

# Crosby Lab Code

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Scripts for analyzing evoked currents and action potentials in R

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# Evoked Currents

*Raw plots*

*Pruned individual plots*

*Pruned summary plots*

*Statistical analysis*

Table 1: A multivariate repeated measures ANOVA using Pillai's Trace shows that current amplitude varies significantly with time, and there are no significant interactions between factors like sex and treatment.

Parameters	df	Pillai's Trace	F	<i>p</i> -value
(Intercept)	1, 57	0.91	578.34	< 0.001
Treatment	3, 57	0.02	0.36	0.78
Sex	1, 57	0.01	0.48	0.49
Treatment:Sex	3, 57	0.03	0.54	0.66
Times	4, 54	0.38	8.26	< 0.001
Treatment:Times	12, 168	0.15	0.73	0.72
Sex:Times	4, 54	0.03	0.43	0.79
Treatment:Sex:Times	12, 168	0.26	1.31	0.22

*Levene's test*

*Univariate Shapiro*

*Box's  $M$*

*Multivariate normality*

*T-tests*

*Plot eEPSCs*

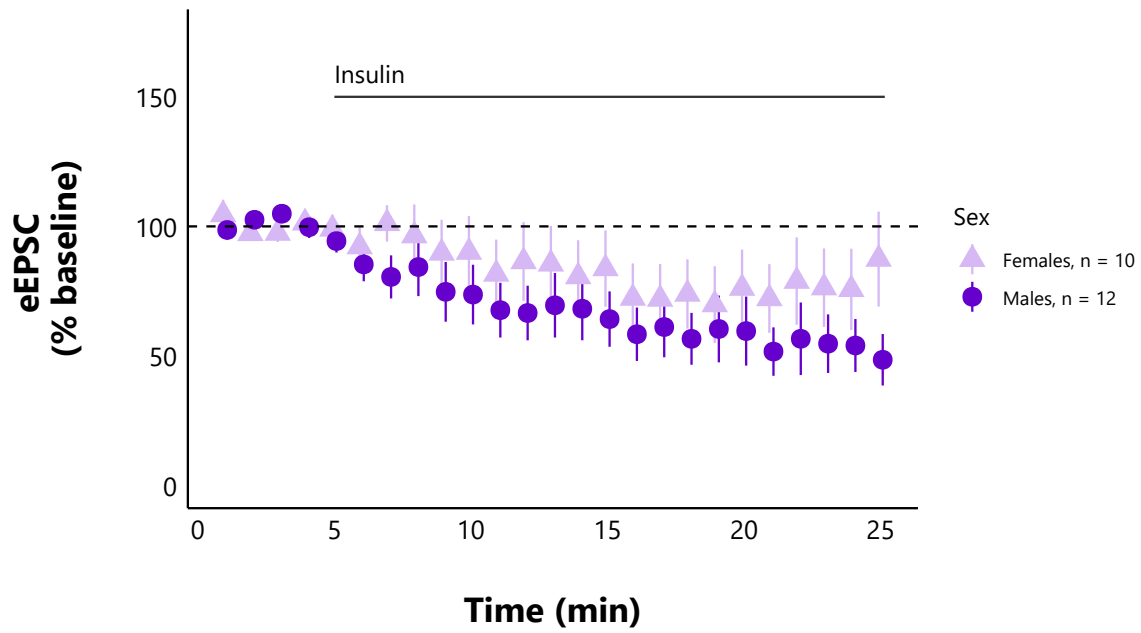


Figure 1: Each point represents the mean eEPSC amplitude ( $\pm$  the standard error) across all cells and  $n$  represents the number of unique cells. The asterisks indicate a statistically significant decrease in current amplitude relative to the baseline (t-test; \* =  $p < 0.05$ , \*\* =  $p < 0.01$ , \*\*\* =  $p < 0.001$ ). The representative traces consist of eEPSCs from one cell averaged over the baseline period (0 to 5 min) and last interval (20 to 25 min) of the recording, and the scale bar represents 50 pA/20 ms.

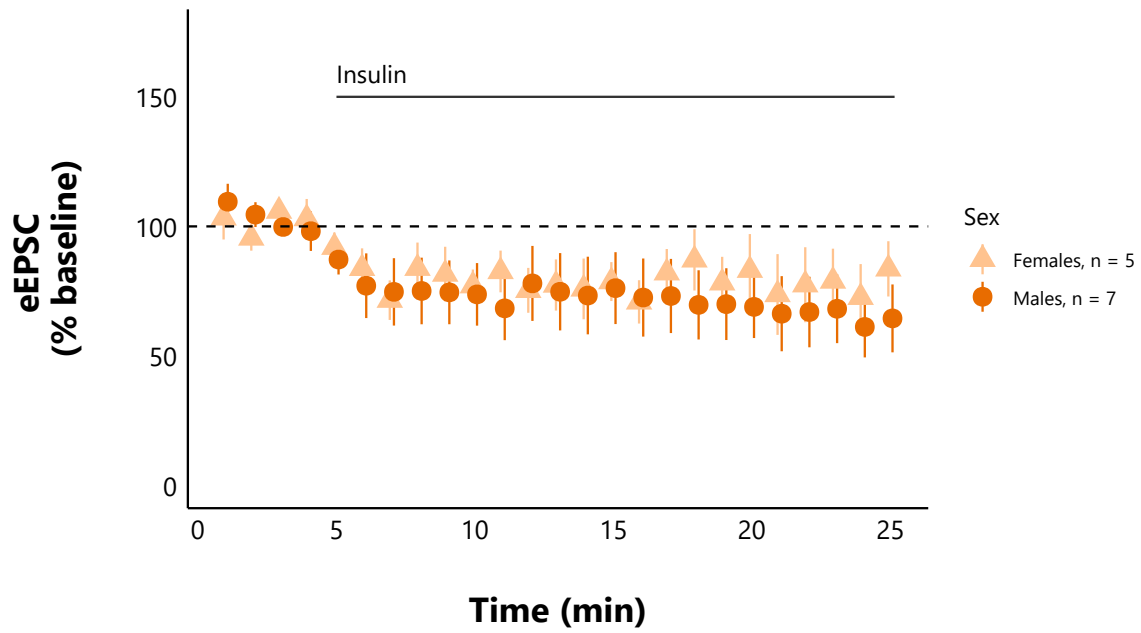


Figure 2: Each point represents the mean  $\pm$  SE eEPSC amplitude and  $n$  represents the number of unique cells. The asterisks indicate a statistically significant decrease in current amplitude relative to the baseline (t-test; \* =  $p < 0.05$ , \*\* =  $p < 0.01$ , \*\*\* =  $p < 0.001$ ). The representative traces consist of eEPSCs from one cell averaged over the baseline period (0 to 5 min) and last interval (20 to 25 min) of the recording, and the scale bar represents 50 pA/20 ms.

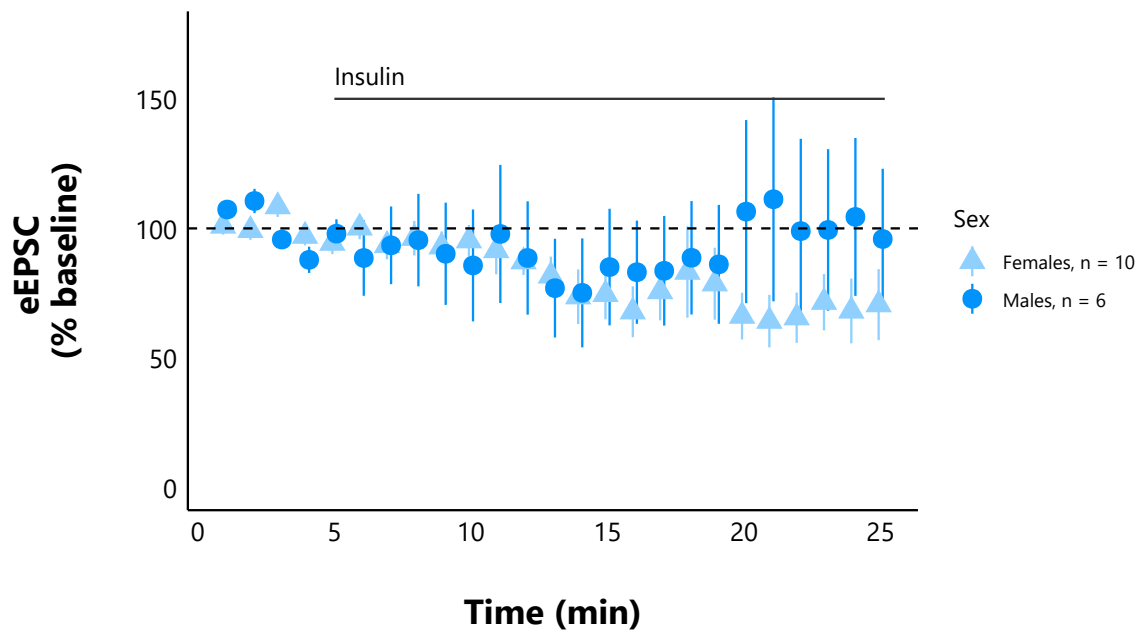


Figure 3: Each point represents the mean  $\pm$  SE eEPSC amplitude and  $n$  represents the number of unique cells. There are no asterisks because the decrease in eEPSC amplitude was not statistically significant relative to the baseline. The representative traces consist of eEPSCs from one cell averaged over the baseline period (0 to 5 min) and last interval (20 to 25 min) of the recording, and the scale bar represents 50 pA/20 ms.



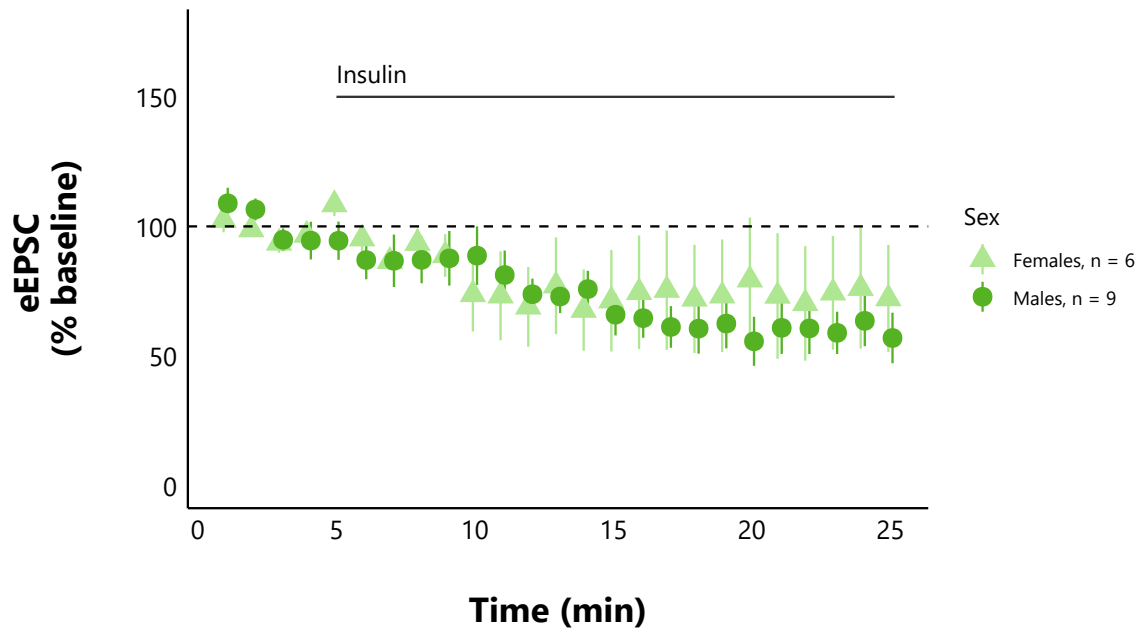


Figure 4: Each point represents the mean  $\pm$  SE eEPSC amplitude and  $n$  represents the number of unique cells. The asterisks indicate a statistically significant decrease in current amplitude relative to the baseline (t-test; \* =  $p < 0.05$ , \*\* =  $p < 0.01$ , \*\*\* =  $p < 0.001$ ). The representative traces consist of eEPSCs from one cell averaged over the baseline period (0 to 5 min) and last interval (20 to 25 min) of the recording, and the scale bar represents 50 pA/20 ms.

# Cell Coordinates Plot

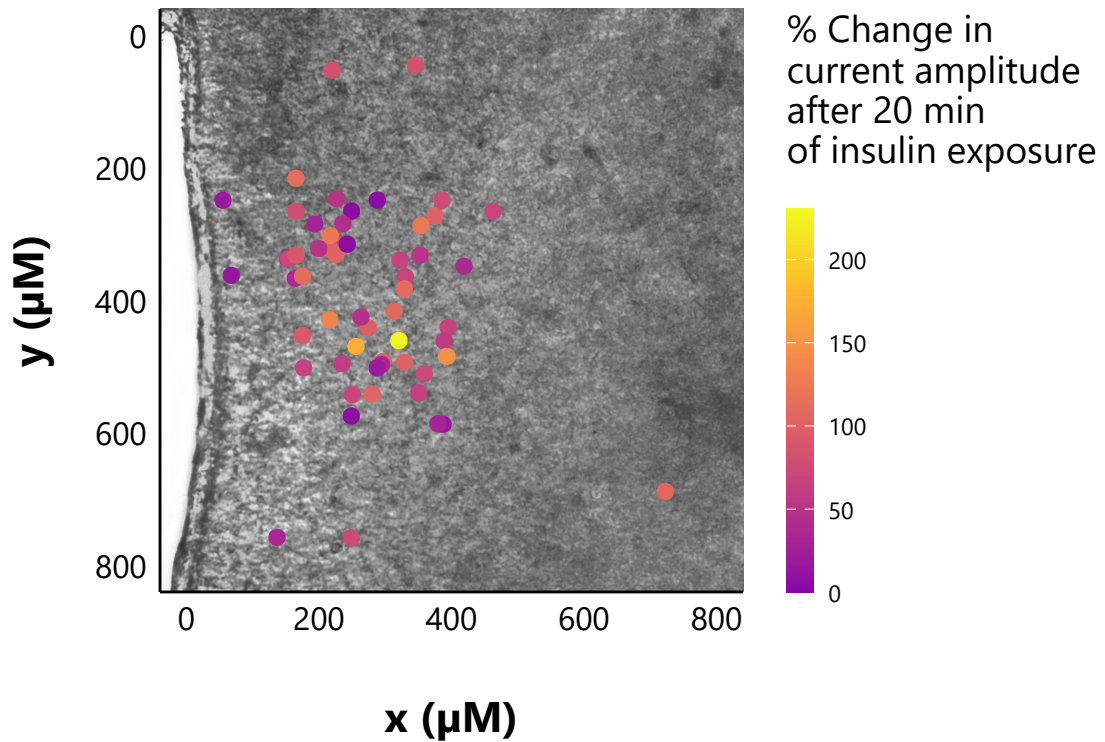


Figure 5: Each neuron is coloured according to the relative decrease in eEPSC amplitude after 20 minutes of insulin exposure. Neurons with similar responses to insulin did not cluster together, and there were no regions within the DMH that had a greater proportion of highly responsive neurons. In this figure (magnified 5X), the top of the third ventricle is located at (0,0).

# Final Treatment Comparisons Plot

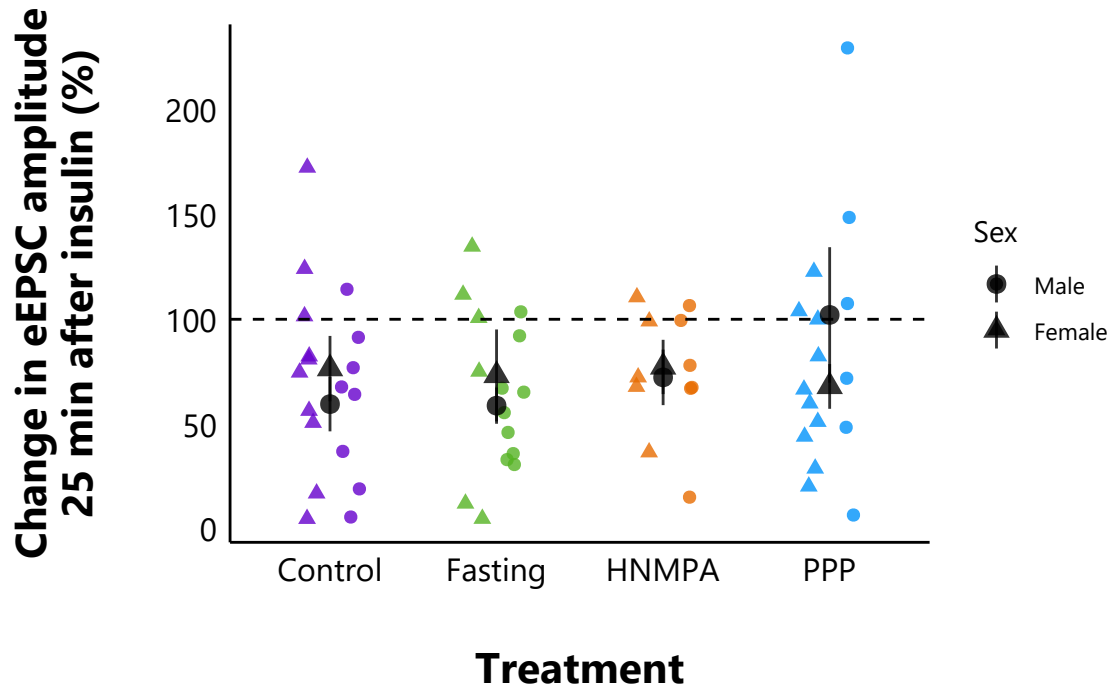


Figure 6: In this summary figure, each dot represents the mean eEPSC amplitude of a cell relative to the baseline (%) during the last 20-25 minutes of the evoked currents protocol. Black solid shapes indicate the mean  $\pm$  SE for each sex.

# Paired Pulse Ratio

*Shapiro*

*PPR Plots*

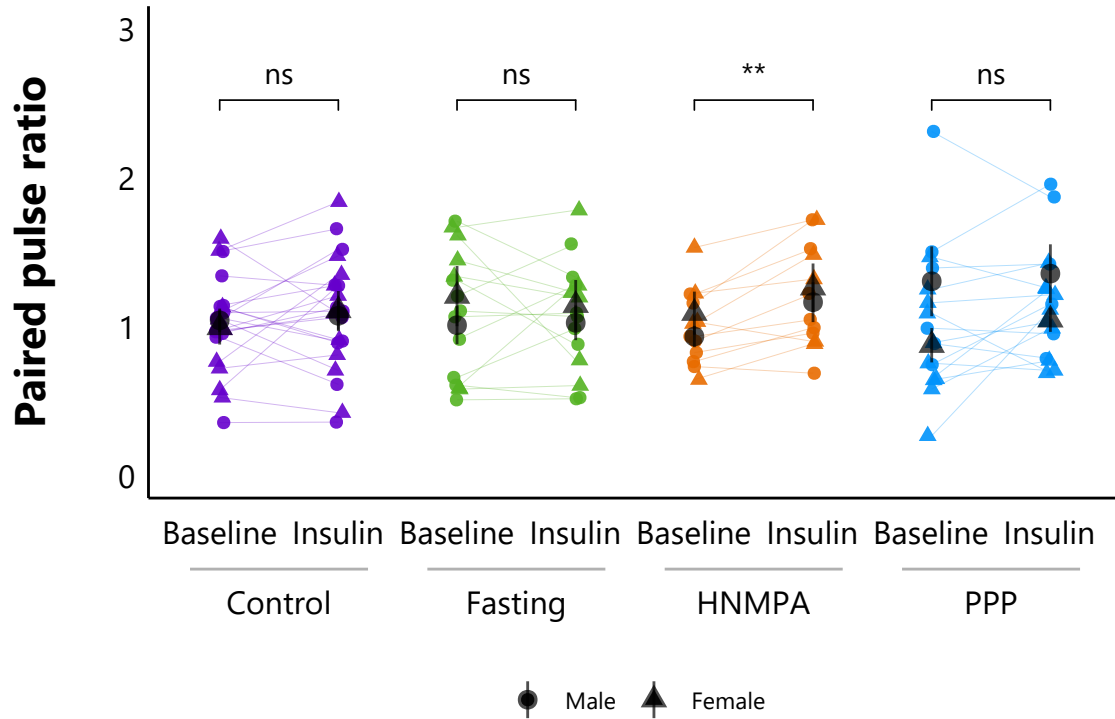


Figure 7: The paired pulse ratio (PPR) did not change significantly after insulin exposure for most of the treatment groups. The change in the PPR relative to the baseline was non-significant for the control, fasting, and PPP treatment groups, but it was statistically significant for the HNMPA-treated cells. Each dot represents the mean PPR of a single cell averaged over the baseline period (0 to 5 minutes) or the last 20 to 25 minutes of the evoked currents protocol. The asterisks indicate a statistically significant difference in the PPR between the two intervals (t-test; \* =  $p < 0.05$ , \*\* =  $p < 0.01$ , \*\*\* =  $p < 0.001$ ). The solid black circle and triangle overlay represents the mean PPR ( $\pm$  SE) per sex.

Table 2: A paired t-test comparing the mean paired pulse ratio (PPR) per cell during the baseline period (0-5 min) to the mean PPR after insulin exposure (20-25 min) shows that insulin does not significantly affect the PPR during experiments with no additional treatments, fasting, or PPP. Insulin significantly increases the PPR when the insulin receptor blocker HNMPA is applied.

Parameter	Statistic	DF	<i>p</i> -value
Control	-1.05	21	0.30
Fasting	0.15	14	0.88
HNMPA	-3.64	11	0.004
PPP	-1.43	15	0.17

# Action Potentials

*AP Frequency t-test*

*Shapiro test*

*Wilcoxon Test*

*Summary plot*

*Individual AP plots*

Table 3: A Wilcoxon signed-rank test shows that insulin significantly affects action potential amplitudes, thresholds, and half-widths. Insulin does not significantly affect the latency to fire, afterhyperpolarization amplitude, or the time of afterhyperpolarization.

Parameter	df	Statistic	<i>p</i> -value
Peak amplitude	15, 15	120	< 0.001
Threshold	15, 11	63	0.005
Latency to fire	15, 11	31	0.90
Half-width	15, 11	2	0.003
After-hyperpolarization amplitude	15, 11	35	0.90
After-hyperpolarization time	15, 11	17	0.17

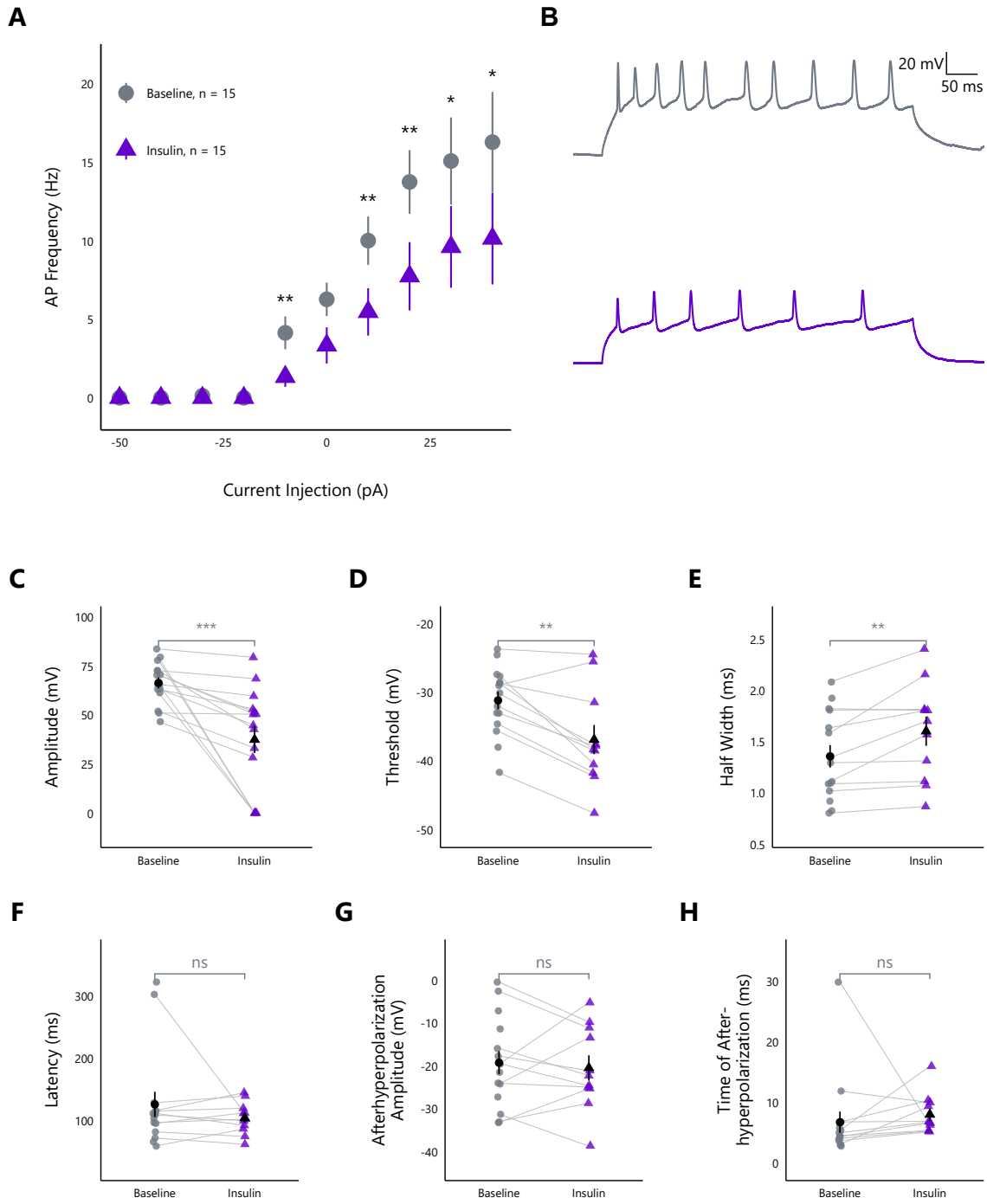


Figure 8: Insulin decreases the excitability of DMH neurons. Data were recorded before and then after 25 minutes of insulin exposure. *A*) Insulin significantly decreases action potential frequency (mean  $\pm$  SE;  $n$  is the number of cells). *B*) Representative traces from a current injection of 40 pA (top: baseline, bottom: insulin). *C*) Insulin significantly decreases action potential amplitudes *C*) and thresholds *D*), while significantly increasing half-widths *E*). Insulin does not significantly affect latency to fire *F*), after-hyperpolarization amplitude *G*) or after-hyperpolarization time *H*). Overlay on Figures C-H: mean  $\pm$  SE.  $n = 15$  for baseline and  $n = 11$  for insulin because 4 cells did not fire any action potentials after insulin exposure. Wilcoxon signed-rank test, \* =  $p < 0.05$ , \*\* =  $p < 0.01$ , \*\*\* =  $p < 0.001$ .