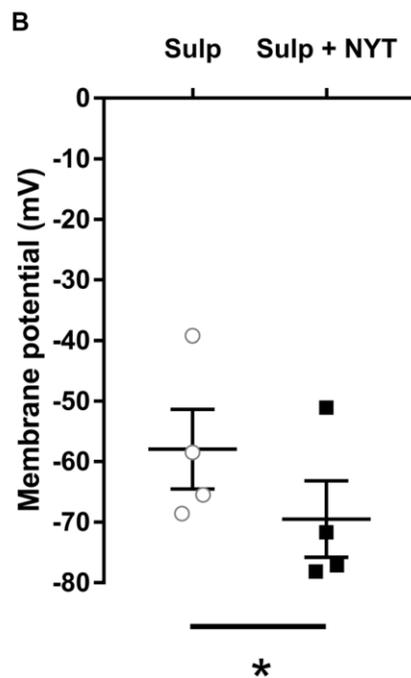
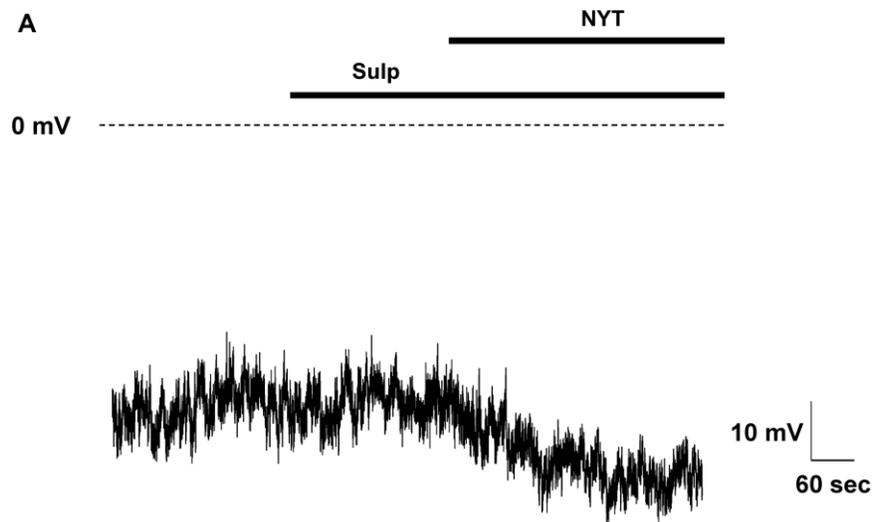
Supplementary Figure 1. The effects of NYT on the firing DAergic neurons in the VTA and SNpc. (A) The representative membrane potential recordings of firing DAergic neuron in the VTA before and after NYT (100 μg/mL) application. (B) The effect of NYT (0.1, 1, 100 μg/mL) on the firing frequency of VTA DAergic neurons (n = 3 from three mice). (C) The representative membrane potential recordings of the firing DAergic neuron in the SNpc before and after NYT (100 μg/mL) application. (D) The effect of NYT (100 μg/mL) on the firing frequency of SNpc DAergic neurons (n = 3 from three mice). Error bars are expressed as mean ± SEM. Statistical analyses were performed by (B (left)) one-way RM ANOVA followed by Dunnett’s multiple comparisons test and (B (right) and D) paired *t*-test.



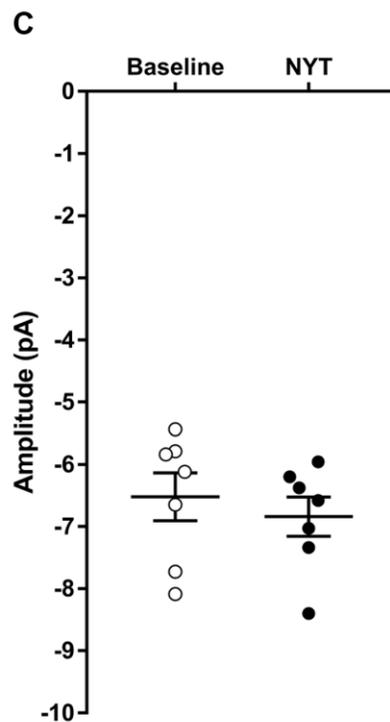
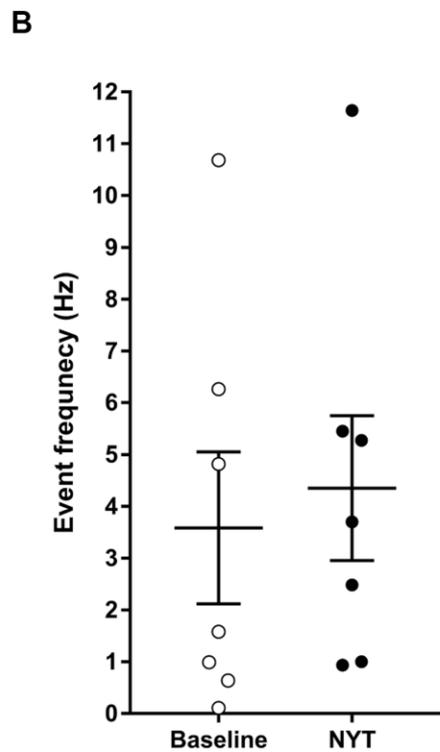
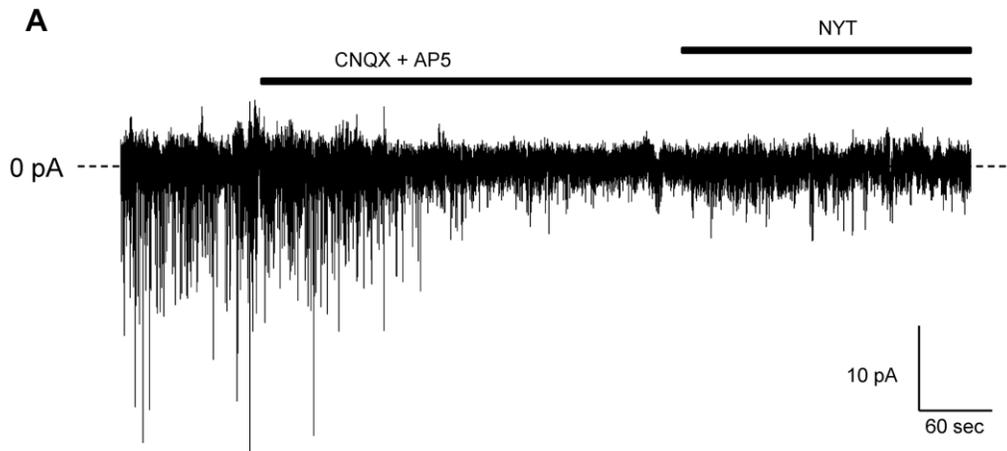
Supplementary Figure 2. The effects of NYT on the electrophysiological properties of MSNs in the NAc core subregion. (A) The representative NYT-sensitive membrane potential recording and the I-V relationship of NYT-sensitive membrane potential ($n = 11$ from seven mice). (B) The protocol of voltage pulses (10 mV increments from -110 to -70 mV, 100 msec, holding potential = -70 mV) for inward currents and the representative current recordings obtained by voltage pulses. The I-V relationship obtained by the voltage pulses ($n = 5$ from two mice). (C) The representative NYT-sensitive inward current recording and the I-V relationship of NYT-sensitive currents by voltage pulses ($n = 5$ from two mice). (D) The representative inward current recordings before and after CsCl application and the I-V relationship obtained by the voltage pulses ($n = 4$ from four mice). (E) The representative NYT-sensitive outward current recording and I-V relationship obtained by voltage pulses ($n = 5$ from two mice). The NYT-sensitive membrane potential and inward/outward currents were obtained by subtracting the trace after NYT application from the trace of baseline. Error bars are expressed as mean \pm SEM. Statistical analyses were performed by (B) two-way RM ANOVA followed by Dunnett's multiple comparisons test (** $p < 0.01$, *** $p < 0.001$; Baseline vs. NYT) or (D) two-way RM ANOVA followed by Bonferroni's multiple comparisons test (* $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$; Baseline vs. CsCl). The arrows indicate the time points of analysis.



Supplementary Figure 3. The effects of NYT on the electrophysiological properties of MSNs in the NAc shell subregion. (A) The representative NYT-sensitive membrane potential recording and the I-V relationship of NYT-sensitive membrane potential ($n = 11$ from nine mice). (B) The representative current recordings and the I-V relationship obtained by voltage pulses ($n = 6$ from four mice). (C) The representative NYT-sensitive inward current recording and the I-V relationship by voltage pulses ($n = 6$ from four mice). (D) The representative inward current recordings before and after CsCl application and the I-V relationship obtained by voltage pulses ($n = 4$ from 3 mice). (E) The representative NYT-sensitive outward current recording and I-V relationship obtained by voltage pulses ($n = 6$ from four mice). The NYT-sensitive membrane potential and inward/outward currents were obtained by subtracting the trace after NYT application from the trace of baseline. Error bars are expressed as mean \pm SEM. Statistical analyses were performed by (B) two-way RM ANOVA followed by Dunnett's multiple comparisons test (** $p < 0.01$, *** $p < 0.001$; Baseline vs. NYT) or (D) two-way RM ANOVA followed by Bonferroni's multiple comparisons test (* $p < 0.05$, *** $p < 0.001$; Baseline vs. CsCl). The arrows indicate the time points of analysis.



Supplementary Figure 4. The effects of NYT on dopamine D2 autoreceptor of VTA DAergic neurons. (A) The representative resting membrane potential recording of VTA DAergic neuron after sulpiride and NYT application. The bars indicate the duration of sulpiride or NYT application. (B) The resting membrane potential of VTA DAergic neurons after sulpiride and NYT application in TH-GFP mice. * $p < 0.05$; sulpiride vs. sulpiride + NYT ($n = 4$ from three mice, paired t -test). Error bars are expressed as mean \pm SEM.



Supplementary Figure 5. The effects of NYT on IPSCs of VTA DAergic neurons. (A) The representative IPSC recording of VTA DAergic neuron after NYT application. The bars indicate the duration of CNQX, AP5 or NYT application. (B) Frequency and (C) amplitude of IPSC inputs into the VTA DAergic neurons after NYT application. ($n = 7$ from three mice, paired t -test). Error bars are expressed as mean \pm SEM.