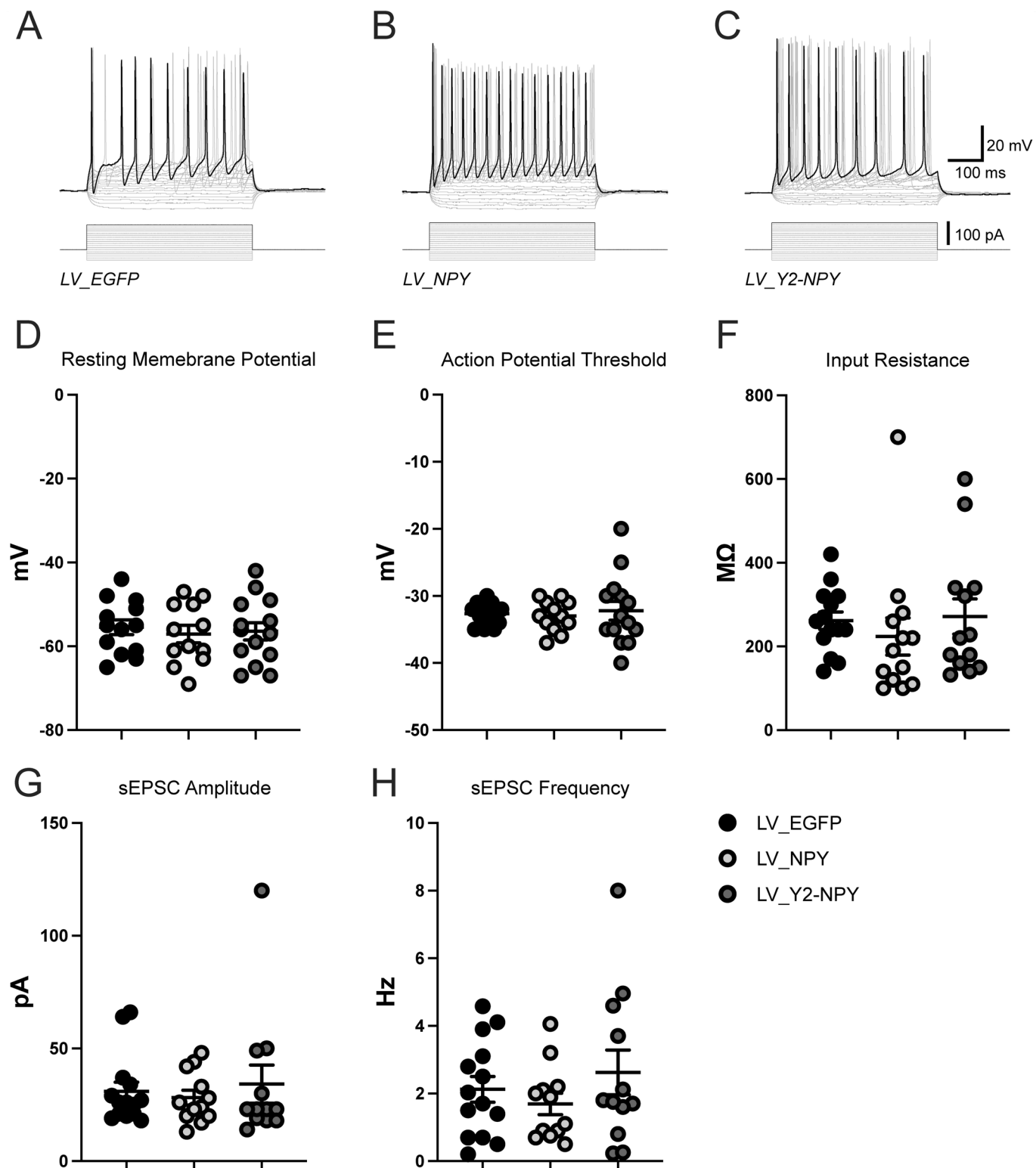


Expanded View Figures

Figure EV1. Electrophysiological characterization of LV transduced primary rat hippocampal neurons.

(A–C) Voltage response of LV transduced neurons to suprathreshold depolarizing current steps. (D–H) Dot plots summarizing average resting membrane potential, action potential threshold, input resistance, spontaneous excitatory post-synaptic currents (sEPSCs) amplitude, and sEPSCs frequency of transduced neurons. Data were shown as mean \pm SEM of 3 independent experiments (biological replicates). (D) LV_EGFP = -55.46 ± 1.77 mV; LV_NPY = -57.08 ± 2.07 mV; LV_Y2-NPY = -56.43 ± 2.07 mV. (E) LV_EGFP = -32.64 ± 0.45 mV; LV_NPY = -33.00 ± 0.66 mV; LV_Y2-NPY = -32.21 ± 1.4 mV. (F) LV_EGFP = 261.4 ± 20.94 M Ω ; LV_NPY = 223.8 ± 44.47 M Ω ; LV_Y2-NPY = 271.5 ± 42.14 M Ω . (G) LV_EGFP = 30.93 ± 4.12 pA; LV_NPY = 28.17 ± 3.25 pA; LV_Y2-NPY = 34.17 ± 8.49 pA. H, LV_EGFP = 2.12 ± 0.38 Hz; LV_NPY = 1.69 ± 0.32 Hz; LV_Y2-NPY = 2.63 ± 0.66 Hz. Source data are available online for this figure.



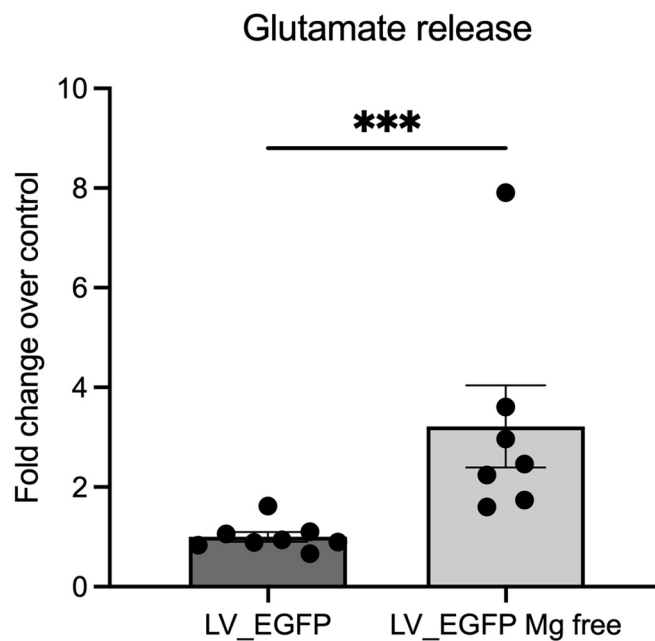


Figure EV2. Effect of Mg^{2+} removal on glutamate release.

Measurement of extracellular glutamate in transduced primary hippocampal neurons incubated for 20 min in Kreb's Ringer Hepes in the presence (LV_EGFP) or absence of Mg^{2+} (LV_EGFP Mg free). Values were normalized for total protein content and are presented as means \pm SEM of at least three independent experiments (biological replicates), with glutamate levels shown as fold change over control (LV_EGFP = 1.00 ± 0.10 ; LV_EGFP Mg free = 3.22 ± 0.82 ; $p = 0.0006$). Statistical significance was calculated using the Mann-Whitney test. *** $p < 0.0001$. Source data are available online for this figure.