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Extended Data Fig. 1	Unit classification and physiological properties of DOWN-State Active (DSA) neurons	Supp_figure1.jpg	<p>(A) Units (n = 2,279 from 11 mice) were first classified based on trough-to-peak latency and firing rate. Each dot corresponds to one unit. Units with trough-to-peak latency < 0.55 ms were tentatively classified as narrow-waveform putative interneurons (INT; blue). Wide waveform units were grouped into putative excitatory cells (PYR cells; red) and inhibitory interneurons (INT; blue), on the basis of the bimodality of the marginal distribution of trough-to-peak latencies (top histogram), firing rate (right histogram) and burstiness (dot size) (Methods). Optogenetically identified neurons (Sst, PV, nNos. And ID2/Nkx2.1) are superimposed on the clouds. PV cells were recorded from V1. (B) DOWN-state rate ($P < 10^{-101}$, KW test), DOWN-state participation ($P < 10^{-69}$, KW test) and UP-state rate ($P < 10^{-136}$, KW test) distributions for putative pyramidal cells (PYR, red), interneurons (INT, blue) and DSA neurons (gray). (C) Similar layout as in C but for REM sleep (left) and walking behavior (right). (D) Peri-DOWN-state Z scored firing raster plot for all putative principal cells (PYR, left) and putative interneurons (INT, middle) for events detected during quiet waking state, as ranked according to Fig. 1D. Dashed lines delimit the ± 50ms window used to estimate firing rate. (E) DSA neurons in the rat PFC (left) and in mouse primary visual cortex, V1 (right). Left (top to bottom): Spike auto-correlogram and spike-waveform, peri-DOWN-state raster plot and average spike cross-correlation (CCG) between a reference neuron and all other simultaneously recorded units in a single session (each row corresponds to spikes of a target neuron referenced to the spike of DSA neuron at time 0. Black line is average). Right, autocorrelogram of 6 V1 DSA neurons. Middle panel: Cross-correlograms between DOWN-state (time 0 is the trough of DOWN state) and 6 DSA neurons. Bottom: average spike cross-correlation (CCG) between a reference DSA neuron (time zero) and all other simultaneously recorded units in that session.</p>

			***P<0.001.
Extended Data Fig. 2	Network property statistics of DSA interneurons	Supp_figure2.jpg	<p>(A) Statistical contrast matrices (two-sided Tukey's test) for firing rates of pyramidal cells (PYR), interneurons (INT) and DSA interneurons during waking quiescence (QWake), NREM sleep, REM sleep and active behavior (Walk). (B) Same layout and statistical comparisons as in A but for the average unit CCGs as a function of brain state. (C) Pearson correlations (tested using a Student's t distribution) between average CCG responses of NREM sleep, QWake, REM sleep and walking behavior for all groups. Note that spike vs population relationships are preserved across brain states. (D) Top: Average joint Z-score rate density between 10% of the interneurons (100 shufflings), 90% remaining interneurons and pyramidal cells neurons (left) and between 10% of the pyramidal cells (100 shufflings), interneurons and remaining pyramidal neurons. Bottom: Spearman correlation and statistical contrast matrices (two-sided Tukey's test) for interneuron and pyramidal neurons rate and Z-scored population for all shown joint histograms. (E) Average (mean \pm IC95) partial correlation values of the Z-scored rate for all groups (blue for pDSA,INT controlling for PYR; red for pDSA,PYR controlling for INT; magenta for pPYR,INT controlling for DSA), after truncating high firing rate units to match median the spikes number of pyramidal cells (average from n = 32 sessions, $P < 10^{-25}$, $F(2, 2787) = 57.42$, repeated measures ANOVA). (F) Left: distribution of excitatory divergence in all groups. Only putative pyramidal units excited their postsynaptic target cells (incidence probability for all groups in top-inset; $P < 10^{-110}$, $\chi^2(2) = 504.24$, χ^2 test). Right: distribution of excitatory convergence for all cell groups ($P < 10^{-67}$, $\chi^2(2) = 306.15$, χ^2 test). DSA neurons have fewer excitatory connections than the interneurons group ($P < 10^{-4}$, $\chi^2(1) = 11.36$, χ^2 test). ***P<0.001.</p>
Extended Data Fig. 3	Mechanisms of DSA neuron firing during DOWN states – model results	Supp_figure3.jpg	<p>(A) Spiking neural model containing 100 leaky DSA neurons receiving an asymmetric inhibitory/excitatory drive (top-left scheme). Bottom, UP/DOWN transitions with DOWN-selective firing DSA neurons (gray dots in the rastergram at the top). Top right, log firing rate distributions in the model corresponded to the those of the recorded neurons. (B) Peri-DOWN-state Z scored firing raster plot for all simulated principal cells (PYR, left) and</p>

			<p>interneurons (INT, right), including (incl) 10% of DSA_n, ranked according to their event response. Bottom, Z-scored average CCG for each model neuron group (compare with Fig. 1E). The peri-DOWN state raster plots and unit cross-correlograms (Extended Data Fig. 3B) were similar to experimental results (compare with Fig. 1D, E) for both pyramidal and interneuron populations (Supp. Fig. 3C-D; PYR rate mean \pm SD: 0.2 ± 0.09 Hz during DOWN, 1.31 ± 0.41 Hz during UP; $P \sim 0$; INT: 0 ± 0.1 Hz during DOWN, 8.38 ± 2.52 Hz during UP, $P \sim 0$) as well as for the DSA group (7.43 ± 0.90 Hz during DOWN, 1.29 ± 0.34 Hz during UP, $P < 10^{-83}$; Kruskal-Wallis test). (C) Magnitude of DOWN-state response as a function of the CCG response. Marginal CCG (top, $n = 20,000$, 5000 and 500 PYR, INT and DSA_n; $P \sim 0$, KW test) and DOWN-state response (right, $P \sim 0$, KW test) distributions for simulated pyramidal cells, interneurons and DSA neurons in each group. (D) DOWN-state/UP-state firing rate ratio ($P \sim 0$, KW test), DOWN-state rate ($P \sim 0$, KW test), DOWN-state participation ($P \sim 0$, KW test) and UP-state rate ($P \sim 0$, KW test) distributions for all simulated pyramidal cells (PYR, red), interneurons (INT, blue) and DSA neurons (gray). (E) Average (mean \pm IC95) partial correlation values of the Z-scored rate for DSA and INT (blue), DSA and PYR (red), and PYR and INT (magenta) for different values of inhibitory weight simulated in the INT-DSA connection (0 to -3 mV, $n = 2$ simulations of 5100 cells per condition; $P \sim 0$, $F(2, 44.094) = 278.36$, repeated measures ANOVA). Model neurons showed that partial correlation between pyramidal cells and interneurons was positive at all time bins but negative for DSA neurons versus pyramidal cells and interneurons (compare with Fig. 2I), and it was strongly dependent on the inhibitory drive imposed on the DSA neurons by other inhibitory neurons. (F) DSA neurons Z-scored DOWN-state responses (mean \pm IC95) for different values of inhibition simulated on the DSA neurons ($n = 200$ DSA_n per condition). DSA neurons rate during UP-states, but not during DOWN-states, depends on the inhibitory strength between INT and DSA neuron pairs. (G) The peak delay of DSA neurons during DOWN-states correlates with the inhibitory strength of the INT-DSA connection (mean \pm IC95, $n = 2$ simulations of 5100 cells per conditions; Pearson Correlations correlations tested using a Student's t distribution). (H) Partial correlation spectra (mean \pm IC95) of the Z-scored simulated rate for each group as in Fig. 2D ($n = 5$ simulations of 5100 cells using a</p>
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			<p>DSA Inh Drive = 1.5 mV; $P \sim 0$, $F(2, 31.495) = 921.33$, repeated measures ANOVA). (I) Temporal dynamics of the Z-scored DOWN-state responses across all model neurons from H (top-left, mean \pm IC95; $P < 10^{-104}$, KW test). Simulated DSA neurons reach peak firing rate ~ 40 ms after maximum rate decrease of pyramidal cells and interneurons during DOWN-states (top-right; medians, interquartile ranges and error bars). Bottom, Likewise, the average CCG delay between simulated DSA neuron spikes and spikes of other cells (bottom, dashed gray line; $P < 10^{-315}$, KW test) matches DOWNs states responses (~ 40ms, left avCCG average curves, right, medians, interquartile ranges, maxima and minima). (J) Similar displays as in I but generated from recorded neurons (mean \pm IC95; $n = 1457$, 775 and 47 PYR, INT and DSA_n, respectively, from 11 mice; $P < 10^{-22}$ for the temporal dynamic of the Z-scored DOWN-state responses and $P < 10^{-7}$ for the average CCG delay, KW test). The trough of the cross-correlogram of DSA neurons versus other neurons occurred earlier than the peak of the cross-correlograms of other neurons as in the model. *$P < 0.05$; **$P < 0.01$; ***$P < 0.001$. For model details, see Methods.</p>
Extended Data Fig. 4	Mechanisms of DSA neurons firing during the DOWN-states	Supp_figure4.jpg	<p>(A) Top: Z-scored peristimulus histogram (PSTH) for cells recorded in the Sst::ChR animals ($n = 2$ mice), grouped as pyramidal cells (PYR, red), interneurons (INT, blue) and DSA neurons (gray). Horizontal bar: duration of optogenetic stimulation. Middle: firing rate trajectory for the DSA – INT response ($P < 10^{-29}$ between baseline and rebound epochs, KW test). Bottom: Firing trajectory for the DSA – PYR response ($P < 10^{-11}$, KW test). (B) Same than A but for PV::ChR animals ($n = 4$ mice; $P = 0.009$ and $P = 0.39$ for ‘INT resp vs DSA resp’ and ‘PYR resp vs DSA resp’, respectively, KW test). Non-significant, n.s. (C) Spike auto-correlogram (top, mean \pm IC95) and spike waveform (bottom, mean \pm IC95) of ID2/Nkx2.1-expressing DSA neurons (gray, $n = 6$) in optogenetic experiments are similar to non-responding DSA neurons (dark yellow, $n = 3$, ID2-/Nkx2.1- or non-recombined neurons) as well as to DSA neurons observed in other mice (black, $n = 41$; KW test). (D) Depth distribution of all recorded units (as in Fig. 1I), highlighting the position of the nNos⁺ neurons (yellow diamonds; $n = 25$ cells optogenetically responding cells from 4 Nos1::ChR mice). Note the deep position of the two DSA/nNos⁺ neurons (encircled). (E) Same layout as in D but highlighting the position of the ID2+/Nkx2.1+ cells</p>

			in black circles (n = 8 from 6 ID2/Nkx2.1::Ai80 mice). Note that the two ID2+/Nkx2.1+ cells, residing above layer 5 (encircled), were not DSA neurons. **P<0.01; ***P<0.001, after Tukey's honesty <i>post hoc</i> multiple comparisons.
Extended Data Fig. 5	Intrinsic electrophysiological features of L5/6 ID2/Nkx2.1 neurons	Supp_figure5.jpg	(A) Group differences (mean \pm SD) for three electrophysiological parameters between L1 neurogliaform cells (green, n = 25 L1 NGFC), L5/6 ID2/Nkx2.1 (gray, n = 18 cells) and delayed fastk spiking cells (in blue, from ref ³⁰ , n = 23 dFS). Note that Goldberg et al used 600 ms second square pulses on the delayed respond tests. (B) Example of rebound spiking test protocol. Average traces from a L5/6 ID2/Nkx2.1 cell held at -70mV, then hyperpolarized to -90mV for 3s, then depolarized to either -70 mV (black) or -50 mV (red). If present, rebound spikes should be apparent immediately following the depolarizing step (n = 11 L5/6 ID2/Nkx2.1 cells).
Extended Data Fig. 6	Anatomical features of deep and superficial cortical ID2/Nkx2.1 interneurons	Supp_figure6.jpg	(A)Depth distribution of ID2+/Nkx2.1+ neurons (tdTomato+ or GFP+) in the medial prefrontal cortex (mPFC), PPC and anterior and posterior regions of the primary visual cortex (V1). (B) Photographs showing distribution of GFP+ (or tdTomato+) neurons in the mPFC of an ID2/Nkx2.1::Ai80 mouse (Ai65) (top and middle images, reverse contrast epifluorescence; n = 47 neurons in 9 samples from 3 mice). Most neurons were found in layer 6. (C) A layer 6 tdTomato+ ID2/Nkx2.1 neuron (red) at two different depth levels. Note the putative PV+ (cyan) puncta in close apposition to the tdTomato+ cell (arrows). Top, 0.38 μ m-thick single optical section. Bottom, 0.76 μ m-thick maximum intensity z-projection (representative micrograph, tested in 3 mice). (D) Top: Z projections (confocal maximum-intensity projection) of two intracellularly-filled tdTomato+ ID2/Nkx2.1 neuron from two ID2/NKx2.1::Ai65 mouse. Bottom: reconstruction of an intracellularly-filled tdTomato+ ID2/Nkx2.1 neuron (from a 223.2 μ m thick confocal stack, maximum-intensity projection as inset) of an ID2/NKx2.1::Ai65 mouse. The morphological features of the two reconstructed ID2/Nkx2.1 Neuron corresponded to those described in the literature for neurogliaform cells in L2/3 ^{24,29} and L1 ²⁶ . Total dendritic length: 7968.5 μ m and 11641 μ m for the cell in Fig 5D and the cell in Extended Data Fig. 6, respectively; total axonal length: 77873 μ m and 53669 μ m; membrane surface in dendrites and soma: $1.02 \times 10^5 \mu\text{m}^2$ and $2.81 \times 10^5 \mu\text{m}^2$; membrane surface in axons: $9.21 \times 10^5 \mu\text{m}^2$ and $1.14 \times 10^6 \mu\text{m}^2$; number of dendritic branches: 117 and 105; axonal branches: 747 and

			<p>314. (E) ID2+/Nkx2.1+ neuron (red) in L6b weakly immunoreactive for both NPY and nNos (arrow; n = 64 tdTomato+ ID2/Nkx2.1 neuron from 3 mice were tested against NPY). Note also strongly immunoreactive NPY+ nNOS+ neuron lacking tdTomato expression (arrowhead on top). (F) A tdTomato+ ID2/Nkx2.1 neuron in L5 of PPC immunoreactive for calretinin (white arrowheads) and lacking expression of Calbindin (widefield epifluorescence; n = 54 tdTomato+ ID2/Nkx2.1 neuron from 3 mice were tested, 2 of which were positive for calretinin). (G) PV+ tdTomato+ neuron in L2 of V1 (representative micrograph, tested in 3 mice). (H) Left: a layer 2 tdTomato+ ID2/Nkx2.1 neuron in layer 2 (widefield epifluorescence, reverse contrast). Right, high magnification of the marked rectangle area shows the tdTomato+ ID2/Nkx2.1 neuron axon distribution ('cartridges', some of them marked by arrows) typical of axo-axonic neurons (widefield epifluorescence, reverse contrast; representative micrograph, tested in 3 mice). The ID2/Nkx2.1 cells labeled here are identical to the Lamp5/Lhx6 type described in ref^{21,22}. ID2 and Lamp5 are both broadly expressed in putative neurogliaform interneurons²¹, and Lhx6 is expressed downstream of Nkx2.1 in cortical interneurons. The proportion of the Lamp5/Lhx6 subtype has been reported to be $1.4 \pm 0.2\%$ of total interneurons²², with the majority located in L5/6.</p>
Extended Data Fig. 7	Effector role of the DSA neurons	Supp_figure7.jpg	<p>(A) Cross-correlation (CCG) between Non-Mod (reference, n = 648 units) and -Mod (yellow, n = 118 units) compared to the CCG within only Non-Mod cells (green) (top and bottom-left; mean \pm IC95; P = 0.004, KW test), and CCG asymmetry [before-after]/[before+after] (bottom-right; medians, interquartile ranges, maxima and minima; P < 10^{-5}, KW test). (B) Amplitude response differences (medians, interquartile ranges, maxima and minima) for - Mod (n = 51 units from 9 session with at least 1 DSA neuron from 6 ID2/Nkx2.1::CatCh mice) and No Mod cells (n = 195 units) in LOW and HIGH DSA neuron rate DOWN-UP events. (C) Temporal dynamics of the Z-scored DOWN-UP events responses for LOW and HIGH events (n = 2.954 and 5.317 events, respectively; medians, interquartile ranges, maxima and minima). (D) Averages (mean \pm IC95) of the Fourier-transform spectra for non-stimulated (control, black) and stimulated (green) epochs (top) during NREM (n = 12 sessions from 6 mice). Peak frequency, peak amplitude and delta power (0.5-4 Hz) comparisons for non-stimulated and stimulated epochs (bottom). Note decreased</p>

			peak frequency of in the delta band during stimulated epochs, likely reflecting the prolongation of DOWN state upon activation of ID2/Nkx2.1 neurons (see Fig. 6). (E) Bottom, Z-scored sequential activity of a simultaneously recorded population during DOWN-UP events sorted for 'High DSA rate' events (session mean template), vertically arranged by latency, including two DSA neurons bottom two rows). Top, session mean template for 'Low DSA rate' events but arranged according to 'High DSA rate' classification (see Fig. 6F for 'Low DSA rate'). Note that the two types of event sequences are different when DSA spiking activity in the slow oscillation even is high or low. *p < 0.05; **p < 0.01; ***p < 0.001.
Extended Data Fig. 8	Stimulation of DSA neurons during NREM sleep	Supp_figure8.jpg	(A) Multitaper spectrogram from 0-40 Hz of one entire recording day with NREM optogenetic stimulation (including baseline, behavioral session 1, rest period and behavioral session 2) utilized for state scoring. Middle, electromyogram (EMG) over the same session (au, arbitrary units). Bottom, state scoring of the same session (as in ref ¹⁷). States are coded as label to the left. (B) Three-dimensional plot showing state segregations. Each point corresponds to 1 s of recording time, with colors indicating the identified state during that second as labeled. (C-D) Same than A-B but for a no-simulation (control) day. (E) NREM sleep durations (mean ± SD) between test sessions 1 and 2 are similar in the four different conditions. (F) Normalized fractions of time spent in Wake, NREM and REM in the home cage between behavioral sessions 1 and 2. (G) CCG trace (mean ± IC95) between light pulses and UP and DOWN states (left) for all NREM stimulated sessions (n = 15 sessions). Right, Fraction of stimulated events over all detected events (medians, interquartile ranges, maxima and minima; P < 10 ⁻⁵ , KW test). ***p < 0.001.

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9 **Sleep DOWN state-active ID2/Nkx2.1 interneurons in the neocortex**

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ABSTRACT

Pyramidal cells and GABAergic interneurons fire together in balanced cortical networks. In contrast to this general rule, we describe a distinct neuron type in mice and rats whose spiking activity is anti-correlated with all principal cells and interneurons in all brain states but most prevalently during the DOWN state of non-REM sleep. We identify these DOWN-state active (DSA) neurons as deep-layer neocortical neurogliaform cells that express ID2 and Nkx2.1, and weakly immunoreactive to nNOS. DSA neurons are weakly excited by deep layer pyramidal cells and strongly inhibited by several other GABAergic cell types. Spiking of DSA neurons modified the sequential firing order of other neurons at DOWN-UP transitions. Optogenetic activation of ID2/Nkx2.1 interneurons in the posterior parietal cortex during nonREM sleep, but not during waking, interfered with consolidation of cue discrimination memory. Despite their sparsity, DSA neurons perform critical physiological functions.

INTRODUCTION

The brain often ‘reboots’ itself during sleep from transient silence. The silent epochs, known as delta waves or DOWN states are interleaved among spiking activity-rich UP states during non-REM (NREM) sleep, and the toggling between these states is known as cortical slow oscillation¹⁻⁶. NREM, and particularly slow oscillations, have been implicated in the development of cortical circuits, homeostatic regulation of network dynamics and memory consolidation⁵⁻¹⁰.

Neuronal nitric oxide (nNOS)-expressing neurons show elevated activity during sleep, as measured by the increased expression of the early gene cFos¹¹⁻¹³, and may play a special role in controlling slow oscillations. Deep layer neurons of the cortex^{3,4,14}, a variety of somato-dendritic regenerative currents in cortical pyramidal cells and the excitatory recurrences among neurons have been postulated to be the source of stochastic metastability and persistent UP state activity¹⁵. In turn, spike afterhyperpolarization and synaptic depression, combined with decreasing excitation-inhibition balance have been suggested to be responsible for switching from UP to DOWN state³. The mechanism of the DOWN state is less understood. In addition to the absence of spikes, synaptic activity is also reduced during the DOWN cortical state, leaving both principal cells and inhibitory interneurons transiently in a dis-excited and dis-inhibited state. During the DOWN-UP transition, pyramidal neurons become active rapidly and sequentially¹⁰. While some pyramidal neurons occasionally emit action potentials during the DOWN state⁸, it is a generally held view that no cortical neuron fires selectively during the DOWN state.

Against this currently held scenario, we describe a small set of deep layer neurons in three neocortical areas of mice and rats whose spiking activity is inversely correlated with all principal cells and all other types of interneurons in both waking and sleeping states, with highest firing rate displayed during the DOWN state (“DOWN State Active neurons”, DSA). We identify DSA neurons as ID2/Nkx2.1 neurogliaform interneurons.

RESULTS

In the first set of experiments, we used multiple-shank silicon probes to record local field potential (LFP) and spiking activity in the posterior parietal cortex (PPC) of 11 mice (Fig. 1A-C). The depth distribution of LFPs, current sinks and sources and unit firing were used to

identify cortical layers¹⁶. The most prominent landmark was the largest power in the high frequency band (high-pass filtered >120 Hz), defining mid-layer 5. A layer source in layer 5 identified DOWN state of slow oscillation (Fig. 1C; ¹⁶). Single units were separated into putative excitatory and inhibitory classes (n = 1457 and n = 822, respectively), using a multi-step approach¹⁶ (Extended Data Fig. 1A; Methods).

Physiological characterization of DOWN state active (DSA) neurons

DOWN states were identified using LFP and spiking criteria (Fig. 1B; Methods). As expected, firing rates of most putative pyramidal neurons across all layers were suppressed during DOWN states (Fig. 1D; Extended Data Fig. 1B; $P < 10^{-237}$ between UP and DOWN states; Wilcoxon's paired signed-rank test). Similarly, cells classified as putative interneurons were also mostly silenced ($P < 10^{-128}$). However, we also identified a small subset of neurons (n = 47 in PPC) that were specifically and dominantly active during the DOWN state of NREM (DSA neurons; Fig. 1D; 5.60 ± 1.54 Hz during DOWN, 2.17 ± 1.42 Hz during UP; $P < 10^{-9}$; Wilcoxon's paired signed-rank test). Average cross-correlation (CCG) between DSA neurons and all principal cells and all interneurons showed an inverse relationship during both NREM and waking states (Fig. 1E; for REM sleep and walk see Extended Data Fig. 1C). The firing rate ratio between DOWN and UP states effectively segregated DSA neurons from other neurons (Fig. 1F; Extended Data Fig. 1B). Similar segregation was obtained by plotting the relationship between spikes of DSA neurons and DOWN state epochs against cross-correlation features with other neurons (Fig. 1G and Extended Data Fig. 1D). DSA neurons had lower burst indices (defined as fraction of spikes with ≤ 6 ms window to all spikes), wider autocorrelograms and waveforms than other interneurons and pyramidal cells and had longer duration waveform than other interneurons (Fig. 1H). All DSA neurons were found in deep cortical layers (Fig. 1I).

In a second set of experiments, DSA neurons with similar features were also identified in the visual cortex (n = 6 DSA neurons of 1,326 neurons in 19 mice; Extended Data Fig. 1E; ¹⁶) and prefrontal cortex in the rat (n = 2 DSA neurons of 1,121 neurons; Extended Data Fig. 1E; ¹⁷).

Local circuit control of DSA neurons by pyramidal cells and interneurons

The population firing rates of pyramidal cells and interneurons varied moderately across behavioral states (PYR: 8.46% average change across all states, $P < 10^{-41}$; INT: 12.08%, $P < 10^{-75}$; Friedman test). In contrast, DSA neurons were most active during NREM sleep and quiet waking, coinciding with transiently reduced firing of the remaining neuronal population, and their rate decreased more than 4-fold during walking and REM sleep (Fig. 2A; Extended Data Fig. 2A; 22% change across all states; $P < 10^{-7}$; Friedman test). DSA neurons were negatively correlated with other neurons during their high firing rate states (NREM and quiet wakefulness), whereas population cross-correlation between pyramidal neurons and interneurons were positive and larger during NREM sleep (Fig. 2B; Extended Data Fig. 2B-C; ¹⁸).

To gain insight on the nature of interactions of DSA neurons with the rest of the PPC population, we constructed joint co-activity histograms of pyramidal cells and putative interneurons with DSA neurons (Fig. 2C; Extended Data Fig. 2D). The firing rates of DSA neurons were more negatively correlated with the activity of interneurons (Fig. 2C, *y-axis* and blue trace in top panel) than with those of pyramidal cells (*x-axis*, red trace), suggesting that DSA neuron are inhibited by GABAergic cells. Moreover, partial correlation of the firing rates between DSA neurons and interneurons was more negative than the partial correlation rate between DSA neurons and pyramidal cells for all time bins tested (Fig. 2D), suggesting that DSA neurons are disinhibited during DOWN states. Firing rate differences between pyramidal cells and interneurons could not explain these effects (Extended Data Fig. 2E). In contrast, firing rates of pyramidal neurons and interneurons were positively correlated from 10 ms to 1 s time bins (Fig. 2D). To further support these findings, we quantified putative monosynaptic connections and spike transmission probabilities between pairs of neurons ^{16,19}. As expected, we found stronger spike transmission between pyramidal cells and interneurons compared to pyramidal-pyramidal pairs (Fig. 2E-G; Extended Data Fig. 2F; ¹⁶). Spike transmission between pyramidal-DSA neuron pairs was weakest (Fig. 2F-G) and less probable than between pyramidal cell–interneuron pairs (Fig. 2F; Extended Data Fig. 2F; $P < 10^{-190}$; Fisher's exact test), supporting the population co-activity results. Finally, DSA neurons were strongly coactive with other DSA neurons and this synchrony was stronger than observed within interneuron and pyramidal neuron populations, respectively (Fig. 2H).

To account for the unusual firing patterns of DSA neurons, we adopted a model of slow oscillations¹⁵ to provide a proof of principle regarding how the observed spike correlations among pyramidal cells, interneurons and DSA neurons might explain their physiological connectivity (Extended Data Fig. 3). In line with our experimental observations, the model predicted that DSA neurons are strongly inhibited by other interneurons and weakly excited by nearby deep layer pyramidal cells.

DSA neurons comprise deep layer ID2/Nkx2.1 interneurons

In the search for the cellular identity of DSA neurons as well as the interneuron types that may inhibit them, we optogenetically tested four promoter-specific transgenic mouse lines expressing light-sensitive opsins (ChR or CatCh) in GABAergic subpopulations (Fig. 3A-F), including nNOS-expressing neurons¹¹. None of the DSA neurons recorded in these mice were somatostatin (Sst)-expressing (n = 0/7 units from two Sst:ChR mice) or parvalbumin (PV)-expressing (n = 0/5 V1 units from 4 PV::ChR mice¹⁶) interneurons, which together account for ~90% of the infragranular layer inhibitory neurons in mice²⁰. Two of twenty five light-responding neuronal nNOS-expressing cells (from 4 Nos1::ChR mice) were DSA neurons (Fig. 3D), and were located below layer 5 (Extended Data Fig. 4D). Based on this clue, we searched further by screening lines that marked sparse subpopulations of nNos-expressing interneurons preferentially in deep cortical layers. A small subpopulation of GABAergic neurons defined by Lamp5 and Lhx6 co-expression are found in both Nos1-CreERT2 and Nkx2-1-CreERT2 driver lines with the majority located in deep cortical layers^{21,22}. By cross-breeding three mouse lines (ID2-CreER, Nkx2.1-Flpo and the Cre-Flpo dependent reporter line Ai80), we created an intersectional line expressing the channelrhodopsin CatCh in neurons expressing ID2 and Nkx2.1, corresponding to Lamp5 and Lhx6, respectively²³. We found that the majority of the recorded DSA neurons (66%, 6 of 9 DSA neurons from 5 ID2/Nkx2.1 mice) were of the light-responding ID2/Nkx2.1 type (Fig. 3E-F; 2.98 ± 1.41 Hz baseline; 13.13 ± 1.73 Hz stimulation; $P < 10^{-3}$ for all units; Wilcoxon's paired signed-rank test). Conversely, 6 of the 8 light-responding ID2/Nkx2.1 neurons (75%) were DSA cells (Extended Data Fig. 4E).

Optogenetic stimulation of Sst- and PV-expressing neurons suppressed firing rates of most DSA neurons (Fig. 3F; Sst stimulation: 6/7 DSA units decreased; PV stimulation: 4/5 DSA units decreased significantly), in addition to inhibiting other interneurons and pyramidal cells (Fig. 3B, C; Sst: 67% of the pyramidal cells and 46% of the interneurons decreased; PV: 81% of the pyramidal cells and 48 % of the interneurons decreased). Upon release from optogenetically induced suppression of DSA neurons by PV and Sst interneurons, DSA neurons responded with excess spiking lasting 70 to 200 ms (Extended Data Fig. 4A, B). In *Nos1::ChR* mice, in addition to the two light-excited DSA neurons, optogenetic stimulation of nNOS-expressing neurons significantly suppressed firing of 10 of 25 DSA units (Fig 3D,F). The same intensity light stimulation in *ID2/Nkx2.1::Ai80* mice was less effective in suppressing other interneurons and pyramidal cells (Fig. 3E; 9.7% of the pyramidal cells and 24% of the interneurons), possibly due to sparseness of the light-stimulated *ID2/Nkx2.1* neurons. The autocorrelograms and spike waveforms of opto-tagged *ID2/Nkx2.1* neurons were indistinguishable from those of other DSA neurons (Fig. 3G and Extended Data Fig. 4C), indicating that all or the majority of DSA neurons likely belong to the *ID2/Nkx2.1* type.

In vitro electrophysiological characterization of *ID2/Nkx2.1* neurons ($n = 22$ tdTomato+ cells in 4 *ID2/Nkx2.1::Ai65* mice) identified them as ‘late spiking’ neurons, with > 0.6 second of delay to the first spike at near-threshold current steps, large spike after-hyperpolarization (15 – 20 mV), and regular, non-adapting spikes in response to supra-threshold current steps (Fig. 4A, B). Since these electrophysiological features of *ID2/Nkx2.1* neurons are similar to neurogliaform cells^{24–28}, we compared six measured parameters with those of layer 1 neurogliaform cells²⁶. Although several parameters of the two groups differed quantitatively from each other (Fig. 4B), delayed ‘late spiking’ nature of both groups classifies deep layer *ID2/Nkx2.1* neurons as neurogliaform cells^{26,29}. Only one other neuron type, the delayed fast-spiking cell, shows late spiking but its other features differ substantially from both deep layer *ID2/Nkx2.1* and layer 1 neurogliaform cells (Extended Data Fig. 5A; ³⁰). *ID2/Nkx2.1* neurons ($n = 11$) did not show hyperpolarization-induced rebound spikes (Extended Data Fig. 5B).

Anatomical features of DSA ID2/Nkx2.1 interneurons

Our physiological and optogenetic findings indicated that DSA neurons resided mainly below layer 5 and they were weakly excited by local pyramidal cells and strongly inhibited by Sst- and PV-expressing interneurons, compared to other neurons. Commensurate with these results, we found that the layer distribution of DSA neurons in the PPC (see Fig1I) matched that of tdTomato-expressing ID2/Nkx2.1 interneurons (n = 54 neurons, 3 mice), with most neurons residing in layer 6 (~ 66%, Fig. 5A; $P < 10^{-4}$, across layer comparison; Kruskal-Wallis (KW) test), similar to the distribution of Lamp5-Lhx6 neurons in other areas of the mouse cortex (Extended Data Fig. 6A-B)^{21,22}. The majority (~70%) of tdTomato-expressing neurons were weakly immunoreactive for nNOS (0.33 ± 0.20 relative nNos intensity for n = 12 tdTomato+/nNos+ cells; 0.85 ± 0.09 for n = 9 Sst+/nNos+ cells; $P < 10^{-8}$, KW test), mainly in deep layers (Fig. 5B, C; 83% in layer 6; $P < 10^{-31}$, Fisher's exact test; n = 3 ID2/Nkx2.1 mice), and lacked immunoreactivity for Sst and PV (n = 0/54 for both PV and Sst; Fig. 5B and Extended Data Fig. 6C) and most of them lacked NPY (n = 11/64, Extended Data Fig. 6E) and calretinin (CR, n = 2/54, Extended Data Fig. 6F). nNOS+ deep layer tdTomato-expressing interneurons (118 cells from 3 ID2/Nkx2.1::Ai65 mice) had small somata, compact dendritic fields and dense local axon arborizations (inter-varicosities median distance average = $1.5 \pm 0.14 \mu\text{m}$, n = 2 neurons; Fig. 5C-D and Extended Data Fig. 6D). These feature are similar to those described for neurogliaform cells in layer 2/3^{24,31}, layer 1²⁶ and layer 6²³. We also observed PV-immunoreactive puncta in close apposition to the somata of ID2/Nkx2.1 cells, supporting the physiology-based connectivity measures (Extended Data Fig. 6C). In superficial layers, ID2/Nkx2.1 neurons immunonegative for nNOS (58% from all tdTomato+ cells in layers 1 and 2/3) were axo-axonic cells and other kinds of neurons, including PV+ cells with basket-like axon endings (Extended Data Fig. 6G-HG; ²¹), indicating that they are distinct from deep layer ID2/Nkx2.1 cells.

Targets of DSA - ID2/Nkx2.1 interneurons

An important question that we addressed next is the impact of DSA neurons on their targets. We identified potential target pyramidal cells and interneurons by their significantly decreased firing rates in response to optogenetic stimulation of ID2/Nkx2.1 interneurons (Fig. 4E; negatively modulated or -Mod; Methods). Those whose rates did not change were designated as non-Modulated (Non-Mod). Both -Mod pyramidal cells and -Mod interneurons were located, on average, deeper in the cortex (below layer 5) than the Non-Mod pyramidal cells and interneurons

(Fig. 6A). Both –Mod pyramidal cells and –Mod interneurons fired later during the DOWN–UP transition than Non-Mod cells (Fig. 6B and Extended Data Fig. 7A), with minor effects on the response magnitude (Fig. 6B).

The above analysis suggested that activity level of DSA neurons in the DOWN state can delay the timing of their targets during the emerging UP state, and thus, their firing order at the DOWN-UP transition^{32,33}. To test this hypothesis, we classified each slow oscillation event (DOWN state/ UP state pairs) based on the level of DSA neuronal activity. In each sleep session, we calculated the mean firing rate of DSA neurons and classified each DOWN state as ‘LOW’ (below session mean) or ‘HIGH’ (above mean) contribution of DSA spiking (Fig. 6C). With such separation, we re-investigated the relationship between – Mod and Non-Mod neurons (pyramidal cells and interneurons combined). We found that when DSA neurons were active, the recruitment of the –Mod neurons was significantly delayed during DOWN–UP transitions, whereas no such difference was present for the No–Mod neurons (Fig. 6D and Extended Data Fig. 7B), supporting the hypothesis that activity of the DSA neurons affect spike timing of their putative targets. In further support of this view, HIGH DOWN-UP events transitioned slightly, but significantly, slower than LOW events (20-80% rise time (mean \pm SD: 0.10 ± 0.08 and 0.11 ± 0.08 , for LOW and HIGH events, respectively; $P = 0.0036$; Wilcoxon’s paired signed-rank test; Extended Data Fig. 7C). Conversely, delta power peak frequency was significantly lower during optogenetically simulated NREM epochs than during non-stimulated ones (1.70 ± 0.32 and 1.96 ± 0.18 , respectively; $P = 0.0144$; Wilcoxon’s paired signed-rank test; Extended Data Fig. 7D).

In addition, we also computed the session mean sequence rank order of all neurons for the DOWN-UP events separately for the HIGH and LOW groups. In turn, these LOW DSA and HIGH DSA session sequence templates were correlated with every individual slow oscillation event (Fig 6E; the tested event was not used for the template generation). We found that when the HIGH session mean DSA template was correlated with HIGH individual DSA events or when the LOW session mean DSA template was correlated with LOW DSA events, the correlation coefficients were significantly higher than comparisons across group events (HIGH template with LOW DSA events and LOW template with HIGH DSA events; Fig. 6F, G and

Extended Data Fig 7E). As further controls, within-group (HIGH-HIGH and LOW-LOW) rank order correlations were also higher than the correlation of rank orders between events of the first and second halves of sleep sessions (irrespective of DSA activity level) or the correlations between randomly shuffled events (average of 100 shufflings). In summary, DSA neurons can modify the firing order sequences of their target neurons in DOWN-UP transitions.

Behavioral effects of DSA – ID2/Nkx2.1 interneurons

Our second approach to examine the impact of DSA neurons on their targets was based on a behavioral readout. Recent imaging work has demonstrated the importance of the PPC in visual pattern discrimination and memory-guided behavior³⁴, and we adopted that paradigm to test the potential role of DSA neurons in the consolidation of cue discrimination memory. Five mice, implanted bilaterally with optic fibers in deep layers of the PPC were trained to perform a visual cue-guided choice task (Fig. 7A). The mice first learned the correspondence between the correct visual pattern and correct choice in the T maze. Two identical sessions were performed each day, session 1 in the morning and session 2 in the afternoon so that the animals could utilize the information learned in the morning in their afternoon session. After the animals had learned the task, the visual discrimination patterns were varied from day to day, thus requiring new learning each day. Between sessions, animals were allowed to sleep in their home cage either unperturbed or with bilateral activation of PPC ID2/Nkx2.1 neurons during NREM sleep or waking. Non-stimulation and optogenetic stimulation days varied in a semi-random manner within and across mice. The same intensity and duration pulses (100 ms) that were found effective in driving ID2/Nkx2.1 neurons in the physiological experiments were randomly (every 500 ms \pm 400 ms) delivered during the entire NREM sleep in the home cage after the morning session (7496 \pm 1600 pulses).

As expected, on days with no perturbation in the PPC, mice showed a significant improvement of discrimination performance (probability of correct trials, computed as correct trials/all trials) in the afternoon session (Fig. 7B, C, gray color; from 0.57 \pm 0.09 in morning session to 0.68 \pm 0.06 in afternoon session, $P = 6.10 \times 10^{-5}$, Wilcoxon's paired signed-rank test). In contrast, the memory boost from the morning session was abolished by stimulation of ID2/Nkx2.1 neurons during NREM (Fig. 7B). However, when optogenetic stimulation was applied during waking

(mostly quiet wake state) in the home cage ($500 \text{ ms} \pm 400 \text{ ms}$ random pulses; 7487 ± 1333 pulses; $P = 0.35$ between NREM and awake pulses number, KW test), memory gain from sleep was not affected (Fig. 7B; $P = 0.94$; Tukey-Kramer's range test). As a further control, when ID2/Nkx2.1 neurons were activated while the mouse ran through the center arm during the morning task, memory gain from the morning to afternoon session was not affected (7B, C; $P = 0.62$; Tukey-Kramer's range test).

Differences between morning session performance across days (Fig. 7B; $P = 0.57$, KW test), or time spent in NREM state (Extended Data Fig. 8A-E; $P = 0.37$, KW test) or changes in other brain states (REM and waking behavior, see Extended Data Fig. 8F) could not explain these results. Light stimulation at the same intensity in a second group of mice ($n = 3$), which lacked ChR expression in ID2/Nkx2.1 ('photostimulation control'), did not prevent memory improvement from morning to afternoon sessions, excluding potential non-specific effects (Fig. 7D). Our stimulation protocol impacted approximately 20% and 5% of all detected UP and DOWN-states, respectively (Extended Data Fig. 8G), which is largely proportional to the duration distributions of UP and DOWN states.

Finally, we explored the effect of optogenetic stimulation of ID2/Nkx2.1 cells on sleep replay of neuronal correlations using the explained variance (EV) method (Fig. 7E-F; see Extended Methods)³⁵. We found significant reactivation for non-stimulated days (Fig. 7G), an effect which was abolished by optogenetic stimulation of ID2-Nkx2.1 cells during NREM. Moreover, EV was significantly higher on non-stimulated days than on optogenetic stimulation days. In summary, artificial activation of ID2/Nkx2.1 neurons during NREM sleep eliminated the benefit of sleep-assisted memory consolidation in PPC, likely interfering with the physiological structure of UP and DOWN states.

DISCUSSION

We described a unique set of deep layer neurons in PPC, V1 and frontal neocortical areas whose spiking activity is inversely correlated with all principal cells and all other types of interneurons across all brain states. This anticorrelation was most prevalent during the DOWN state of NREM sleep where these neurons fired specifically and selectively ("DOWN State Active neurons",

DSA). We demonstrated that DSA neurons belong to a special subset of neurogliaform, weakly immunoreactive for nNOS with dense local axon arbors, which we identified as ID2/Nkx2.1 cells. They were inhibited by parvalbumin and somatostatin- and nNO-expressing interneurons and weakly excited by deep layer pyramidal cells, relative to other interneurons. Spiking of DSA neurons during the DOWN state delayed the sequential order of their target neurons at the DOWN-UP transition and prolonged the DOWN state. Finally, we found that optogenetic activation of ID2/Nkx2.1 interneurons in the posterior parietal cortex during NREM sleep interfered with consolidation of cue discrimination memory and learning-induced sleep replay of neuron pairs.

Unique features of deep layer ID2/Nkx2.1 neurogliaform interneurons

We identified DSA neurons as deep-layer neocortical ID2/Nkx2.1 interneurons, also known as Lamp5+/Lhx6+ neurons^{21,23}. Although only a subset of the physiologically characterized DSA neurons were optogenetically identified as ID2/Nkx2.1 interneurons, all DSA neurons shared similar physiological features, including spike waveform, spike autocorrelation dynamic, brain state-dependent firing rates, anti-correlation with spiking activity of all other neuron types and their anatomical layer distribution. A small subset of DSA cells in Nos1::ChR mice were nNOS-expressing neurons. Anatomically, ID2/Nkx2.1 interneurons had weak nNOS expression but did not express PV or Sst. Neurons expressing nNOS have previously been classified in two major types: ‘NOS-type I’, with strong NOS expression, large somata and Sst co-expression; and ‘NOS-type II’, a more diverse group, with smaller sized somata and low level of NOS expression. This latter group includes neurogliaform, VIP, PV and Sst interneurons^{36,37}. Thus, ID2/Nkx2.1 interneurons are within the NOS-type II group.

In earlier experiments, nNOS-expressing neurons in multiple cortical regions of mice, rats and golden hamsters showed increased c-Fos expression following sleep deprivation, and the magnitude of nNOS expression correlated with delta power of NREM sleep¹¹. Non-selective nNOS knockout mice show reduced NREM sleep time, shorter NREM bouts, decreased slow oscillation power and impaired NREM compensation after sleep deprivation¹². In contrast, when nNOS deletion is confined to Sst neurons in the neocortex and striatum, these mice show blunted slow oscillation power in response to sleep deprivation but without altered NREM

duration or sleep structure¹³, suggesting that while power in the slow oscillation band is primarily regulated by cortical circuits²⁻⁴, control of sleep state choreography is mainly subcortical¹³. Mice lacking nNOS expression in Sst positive neurons exhibit impairment in recognition memory but not in a hippocampal-dependent object recognition task¹³. In all these studies, the critical c-Fos expressing neurons were of the nNOS-type 1.

In our experiments, most nNOS neurons fired together with other interneurons and pyramidal cells during the UP state of slow oscillation. Thus, while type 1 nNOS neurons, present in all cortical layers, and deep layer ID2/Nkx2.1 interneurons are quite distinct, they appear to complement their contributions to NREM. Type 1 nNOS neurons may release nitric oxide¹³ and bring about deepening of NREM sleep by increasing the power of slow oscillations. ID2/Nkx2.1 interneurons, on the other hand, become selectively active during the DOWN state, prolong the duration of DOWN state and bias the sequential organization of pyramidal neurons during the DOWN-UP transition, and may contribute to other, yet to be discovered, functions.

ID2/Nkx2.1 interneurons showed typical features of neurogliaform cells, including small somata and dense local arborization with small boutons²⁴⁻²⁶. The physiological features of neurogliaform neurons in the superficial cortical layers have been characterized previously. Uniquely among all known GABAergic interneurons, single presynaptic action potentials of neurogliaform neurons can activate postsynaptic metabotropic GABA_B receptors^{24,38} and halt activity of principal cells through dendritic calcium channels³⁹. In addition, they affect their targets via slow inhibitory postsynaptic currents (IPSCs) mediated by GABA_A receptors localized on the dendrites of pyramidal neurons²⁹. In response to near threshold current steps, they fire following a noticeable delay, while stronger depolarizing pulses induce sustained non-accommodating spike trains with large spike after-hyperpolarizations²⁶⁻²⁸. ID2/Nkx2.1 interneurons shared these biophysical features with neurogliaform neurons in superficial cortical layers, although the two groups differed in quantitative terms.

In contrast to other forebrain GABAergic interneuron types (e.g., PV, Sst, VIP subtypes), neurogliaform neurons originate from several distinct embryonic brain regions. Most hippocampal neurogliaform cells arise from Nkx2.1+ lineages (Lhx6+) whereas the vast majority

of neocortical neurogliaform cells have a non-Nkx2.1 CGE/POA origin^{25,40,41}, the exception being the ID2/Nkx2.1 type described here. Neurogliaform cells are present in all cortical layers but are most abundant in layer 1 and layer 2/3²⁰. We observed ID2/Nkx2.1 interneurons, but not DSA neurons, scattered in all layers. Some of these superficial neurons were chandelier (axo-axonic) cells, which have been shown to derive from late born Nkx2.1-expressing cells in a restricted region of the median ganglionic eminence⁴². Another unique and important feature of ID2/Nkx2.1 interneurons is that their relative prevalence, compared to other interneurons, is ten-fold higher in the primate neocortex compared to the mouse²³, suggesting an expanded role for this neuron type in a more complex primate neocortical mantle.

DSA neurons were more weakly excited by neighboring pyramidal cells, compared to other interneurons, but they might receive excitatory inputs from other sources as well. At the same time, they were more strongly inhibited by PV, Sst, and to a lesser extent, by NOS-expressing interneurons, relative to other interneurons. During the DOWN state of slow oscillations, all other neurons are silent, with exceptional sporadic spiking of some pyramidal cells⁸. Therefore, the source or mechanisms which sustains spiking of DSA neurons remains unknown. One possibility is that upon release from inhibition ID2/Nkx2.1 interneurons can induce rebound spikes. Although ID2/Nkx2.1 interneurons did not display biophysical properties of rebound spiking, in our optogenetic experiments, DSA neurons emitted excess spikes for 50 to 200 ms upon release from PV and Sst-induced silencing, a duration that typically corresponds to that of typical DOWN states. A second option is that intrinsic properties of ID2/Nkx2.1 interneurons maintain sustained spiking at resting membrane potential. Our intracellular recordings *in vitro* did not support this scenario but it remains to be examined whether ID2/Nkx2.1 interneurons possess membrane channels that endow them with special properties to generate action potentials at rest. Yet another explanation is that hitherto unidentified subcortical sources, specifically innervating deep layer ID2/Nkx2.1 interneurons, are also selectively active during the DOWN state. One such potential source could be some of the intralaminar and midline nuclei of the thalamus, several of which preferentially innervate deep cortical layers and have been related to arousal and awareness⁴³. Another subcortical source is the cholinergic nucleus basalis, whose neurons display low level but phasic activity during NREM sleep¹. ID2/Nkx2.1 cells may

respond differently to acetylcholine, which is known to suppress the activity of superficial layer neurogliaform cells ⁴⁴.

Effector characteristics of ID2/Nkx2.1 interneurons

Our optogenetic activation of ID2/Nkx2.1 interneurons exerted relatively modest effects on the activity of nearby recorded neurons. This is perhaps not surprising given their 10 to 20-fold less density compared to PV and Sst interneurons in deep cortical layers ^{20,22}. Therefore, focal light delivery likely excited relatively few ID2/Nkx2.1 cells, which may explain why their optogenetic excitation exerted a relatively weak effect on other cells, compared to activation of other interneurons.

Yet, a clear indication of the important contribution of DSA neurons to the DOWN–UP transition is illustrated by our observation that the spike timing of layer 6 neurons suppressed by optogenetic activation of ID2/Nkx2.1 neurons occurred later during the transition from DOWN to UP state compared to those neurons which were not affected by the optogenetic pulses. This effect may also explain why during optogenetic activation of ID2/Nkx2.1 neurons, the delta frequency peak decreased, an indication of prolonged DOWN states. Furthermore, within the ID2/Nkx2.1 neuron-suppressed group, neurons spiked later in the DOWN-UP transition when in the preceding DOWN state spiking activity of DSA neuron was high compared to absence or low spiking activity.

A simple mechanism that could account for the impact of DSA neurons on the sequential organization of neurons in the wake from the DOWN state ¹⁰ is a non-uniform innervation of their targets and/or the different biophysical properties of their target deep layer pyramidal cells. This latter hypothesis is supported by previous observations showing that the rank order of pyramidal neuron firing during the DOWN-UP transition correlates with their baseline firing rates ³². The potential targeting of thalamus-projecting layer 6 pyramidal cells by ID2/Nkx2.1 neurons might be of special significance. Previous work has shown that these layer 6 neurons play a critical role in cortical gain control by affecting activity in all other layers directly and indirectly, through the corticothalamic circuit ⁴⁵.

Additional support for the importance of DSA neurons in affecting populations of neurons comes from our optogenetic manipulation of these neurons in a memory task. While remembering cue discrimination benefited from intervening rest/sleep between morning and afternoon sessions, this advantage was abolished by bi-hemispheric optogenetic activation of ID2/Nkx2.1 interneurons in PPC, specifically during NREM sleep. Non-specific effects of light stimulation, such as activation of the retina or heat activation of nearby neurons, are not likely since the same intensity of light stimulation on the ID2/Nkx2.1-GFP mice were not different from non-stimulated controls. One possible explanation for abolishing the memory consolidation benefit of sleep is that optogenetic perturbation of DOWN states in PCC, a region specifically involved in visual discrimination learning³⁴, interfered with the physiological patterning of neural firing. Support for this contention was provided by the finding that replay of neuron pairs active during maze learning was significantly stronger during control days than on days when ID2/Nkx2.1 neurons were optogenetically stimulated during NREM. Although it is tempting to suggest that perturbation of the sequential order of firing of pyramidal neurons, brought about by artificial activation of ID2/Nkx2.1 interneurons is related to the memory deficit³⁵, further experiments are needed to firmly establish this potential link. The modest goal of our behavioral experiment was to show that artificial activation of DSA neurons can perturb memory consolidation and, thus, demonstrate the physiological effectiveness of ID2/Nkx2.1 interneurons. It is expected that the physiological consequences of ID2/Nkx2.1 neuron perturbation in different cortical regions will depend on the specific circuit function of the perturbed region.

Conclusions

We demonstrate that deep layer neocortical ID2/Nkx2.1 interneurons comprise a distinct neuron type, characterized by an inverse activity relationship with all other cortical neurons. They share physiological features of the broader class of neurogliaform neurons, yet they possess unique and distinct features. ID2/Nkx2.1 neurogliaform cells are strongly inhibited by several other interneuron types. High activity of DSA neurons delayed the onset of DOWN-UP transition and affected the sequential ordering of neurons, and their optogenetic activation in the posterior parietal cortex during nonREM sleep interfered with consolidation of cue discrimination memory. It may be of special significance that the prevalence of ID2/Nkx2.1 interneurons

increases with the evolution of neocortex. Thus, despite their small numbers, the physiological importance of ID2/Nkx2.1 interneurons is high.

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Author Contributions

MV and GB designed the experiments. MV, SM, IZ and YS performed the *in vivo* physiological experiments, BS performed the *in vitro* physiological experiments, TV and MV did the anatomical characterization of DSA neurons, SM and IZ helped with behavioral testing. RM and BR designed and performed the breeding of the ID2/Nkx.21 line, MV generated *in silico* experiments and analyzed results. GB and MV wrote the paper with contributions of all authors.

Declaration of Interests

The authors declare no competing interests.

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Figure Legends:

Figure 1. Identification and physiological properties of DOWN-State Active (DSA) neurons in the PPC

(A) Histological verification of probe shank locations (dashed white lines) in Posterior Parietal Cortex (PPC; left). Immunoreactivity for VGLUT2 delineates layers (reverse contrast) on a coronal section of the PPC (right). (B) LFP depth-profile (gray) recorded with one of the shanks in PPC during a 6-sec slow-wave sleep epoch and a 4-sec waking epoch during waking quiescence [QWake] and walking, with simultaneously recorded spikes of isolated units (each dot represents a single spike; red, pyramidal cells; blue, interneurons; black, a single DSA neuron). Light gray LFP segments correspond to identified DOWN-states. (C) Average current source density (CSD) map and superimposed LFP of all the detected DOWN-states from the recording session illustrated in B. Depth distribution of high-frequency ($>120\text{Hz}$) spectral power identifies layer 5. Vertical line, time maximum of the layer 5 source. (D) Peri-DOWN-state Z scored firing raster plot for all putative principal cells (PYR, left), putative interneurons (INT, bottom-middle) and DOWN-state active (DSA) neurons (top-middle), as ranked according to their magnitude of event response (right). Dashed lines delimit the $\pm 50\text{ms}$ window used to estimate firing rate. Note that DSA neurons increase their rate selectively during DOWN-state. SD, standard deviation (Z-scored). (E) Similar layout as in D but for the average spike cross-correlation (CCG) between a reference neuron and all other simultaneously recorded units in NREM sleep (left) and quiet waking (QWake, right). Dashed lines delimit the $\pm 35\text{ms}$ window used to quantify CCG neuron response (right). (F) Distributions of DOWN-state/UP-state firing rate ratio for putative pyramidal cells (PYR, red), interneurons (INT, blue) and DSA neurons (gray) ($P < 10^{-80}$, Kruskal-Wallis (KW) test). (G) Magnitude of DOWN-state response as a function of the CCG response. Grey dots show 47 units (out of 2279) defined as DSA neurons. Marginal CCG (top; $P < 10^{-34}$, KW test) and DOWN-state response (right; $P < 10^{-41}$, KW test) distributions for pyramidal cells, interneurons and DSA neurons (top) in each group (right). (H) Top: average firing auto-correlogram (top-left; mean \pm 95% confidence interval, IC95) and median burst index (medians, interquartile ranges, maxima and minima; $P < 10^{-19}$, KW test) for each group. Bottom: average spike-waveform (mean \pm IC95) and through-to-peak spike duration (spk dur; $P < 10^{-306}$, KW test) for each group ($n = 1457$ pyramidal cells, 775 interneurons 47

DSAns from 11 mice). **(I)** Average high-frequency ($>120\text{Hz}$) spectral power depth profile (mean \pm IC95) was used to align all sessions, relative to mid-layer 5 as reference. Depth distribution for all unit (right; dots are scattered in the x-axis to allow better visibility). DSA neurons are located in deep cortical layers ($P < 10^{-15}$, KW test). Arrowhead, DSA neuron shown in **B**. *** $P < 0.001$.

Figure 2: Network properties of the DSA neurons

(A) Log distribution of average firing rates (top, $P < 10^{-191}$, KW test) and median firing rates (circles; lines, interquartile ranges) of pyramidal cells, interneurons ($n = 1457, 775$ and 47 cells, respectively, from 11 mice) and DSAns during waking quiescence (QWake), NREM sleep, REM sleep and active behavior (Walk) (bottom). **(B)** Average unit CCGs as a function of brain states. Top insets, mean CCG (mean \pm IC95) for all groups in **A** across brain states. **(C)** Average joint Z-score rate density between pyramidal cells, interneurons and DSA neurons. Top, Firing rates of DSA neurons are more strongly anti-correlated with interneuron rates than with firing rates of pyramidal cells (mean \pm IC95; $n = 32$ session with at least one DSA, $P < 10^{-18}$, $F(1, 564) = 78.75$, repeated measures ANOVA and Spearman correlation (ρ) tested using a Student's t distribution). **(D)** Average (mean \pm IC95) partial correlation values of the Z scored average rate for all groups (blue for $\rho_{\text{DSA,INT}}$ controlling for PYR; red for $\rho_{\text{DSA,PYR}}$ controlling for INT; magenta for $\rho_{\text{PYR,INT}}$ controlling for DSA, $n = 32$ sessions, $P < 10^{-27}$, $F(2, 2787) = 61.23$, repeated measures ANOVA). **(E)** Cross-correlation histograms of an example putative monosynaptic between a pyramidal cell–interneuron pair (top; red, pyramidal cells; blue, interneurons) and a pyramidal-DSA neuron pair (bottom; gray star, DSA neuron). Insets show auto-correlogram for each pair member (± 20 ms). Green bins depict 1-3ms monosynaptic window. **(F)** Z scored cross-correlograms for all identified monosynaptic pairs between PYR-DSA, PYR-INT and PYR-PYR. Bottom, average spike transmission probability (trans prob, mean \pm IC95) for each group. **(G)** Group difference (medians, interquartile ranges, maxima and minima) of spike transmission probability for all detected putative monosynaptic pairs in **F** ($n = 149, 957$ and 17 for PYR-PYR, PYR-INT and PYR-DSA, respectively; $P < 10^{-6}$, KW test). **(H)** Upper panels, DOWN-state responses (left) and average CCG (right) for all simultaneously recorded units in a representative session with two DSA neurons (arrowhead). Bottom, Z-scored within-cell type cross-correlograms (mean \pm IC95, all cell pairs from the same group recorded

simultaneously: $n = 82972$ PYR-PYR, 17392 INT-INT and 18 DSA_n-DSA_n pairs, $P < 10^{-292}$ KW test). * $p < 0.05$, ** $p < 0.01$; *** $p < 0.001$.

Figure 3: Cell type identity of the DSA_n and their inhibitory inputs.

(A) Schematic of the opto-tagging experiments. Combined light fiber-recoding probes were implanted in promoter-specific transgenic mice expressing excitatory (ChR2) opsins. (B) Peristimulus histogram (PSTH) for all cells recorded in the Sst::ChR animals ($n = 2$ mice). Each row is a rate-normalized color coded peristimulus histogram of a unit. 21% were classified as Sst::ChR expressing cells (top in middle panel). Bar graph (bottom right), probability (prob) of optogenetically di(poly)-synaptically silenced neurons (mod, from modulated). (C) Same as a B for PV::ChR mice ($n = 4$ mice in V1). (D) Same as B for Nos1::ChR mice ($n = 3$ mice). (E) Same as in B for the ID2/Nkx2.1::CatCh mice ($n = 6$). (F) Average optogenetic response for all recorded DSA neurons in the different promoter-specific transgenic mouse lines. Most of the DSA neurons were suppressed (– Mod) upon Sst:ChR and PV::ChR stimulation ($P < 0.01$ for all units; Wilcoxon's paired signed-rank test). A majority of the DSA neurons were recruited with light pulses in the ID2/Nkx2.1::CatCh line. (G) Spike auto-correlogram (left) and spike waveform (right) from ID2/Nkx2.1-expressing DSA neurons are similar to those of DSA neurons observed in other mice (mean \pm IC95 in black; $P = 0.38$ and $P = 0.49$, respectively; KW test).

Figure 4. Intrinsic electrophysiological features of L5/6 ID2/Nkx2.1 neurons.

(A) Example voltage traces for a L5/6 ID2/Nkx2.1 cell (gray) and a L1 neurogliaform cell (L1 NGFC, from ref²⁶ green) during suprathreshold (top), just supra- and just-subthreshold (middle), and hyperpolarizing (bottom) current injections (currents shown in black). (B) Six electrophysiological parameters for both L5/6 Id2;Nkx2.1 (gray) and L1 neurogliaform (green) cells. All plots show individual cells as circles (Time to first spike: $n = 22$ and 27 for ID2/Nkx2.1 and L1 NGFC, respectively, $P = 0.004$; adaptation: $n = 20$ and 26 , $P = 0.001$; spike width: $n = 22$ and 26 , $P < 10^{-6}$; sag amplitude: $n = 20$ and 22 , $P = 0.48$; input resistance: $n = 22$ and 25 , $P = 0.002$; resting potential: $n = 18$ and 27 , $P < 10^{-6}$; two-sided Wilcoxon's rank-sum test), mean values as bars, and SEM values as small bars on right. ** $p < 0.01$; *** $p < 0.001$.

Figure 5: Anatomical features of ID2/Nkx2.1 interneurons in the PPC.

(A) Left, sparse distribution of tdTomato+ somata in the PPC of an ID2/Nkx2.1::Ai65 mouse (70µm-thick coronal section, low power widefield epifluorescence image, reverse contrast). Two of them are marked by arrowheads. Top right, Depth distribution of ID2/Nkx2.1 neurons (mean \pm SE). Grey shaded area on the top graph shows depth distribution of DSA (same as Fig. 1I, DSA). Note that the majority of tdTomato+ neurons were found below layer 5 (3 section per mouse for 3 mice, 9 total samples overlaid as blue dots) and were immunoreactive for nNOS (bottom). (B) Two tdTomato+ ID2/Nkx2.1 neurons in L6, weakly immunoreactive for nNOS (white arrowheads; n = 54 tdTomato+ neurons from 3 mice were tested). Note also a strongly nNOS-immunopositive neuron (hollow arrowhead), which is immunoreactive for both calretinin (CR) and Sst but does not express tdTomato (20.8-µm thick confocal maximum-intensity z-projection; n = 12 tdTomato+/nNos+ cells). (C) Axon distribution in deep cortical layers (widefield epifluorescence, reverse contrast) from three tdTomato+ ID2/Nkx2.1 neurons. Note high axon density in L6. Right, detail of L6b. Two somata are indicated by asterisks. (D) Reconstruction of somata, dendrites and axonal arborization of an intracellularly-filled tdTomato+ ID2/Nkx2.1 neuron (from a 183.3 µm thick confocal stack. Inset: detail of the axonal varicosities; three additional filled tdTomato+ ID2/Nkx2.1 cells, one of them reconstructed, in Extended Data Fig5D). wm, white matter.

Figure 6: Effector role of DSA neurons. (A) Depth distribution of pyramidal cells (red, right abscissa) and interneurons (blue, left), separated into groups of negatively modulated cells (–Mod, light red and light blue) or non-modulated neurons (No Mod, dark colors) upon optogenetic stimulation of ID2/Nkx2.1::CatCh neurons (PYR: n = 44 and 410, respectively, $P < 10^{-6}$; n = 74 and 238 for INT, $P < 10^{-15}$, KW test). ID2/Nkx2.1::CatCh modulated cells are located in deep layers. (B) Temporal dynamics of the Z-scored DOWN-UP events responses for –Mod and No Mod pyramidal cells and interneurons (top-left for PYR, bottom-left for INT; mean \pm IC95). Right panels, timing ($P < 10^{-18}$, KW test) and amplitude ($P < 10^{-5}$, KW test) response group differences, respectively (medians, interquartile ranges, maxima and minima). (C) DOWN-UP events were classified as ‘LOW DSA rate events’ or ‘HIGH DSA rate events’ using the average DSA neuron rate during the DOWN epoch of the DOWN-UP event as the separatrix (top: medians, interquartile ranges, maxima and minima; bottom, mean \pm IC95; 32 sessions with at least one DSA neuron from 11 mice, $P < 10^{-7}$, Wilcoxon’s paired signed-rank test). (D)

DOWN-UP temporal group response (top; medians, interquartile ranges, maxima and minima, $P < 10^{-12}$, KW test) and High-Low DSA firing events difference distributions (bottom; $P < 10^{-7}$, KW test) for -Mod (yellow, $n = 51$ units from 9 session with at least 1 DSA from 6 ID2/Nkx2.1::CatCh mice) and No Mod (green, $n = 195$ units) groups. (E) Illustration of mean rank order values for 'HIGH' and 'LOW' rate events, used as templates against remaining DOWN-UP events ($n, n+1, \dots$). (F) Top, Z-scored sequential activity example of a simultaneously recorded population during DOWN-UP events with 'Low DSA rate' (session mean template), vertically arranged by latency, including two DSA neurons bottom two rows). Bottom, similar session mean template for 'High DSA rate' events, but arranged according to 'Low DSA rate' classification. (G) Rank correlation values within 'HIGH' and 'LOW' DSA events, HIGH vs LOW events, randomly chosen events and slow oscillation events in the first vs the second halves of NREM sleep episodes (Mean \pm SD; $n = 32$ sessions with at least one DSA from 11 mice; $P < 10^{-20}$, Friedman test; $P < 10^{-7}$ and $P < 10^{-8}$ for 'within-LOW' against 'HIGH vs LOW' and 'within-HIGH' against 'HIGH vs LOW', respectively, Tukey's test). * $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$.

Figure 7: Perturbation of DSA neurons during NREM sleep impair memory consolidation

(A) Diode/probes were implanted bilaterally in PPC in ID2/Nkx2.1::CatCh mice. They performed in a goal-associated visual cue discrimination task (e.g., blue and yellow stripe patterns) twice daily (morning and afternoon sessions). (B) Performance of each mouse in the morning (session 1) and afternoon (session 2). On randomly chosen days mice were left either unperturbed in the home cage between sessions (grey symbols) or ID2/Nkx2.1 neurons were optogenetically stimulated (100ms duration light pulses every 500 ± 400 ms) during NREM sleep (purple). As further controls, the same optogenetic stimulation during wake in the home cage (green) or during the task in the central arm continuously (yellow) did not interfere with sleep-induced memory improvement. Each symbol depicts a comparison between morning (session 1) and afternoon (session 2) session (3 days averaged per condition). (C) Only optogenetic perturbation of ID2/Nkx2.1 neurons during NREM sleep affected memory consolidation (mean \pm SD, $P = 0.001$, KW test; $P < 10^{-4}$ for 'No stim' against 'NREM stim', Tukey's test). (D) Same stimulation protocol in a different cohort of animals that lacked ChR ('Photostimulation control') did not affect memory consolidation (mean \pm SD; $n = 3$; $P = 0.8946$, KW test). (E) Example of

rate correlation matrices for all neuron pairs from pre-NREM sleep, maze (first session) and post-NREM sleep in an example mouse (non-stimulation, control day). These matrices were used to calculate the explained variance (EV) and its control value, the reverse explained variance (REV) in both non-stimulated and optogenetically stimulated sessions. **(F)** Pearson's correlation coefficients for all neuron pairs for pre-sleep versus Maze learning (left) and postsleep versus Maze learning (right) comparison (same example session as in **E**). **(G)** Group statistics (medians, interquartile ranges, maxima and minima; $n = 5$ mice, at least one session per animal) of the EV and REV on non-stimulated (control) and NREM-stimulated days ($P = 0.002$, Kruskal-Wallis test). EV on non-stimulated days was significantly higher than its corresponding REV (chance, $P = 0.003$, Tukey's test), and also significantly higher than EV on NREM-stimulated days ($P = 0.012$). $*p < 0.05$; $***p < 0.001$.

761 **METHODS**
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Subjects and electrode implantation. All experiments were approved by the Institutional Animal Care and Use Committee (IACUC) at New York University Medical Center. We used several lines of transgenic mice (n = 15 male mice) for optogenetic tagging of the recorded units: n = 2 Sst-Cre (Sst^{tm2.1(cre)Zjh}/J, Jax stock number: 013044), n = 4 PV-Cre::Ai32 (from B6.129P2-Pvalb^{tm1(cre)Arbr}/J, Jax stock number: 017320, and B6.Cg-Gt(ROSA)26Sor^{tm32(CAG-COP4*H134R/EYFP)Hze}/J, Jax stock number: 024109), n = 3 Nos1-Cre (B6.129-Nos1^{tm1(cre)Mgmj}/J, Jax stock number: 017526); n = 6 ID2-CreER/Nkx2.1-Flpo::Ai80 (from B6.129S(Cg)-Id2^{tm1.1(cre/ERT2)Blh}/ZhuJ, Jax stock number: 016222, Nkx2-1^{tm2.1(flpo)Zjh}/J, Jax stock number: 028577, and B6.Cg-Gt(ROSA)26Sor^{tm80.1(CAG-COP4*L132C/EYFP)Hze}/J, Jax stock number: 025109) for optogenetic tagging of the recorded units. Tamoxifen (20 mg/ml in corn oil) was administered to ID2 /Nkx2.1::Ai80 (or Ai65, for the histology section) animals (3 × 5 mg by oral gavage over 5 days) between P30–P40 to activate the CreER. Sst-Cre and Nos1-Cre mice were injected with AAV-FLEX-ChR2-EYFP to express channelrhodopsin-2 in Cre-expressing neurons. Three ID2 /Nkx2.1::Ai80 mice administered only with corn oil (without Tamoxifen) were used as behavioral photostimulation controls.

All animals (28-35 gr, 3-10 months old) were implanted with 64-site silicon probes (NeuroNexus or Cambridge NeuroTech) in the posterior parietal cortex (AP 2.0 mm, ML 1.75 mm, DL 0.6 mm) or V1 (AP 1.0 mm, ML 2.5 mm, DL 1mm, with a 21° angle from dorso-ventral axis and 10° from the mediolateral axis)⁴⁶. Ground and reference wires were implanted in the skull above the cerebellum, and a grounded copper mesh hat was constructed shielding the probes. Probes were mounted on microdrives that were advanced to layer 6 over the course of 5-8 days after surgery. For most of the experiments, a 100-200 μm fiber optic was attached to one of the shanks of the silicon probe. For V1 recordings, an optic fiber was placed right above the skull over the implantation site. The back end of the fiber was coupled to a laser diode (450 nm blue, Osram Inc.). For ID2 /Nkx2.1::Ai80 recordings, a second fiber was implanted in the contralateral hemisphere at the same coordinates. The probe locations were verified histologically for all mice. Animals were allowed to recover for at least one-week prior to testing. Mice were housed under standard conditions (71-73°F and 40-50% relative humidity) in the animal facility and kept on a 12 h reverse light/dark cycle.

Recordings and behavior. Recording session started 1-2 hr after the onset of the dark phase of the light/dark cycle. We recorded from the mice while they slept or walked around freely moving in the home cage. Electrophysiological data were acquired using an Intan RHD2000 system (Intan Technologies LLC) digitized with 30 kHz rate. The wide-band signal was downsampled to 1.25 kHz and used as the LFP signal.

For optogenetic tagging of specific neuron types, blue laser light (450 nm, Osram Inc) pulses were delivered in PPC or above the V1. The maximum light power at the tip of the optic fiber was 1 to 4 mW. 10 – 100 ms light pulses with 40%, 70% and 100% of the maximum power were delivered (n = 500 - 1000 times at each intensity, at 1 Hz or at 400 ± 200 ms random intervals).

For behavior experiments, animals were handled daily and accommodated to the experimenter, recording room and cables for 1 week before the start of the experiments. Prior to the start of the experiment, animals were water restricted. Two (morning and afternoon) 100 trials (30 to 60-minute-long) behavior sessions were conducted daily, separated by 5 hr, and preceded and followed by 1 hour-long sleep sessions. In the goal-associated visual cue discrimination task, mice run in a 70 cm \times 40 cm (one meter above the floor) figure-8 maze to collect water reward each time they reached the correct (left or right) corner (Fig. 7). Visual cues were arbitrary patterns delivered by two (left and right) 8 \times 8 LEDs matrices (Adafruit) at the initial section of the central arm. For each trial, the visual cue/ reward position was randomly assigned, while a different arbitrary pattern (distractor – but same number of LEDs as the cue) was delivered in the LED matrix on the opposite side. Visual cues and distractor patterns were chosen to activate similar number of single LEDs from the matrices and were randomly assigned to the different experimental subjects so that the visual cue for one animal was the distractor of a different animal. The same pair visual cue/distractor was maintained during training, but once the animals learned the task the visual patterns varied from day to day, demanding new learning each day. All subjects underwent the same number of 'No stim' and 'NREM' stimulation sessions in a randomly assigned fashion (and the same for Maze stim and Awake stim, in addition to No stim and NREM, for mice 1-3). Performance was quantified as the correct choice probability: correct choices/total number of trials. Maze doors (to prevent backward displacement of the mice), water delivery, visual patterns and optogenetic stimuli were controlled by a custom-made Arduino-based circuit (<https://github.com/valegarman>).

Unit clustering and neuron classification. Spike sorting was performed semi-automatically with KiloSort⁴⁷ (<https://github.com/cortex-lab/KiloSort>), using our own pipeline KilosortWrapper (a wrapper for KiloSort, DOI; <https://github.com/brendonw1/KilosortWrapper>). This was followed by manual adjustment of the waveform clusters using the software Phy (<https://github.com/kwikteam/phy>) and plugins for Phy designed in the laboratory (<https://github.com/petersenpeter/phy-plugins>). Following parameters were used for the Kilosort clustering: ops.Nfilt: 6 * numberChannels; ops.nt0: 64; ops.whitening: 'full'; ops.nSkipCov: 1; ops.whiteningRange: 64; ops.criterionNoiseChannels: 0.00001; ops.Nrank: 3; ops.nfullpasses: 6; ops.maxFR: 20000; ops.fshigh: 300; ops.ntbuff: 64; ops.scaleproc: 200; ops.Th: [4 10 10]; ops.lam: [5 20 20]; ops.nannealpasses: 4; ops.momentum: 1./[20 800]; ops.shuffle_clusters: 1. Unit clustering generated two separable groups (Extended Data Fig. 1A) based on their autocorrelograms, waveform characteristics¹⁶ and firing rate. Putative pyramidal cells and interneurons were tentatively separated based by these two clusters. Definitive cell identity was assigned after inspection of all features, assisted by monosynaptic excitatory and inhibitory interactions between simultaneously recorded, well-isolated units and optogenetic responses. DOWN-state active (DSA) neurons were identified by the Z scored ± 50 ms DOWN-state and ± 35 ms CCG response (Fig. 1G). Another method of separation of the DSA neurons was obtained by the firing rate ratios during DOWN and UP states (Fig. 1F).

Unit activity analysis. Units were defined as optically tagged^{16,46}, using a p value cutoff of 10^{-3} . Negatively modulated cells (– Mod) were identified by an average firing suppression less than 1 SD for the entire window of light stimulation (50-100 ms) and a p value cutoff of 10^{-2} . Average unit CCG was computed as the mean from all Z-scored cross-correlograms (CCGs; 5 ms binned, 0.6 s window) sharing same cell as reference and spike groups in different brain states. To calculate the depth coordinates of the units, we aligned depth profiles of electrophysiological landmarks. These included the largest amplitude peak of high-frequency LFP power (500 Hz - 5 kHz) corresponding to mid-layer 5 and four prominent sinks and sources from the averaged DOWN-UP current-source density (CSD) maps for each animal (see Fig. 1C). To investigate the rate relationship between unit groups we used joint probability density of the Z-scored rate of a given group (DSA neurons in Fig. 2C and 100 shuffling average of the 10%

pyramidal cells or interneuron in Extended Data Fig. 2C) between Z-scored firing of the pyramidal cells and interneurons in 0.25 SD bins. We also computed the partial Spearman correlation of the Z-scored spike counts across groups for different bin sizes (0.01, 0.15, 0.20... 1.01; ref⁴⁸).

Rank order tests were used to identify sequences of units repeating consistently across DOWN-UP events as tested against chance order⁴⁹. The timing from multiple units recorded simultaneously were transformed in a normalized sequence order from 0 to 1 within a DOWN-UP event. Only events with at least 6 participating units were included. Only the first spike from each unit in a cycle was considered for the rank order. Rank distribution of the correlation values was tested against shuffle correlations (500 sufflings, significance at p=0.05) using Pearson correlations. Only events with more significant correlations than expected by chance were considered for further analysis. Finally, significant events were compared against the average rank order obtained by grouping the remaining events according to some rules (templates for DSA neuron rate higher than average and DSA neuron rate lower than average; or first half of the recording and second half of the recording session; or randomly chosen events) using again Pearson correlation.

Explained variance (EV) and reverse explained variance (REV) were calculated only in sessions with more than 20 simultaneously recorded neurons. Pairwise correlations for EV and REV were calculated using the Pearson correlation coefficient in 100-ms-binned spike trains. The coefficients were separately calculated for pre-sleep (only NREM), maze (only first session), and post-sleep (only NREM) epochs as correlation matrices (Fig. 7A). The correlations between these three matrices were used to assess the percentage of variance in the post-sleep period that could be explained by the patterns established during the maze experience while accounting for pre-existing correlations in the pre-sleep session (EV), according to the following equation:

$$EV = R_{M,P2 \vee P1}^2 = \left(\frac{R_{M,P2} - R_{M,P1} \times R_{P2,P1}}{\sqrt{(1 - R_{M,P1}^2) - (R_{P2,P1}^2)}} \right)^2$$

where R variables are the correlation coefficients between maze (M), pre-sleep (P1) and post-sleep (P2) pairwise correlation matrices³⁵. The control value (REV) was obtained by switching the temporal order of the pre- and post-sleep session.

Detection of monosynaptic connections. CCG peaks can reflect monosynaptic connections¹⁹. This takes the form of short time-lag (1–4 ms) positive deviation from CCG baseline indicating putative excitatory connections. Such detection is based on testing the null hypothesis of a homogeneous baseline at short timescale⁵⁰. To detect putative monosynaptic connections, 0.5 ms binned CCG were convolved with a 7 ms standard deviation Gaussian window resulting in a predictor of the baseline rate. At each time bin, the 99.9999 percentile of the cumulative Poisson distribution (at the predicted rate) was used as the statistical threshold for significant detection of outliers from baseline. A putative connection was considered significant when at least two consecutive bins in the cross-correlograms within +1.5 to +4 ms passed the statistical threshold¹⁹. All putative connections were visually inspected. Corrected spike transmission probability was calculated as described in English et al¹⁹.

Brain state scoring. Brain state scoring was performed as in Watson et al.¹⁷ with the MATLAB resource SleepScoreMaster from the repository of the laboratory (<https://github.com/buzsakilab/buzcode/tree/master/detectors/detectStates/SleepScoreMaster>). In short, spectrograms were constructed with a 1 s sliding 10 s window FFT of 1250 Hz data at log-spaced frequencies between 1 and 100 Hz. Three types of signals were used for score state: broadband LFP, narrowband theta frequency LFP, and electromyogram (EMG). For broadband LFP signal, principal components analysis (PCA) was applied to the z-transformed (1-100 Hz) spectrogram. The first PC in all cases was based on power in the low (< 20 Hz) frequency range and had oppositely weighted power in the (> 32 Hz) higher frequencies. Theta dominance was taken to be the ratio 5-10 Hz and 2-16 Hz power from the spectrogram. EMG was extracted from the intracranially recorded signals by detecting the zero time-lag correlation coefficients (r) between 300-600 Hz filtered signals (using a Butterworth filter at 300 – 600 Hz with filter shoulders spanning to 275 – 625 Hz) recorded at all sites. All states were reviewed by the experimenter and corrections were made when discrepancies between automated scoring and user assessment occurred.

Detection of UP and DOWN states. Slow waves were detected using the coincidence of a three-stage threshold crossing in three signals: a drop in high gamma power (30-600 Hz), a peak in the delta-band filtered signal (0.5-8 Hz) and drop in units normalize firing rate. The gamma power signal was smoothed using a sliding 100-ms window, and locally normalized to account for non-stationaries in the data. Time epochs in which the delta-filtered signal crossed a peak threshold (.5 SD) and coincide with gamma power drops (-0.5 SD) were considered as candidate delta peaks and binned by peak magnitude, with start and end times at the nearest crossing of the delta band threshold with a minimum duration of 40 ms. Time epochs with gamma power above 1 SD and lasting 0.2-1.4 s were considered putative UP events. The peri-event time histogram (PETH) for spikes from all cells was calculated around delta peaks in each magnitude bin and Z scored normalized. Only candidate events with a rate drop below 2 SD in a ± 50 ms window around the delta peak were considered DOWN states. UP states were defined as candidate epoch that immediately followed and were truncated by DOWN states. DOWN-state responses were aligned by the delta peak or the end of the DOWN epoch (DOWN-UP transition), as indicated.

Spiking network simulations. We built a network model of leaky integrate-and-fire neurons based in Jercog et al ¹⁵, and Barrio et al ⁵¹, using the Python-based simulator Brian2 ⁵². We include $n_{\text{PYR}} = 4000$ excitatory, $n_{\text{INT}} = 1000$ inhibitory and $n_{\text{DSA}} = 100$ neurons with the following connection probability (c): $c_{\text{PYR-PYR}} = 0.1$, $c_{\text{PYR-INT}} = 0.1$, $c_{\text{INT-INT}} = 0.2$, $c_{\text{INT-PYR}} = 0.3$, $c_{\text{PYR-DSA}} = 0.05$, $c_{\text{INT-DSA}} = 0.3$, $c_{\text{DSA-PYR}} = 0.1$ and $c_{\text{DSA-INT}} = 0.1$. In our model, the membrane potential of a neuron (i) obeys:

$$\tau_i \frac{dV_i}{dt} = -(V_i - E_L) + I_{\text{rec}}(t) + I_{\text{ext},i}(t)$$

Where τ_i is the membrane time constant, E_L the leak potential, while $I_{\text{rec}}(t)$ and $I_{\text{ext}}(t)$ are the recurrent and external input terms, respectively. The following values were used for the τ and E_L constants: $\tau_{\text{PYR}} = 20$ ms, $\tau_{\text{INT}} = 10$ ms, $\tau_{\text{DSA}} = 50$ ms and $E_L^{\text{PYR}} = -55.6$ mV, $E_L^{\text{INT}} = -55.6$ mV, $E_L^{\text{DSA}} = 40.0$ mV. Whenever the membrane potential of the neuron i reached the threshold V_{Th} , a spike is simulated, and the membrane potential was reset to V_{Res} with 1 ms of refractory period. The thresholds and resets values were $V_{\text{Th}}^{\text{PYR}} = -47$ mV, $V_{\text{Th}}^{\text{INT}} = -37$ mV, $V_{\text{Th}}^{\text{DSA}} = -40$ mV and $V_{\text{Res}}^{\text{PYR}} = -51$ mV, $V_{\text{Res}}^{\text{INT}} = -49.9$ mV, $V_{\text{Res}}^{\text{DSA}} = -57$ mV.

The recurrent input term consisted of the sum of all inhibitory and excitatory synaptic drives experienced by a given cell according to the rate of its presynaptic partners, and governed by:

$$\frac{dI_{rec}}{dt} = -\frac{g_{PYR,POST}}{\tau_E} - \frac{g_{INT,POST}}{\tau_I} - \frac{g_{DSA,POST}}{\tau_{DSA}}$$

Where τ_x are the response time constants and $g_{PRE,POST}$ are the synaptic weights, with the following values for all combinations of synapses: $g_{PYR,PYR} = 0.97$ mV, $g_{PYR,INT} = 3.5$ mV, $g_{PYR,DSA} = 0.5$ mV, $g_{INT,PYR} = -0.3$, $g_{INT,INT} = -1$, $g_{INT,DSA} = 0.0$ to -3.0 , $g_{DSA,PYR} = -0.3$, $g_{DSA,INT} = -1$, $\tau_E = 8.0$ ms, $\tau_I = 2.0$ ms and $\tau_{DSA} = 20.0$ ms.

The external input term is composed by two terms, a Gaussian white noise (ζ_i) term with $\sigma = 3$ mV and a time constant τ_{GWN} equal to the membrane time constant (τ_i), and a source of Poisson (2 Hz) input pulses with 20 ms duration and 45 mV amplitude, called kicks (I_{kicks}), affecting 10% of the population (so $p_i = 0.1$) of pyramidal cells and interneurons¹⁵:

$$I_{ext,i}(t) = \sigma\sqrt{\tau_i}\zeta + p_i I_{kicks}$$

We run five simulation of 100 s each with a $g_{INT,DSA}$ of 1.5 mV to compare with data obtained from *in vivo* recordings. For inhibitory drive parametric experiments ($g_{INT,DSA}$ from 0 to 3 mV, Extended Data Fig. 3C and 3E) we averaged data from 2 simulation for each $g_{INT,DSA}$ value (-3.0, -2.5, -2 ... 0). Code will be uploaded to <https://senselab.med.yale.edu/modeldb/>.

In vitro recordings. Whole-cell patch clamp recordings of 22 tdTomato+ cells from 7 ID2/Nk2.1::Ai65 adult (postnatal day 37 to 198, B6.129S(Cg)-Id2^{tm1.1(cre/ERT2)Blh}/ZhuJ, Jax stock number: 016222, Nkx2-1^{tm2.1(flpo)Zjh}/J, Jax stock number: 028577, and B6;129S-Gt(ROSA)26Sor^{tm65.1(CAG-tdTomato)Hze}/J, Jax stock number: 021875, respectively) mice of either sex were performed as previously described²⁶. In short, mice were killed using pentobarbital sodium and phenytoin sodium. Once unresponsive, mice were perfused with ice-cold sucrose-ACSF containing the following (in mM): 87 NaCl, 75 sucrose, 2.5 KCl, 26 NaHCO₃, 1.25 NaH₂PO₄, 10 glucose, 1.0 CaCl₂, and 2.0 MgCl₂, saturated with 95% O₂/5% CO₂. Then, 300µm-thick coronal brain sections were made using a Leica VT1200S vibratome and were incubated at 35°C for 30 min in ACSF containing the following (in mM): 125 NaCl, 2.5 KCl, 26 NaHCO₃, 1.25 NaH₂PO₄, 10 glucose, 2.0 CaCl₂, and 1.0 MgCl₂. Slices were then kept in this solution for at least 30 min at room temperature before recording. For *in vitro* recordings, slices were continuously perfused with ACSF saturated with 95% O₂/5% CO₂ and kept at 29–32°C.

All whole cell recordings were performed in the current-clamp configuration; the liquid junction potential was not corrected. Pipettes with resistances of 2–6 M Ω were made (Sutter Instrument) using borosilicate glass and were filled with potassium gluconate intracellular solution containing the following (in mM): 130 K-gluconate, 10 KCl, 10 HEPES, 0.2 EGTA, 4 Mg-ATP, and 0.3 Na-GTP, adjusted to a pH of 7.3 with KOH. Biocytin was dissolved into the intracellular solution (final concentration 0.3-0.5%) for later staining and morphological recovery (n = 11 neurons attempted). Before gaining whole-cell access, a gigaseal was obtained and the pipette capacitance was compensated. Access resistances were monitored throughout recordings and were completely compensated; cells with access resistances >35 M Ω were not analyzed. All data were collected using a Multiclamp 700B amplifier (Molecular Devices), a Digidata 1322A digitizer (Molecular Devices), and Clampex version 10.3 software (Molecular Devices); data were sampled at 20 kHz and low-pass filtered at 10 kHz.

Data were analyzed using Clampex version 10.7 software (Molecular Devices), custom written MATLAB software (MathWorks), and Excel (Microsoft). All statistical analysis (only Mann-Whitney U tests were used) was performed using custom written MATLAB software (MathWorks). Six electrophysiological properties were analyzed and compared to the same six properties of layer 1 neurogliaform cells from ref²⁶. The adaption index was calculated as the ratio of the number of spikes in the last 500 ms over the number of spikes in the first 500 ms of a positive current injection that elicited ~30 Hz firing. The time to first spike (in milliseconds) was calculated as the difference in time between the peak of the first AP and the start of the positive current injection. The AP spike-width (in milliseconds) was calculated as the difference in time between the ascending and descending phases of a spike at the voltage found at the time between spike peak and spike threshold. The amplitude of voltage sags (%) were measured using hyperpolarizations to ~-100 mV from a holding potential of ~-70mV and calculated as $100 \times (V_{\text{sag minimum}} - V_{\text{steady-state}}) / (V_{\text{sag minimum}} - V_{\text{holding}})$. The input resistance (in M Ω) was calculated using Ohm's law from averaged traces of 150 ms long negative current injections of -20 pA. The resting membrane potential (in millivolts) was measured with 0 pA current injection soon after gaining whole-cell access. Data from recorded Id2;Nkx2.1 cells was compared to previously published neurogliaform cells in L1²⁶.

Histological processing and microscopy. ID2-CreER/Nkx2.1-Flpo::Ai65 mice (n = 5) were overdosed with pentobarbital injection (100 mg/kg body weight), perfused with saline and 4% paraformaldehyde before their brains were rapidly removed. After overnight post-fixation in 4% paraformaldehyde solution, the brain was washed 3 times in 0.1 M phosphate buffered saline. Coronal sections of 70–100 μ m thickness were cut using a Leica Microsystems VT 1000S vibratome, wash 3-4 times in 0.1 M phosphate buffer (PB) and stored in 0.1 M PB with 0.05% sodium azide at 4°C. For initial visualization and documentation of tdTomato-expressing neurons, sections were mounted on glass slides in VectaShield (Vector Laboratories) under a coverslip and sealed with nail polish, then observed with epifluorescence on a widefield Leitz DMRB microscope (Leica, Openlab version 5.5.0). For the identification of tdTomato+ or GFP+ neuron molecular profiles, layer markers and putative synaptic inputs, immunohistochemistry was performed as described previously^{53,54}. Briefly, sections were permeabilized in Tris-buffered saline (TBS) with 0.1% Triton X-100 (Tx) or subjected to 2-3 rounds of freeze-thaw (FT) over liquid nitrogen after cryoprotecting in 20% sucrose. For FT sections, a TBS base solution was used throughout. Sections were blocked in 20% normal horse serum (NHS) in TBS/TBS-Tx for at least 1 h at room temperature (RT) then incubated in primary antibody solution (containing 1% NHS in TBS/TBS-Tx) for at least 3 nights at 4°C. Control sections from C57Bl6j mice that lacked the primary antibodies were used throughout. We used the following primary antibodies (species, dilution, company, code): We used the following primary antibodies (species, dilution, company, code): Sst, rat clone YC7 antibody (monoclonal), 1:500, EMD Millipore Corporation, MAB354; nNOS, rabbit, 1:1000, EMD Millipore, AB5380; calretinin (CR), goat, 1:1000, Swant, CG1; RFP, rat, 1:500, ChromoTek, 5f8-100; Vesicular glutamate transporter 2 (VGLUT2), guinea pig, 1:2000, Synaptic Systems, 135 404; PV, goat, 1:1000, Swant, PVG-214; PV, guinea pig, 1:5000, Synaptic Systems, 195 004; GFP, rabbit, 1:500, ThermoFisher, A-11122; NPY, sheep, 1:1000, EMD Millipore Corporation, AB1583.. Specificity information is provided in ref⁵⁵. Sections were washed 3 times in TBS/TBS-Tx and incubated in secondary antibody solution in 1% NHS in TBS/TBS-Tx for at least 3 h RT or overnight at 4°C. Secondary antibodies raised in donkey were as previously described^{53,54}: donkey anti-rabbit Alexa Fluor 405, 1:250, Invitrogen; A31556; donkey anti-mouse, anti-goat or anti-guinea pig DyLight405, 1:250, Jackson ImmunoResearch Laboratories, 715-475-151, 705-475-147, 706-475-148, respectively; donkey anti-rabbit or anti-goat Alexa Fluor 488, 1:1000,

Invitrogen, A21206 and A11055, respectively; donkey anti-rabbit, anti-goat, anti-sheep, anti-mouse or anti-guinea pig Cy3, 1:400, Jackson; 711-165-152, 705-165-147, 713-165-147, 715-165-151 and 706-165-148, respectively; donkey anti-rabbit, anti-goat, anti-mouse or anti-guinea pig Cy5, 1:250, Jackson; 711-175-152, 705-175-147, 713-175-147, 715-175-151 and 706-175-148, respectively. Overviews of multi-channel immunofluorescence were acquired using widefield epifluorescence either on an AxioImager.Z1 (Carl Zeiss, AxioVision 2009 software version Rel.4.8.1) or on the Leitz DMRB microscope. Confocal microscopy (LSM 710; Carl Zeiss, ZEN 2008 version 5.0) was used to document tdTomato+ neurons and their putative inputs, as described previously⁵⁴. Native tdTomato fluorescence was superior to that of the RFP antibody. For the quantification of tdTomato+ cell density, PPC layers in reference sections were defined by immunoreactivity for PV and VGLUT2⁵⁶. Cell density was also measured in the anterior/rostral V1 (-2.6 to -2.8 mm posterior of Bregma), caudal/posterior V1 (-3.4 to -3.8 mm posterior of Bregma) and mPFC. For mPFC, counts were in range of 1.9 to 2.1 mm anterior of Bregma, measuring in anterior cingulate, prelimbic and infralimbic, and medial orbital cortex.

In vitro labeling and morphological imaging of intracellularly recorded neurons were described previously²⁶. In short, cells (n = 11 attempted) were recorded with an intracellular K-gluconate solution containing 0.3-0.5% biocytin. After recording, slices were immediately placed into a 4% PFA solution and stored at 4°C overnight, then moved to 30% sucrose solution in 0.1M PB at 4°C until further processing. Slices were thoroughly washed with 0.1M PB and then placed in CUBIC #1 solution for two days for tissue clearing. Then, slices were washed again thoroughly with 0.1M PB and treated overnight with a 1:500 dilution of streptavidin conjugated to Alexa-633 in 0.1M PB. After another round of washing with 0.1M PB, slices were treated with CUBIC #2 solution for ~30min and then mounted in CUBIC #2 solution for confocal imaging (Zeiss) using a 20X objective or a 63X oil-immersion objective. The morphology of recorded cells was reconstructed using NeuTube⁵⁷.

Statistical analysis. All data presented were obtained from experimental replicates with at least three independent experimental repeats for each assay. All attempts of replication were successful. Data collection were not performed blind to the subject conditions, except for the behavior experiments and scoring. Data analysis were performed blinded to the scorer or did not

require manual scoring. Visual cues for the behaviour experiments were randomly assigned to the different experimental subjects so that the visual cue for one animal was the distractor of a different animal. All subjects underwent the same number of conditions (unless stated otherwise) in a randomly assigned fashion. All statistical analyses were performed with standard MATLAB functions. No specific analysis was used to estimate minimal population sample but the number of animals, trials, and recorded cells were larger or similar to those employed in previous works (refs^{16,17,19,26,48,49}). Unless otherwise noted, for all tests, non-parametric two-tailed Wilcoxon's paired signed-rank test, Kruskal-Wallis one-way analysis of variance and Friedman were used. When parametric tests were used, the data satisfied the criteria for normality (Kolmogorov–Smirnov test) and equality of variance (Bartlett's test for equal variance). For multiple comparisons, Tukey's honesty *post hoc* test was employed and the corrected *P < 0.05, **P < 0.01, ***P < 0.001 are indicated, two-sided. NS indicates not significant P-values for Pearson's correlations or Spearman's correlations are computed using a Student's t distribution for a transformation of the correlation. Results are displayed as mean ± IC95 unless indicated otherwise. Boxplots represent median and 25th/75th percentiles and their whiskers the data range. In some of the plots, outlier values are not shown for clarity of presentation, but all data points and animal were always included in the statistical analysis. The exact number of replications for each experiment is detailed in the text and figures.

Reporting Summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

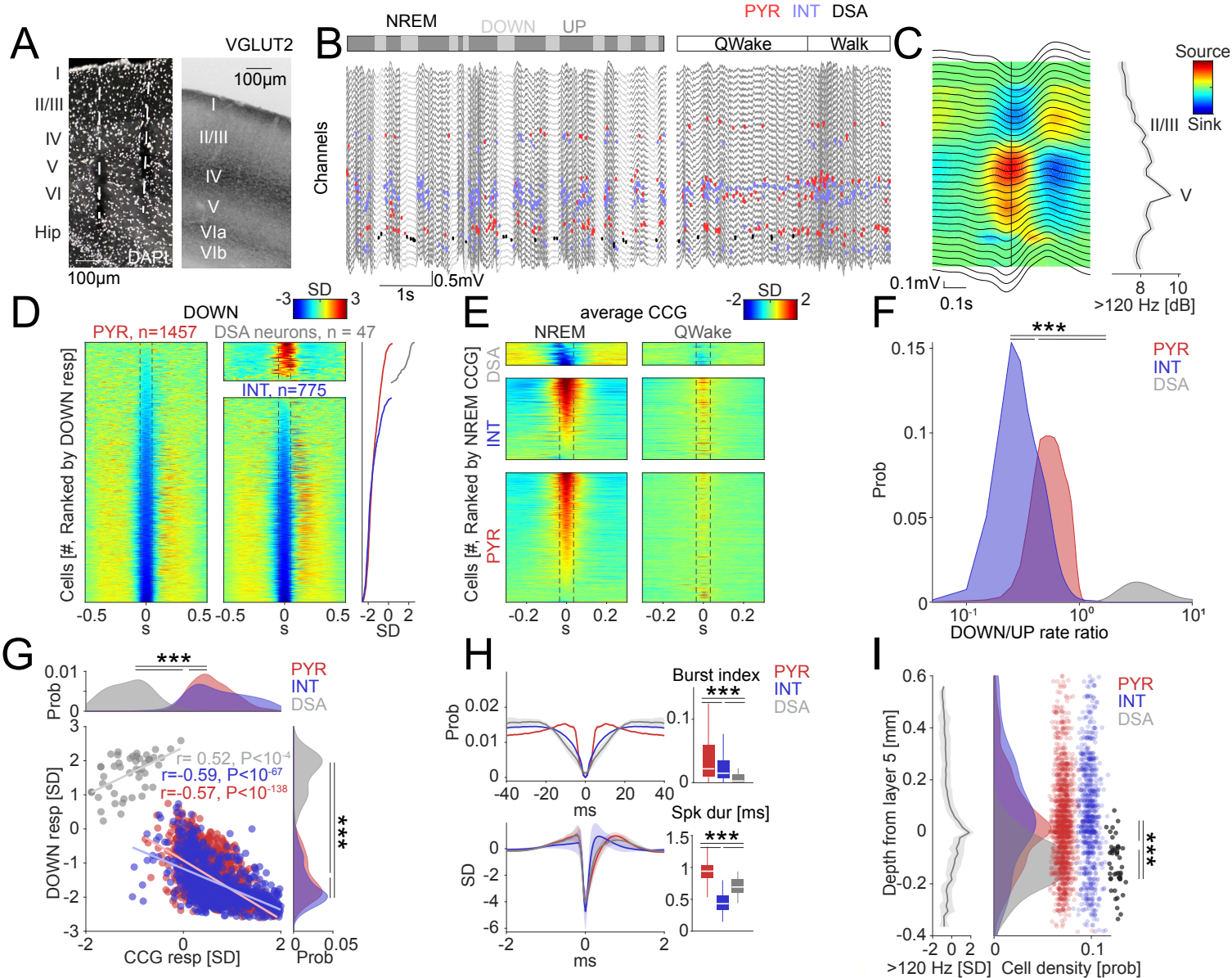
Code availability. All custom code is freely available on the Buzsáki Laboratory repository (<https://github.com/buzsakilab/buzcode>).

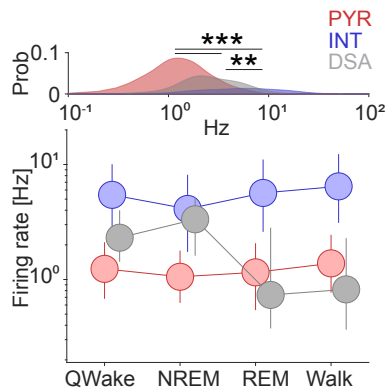
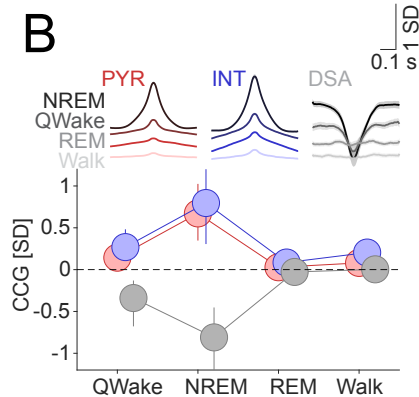
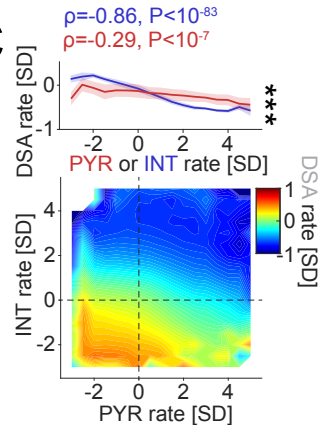
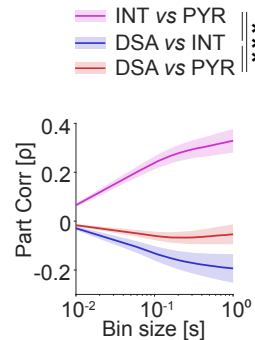
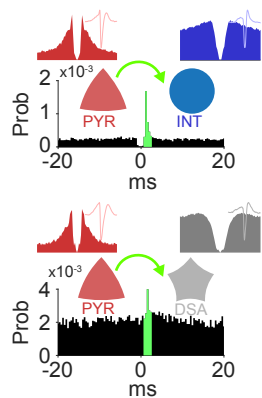
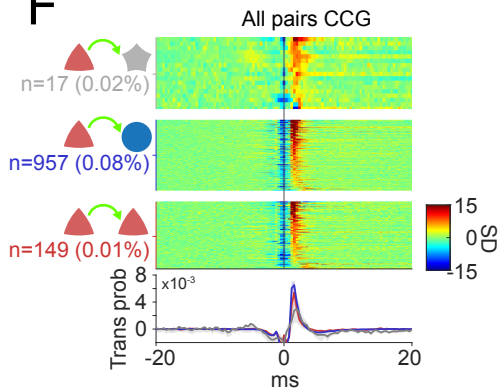
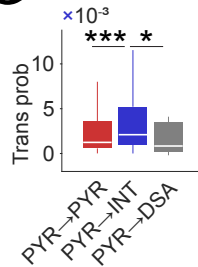
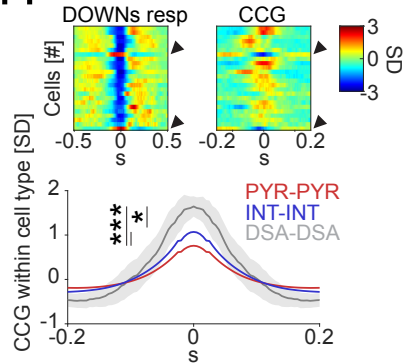
Data availability. The data that support the main findings of this study is publicly available on the Buzsaki Lab Databank, <https://buzsakilab.com/wp/public-data/>.

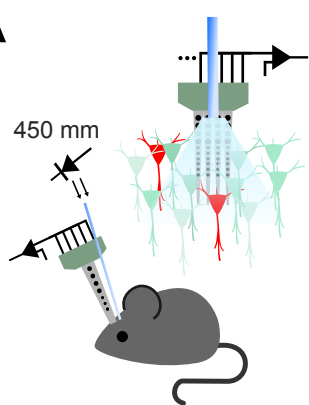
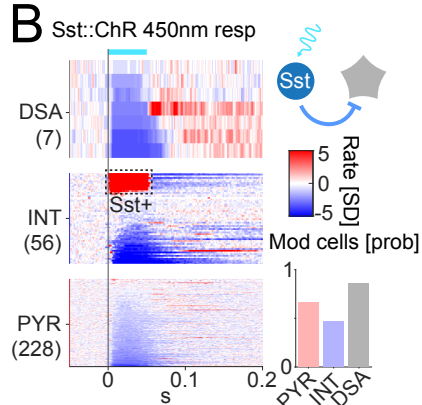
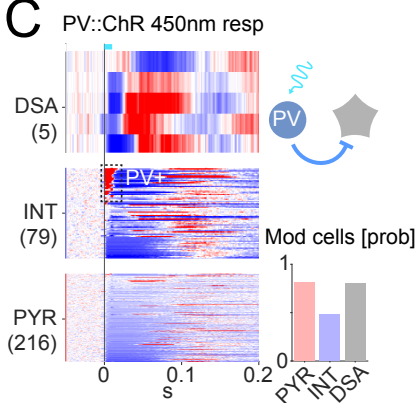
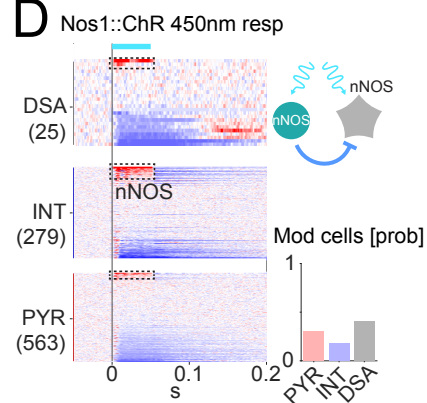
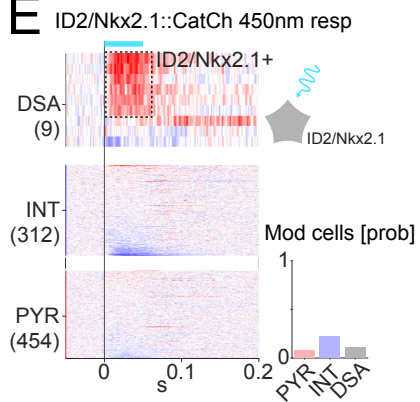
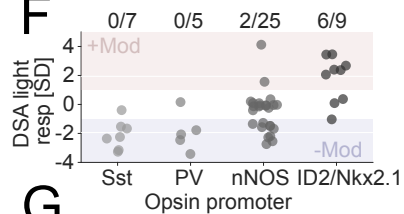
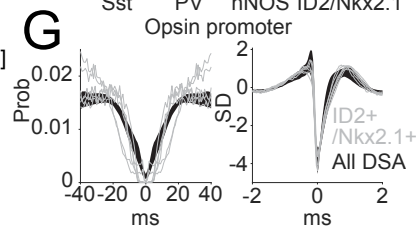
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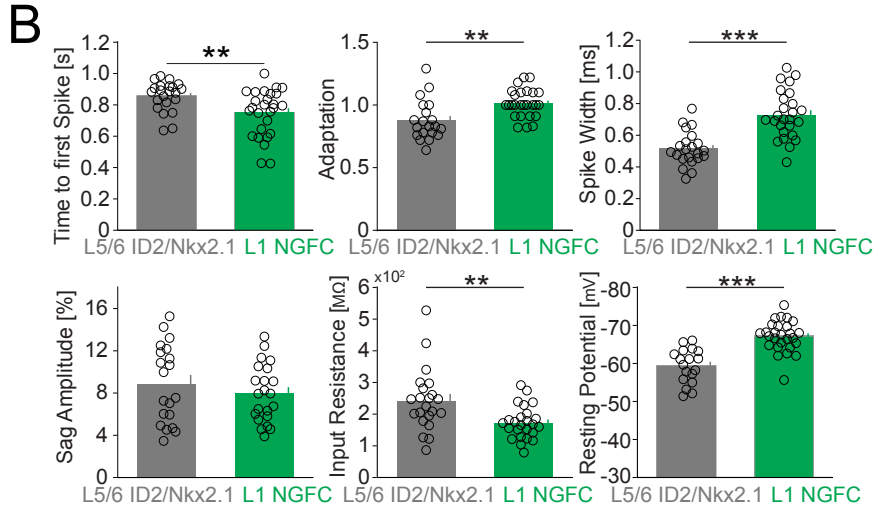
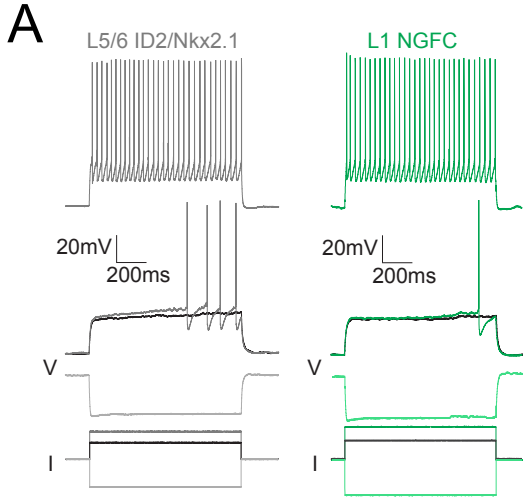
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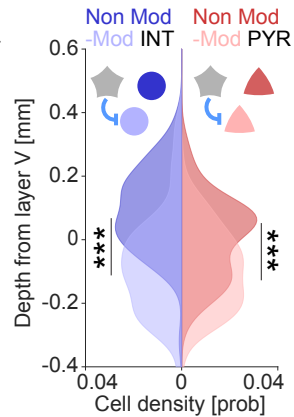
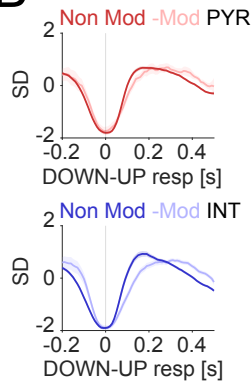
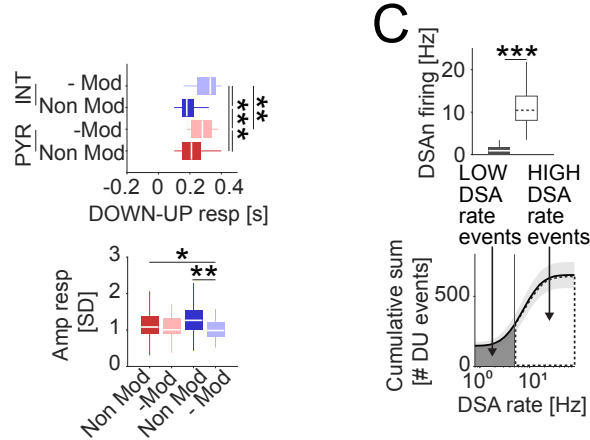
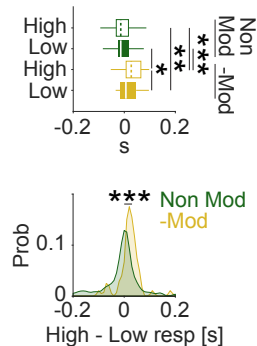
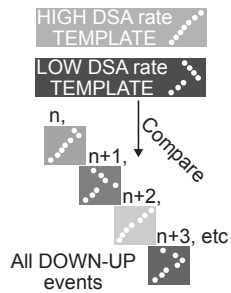
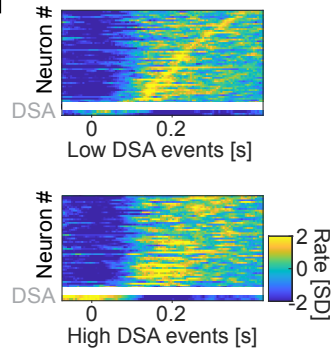
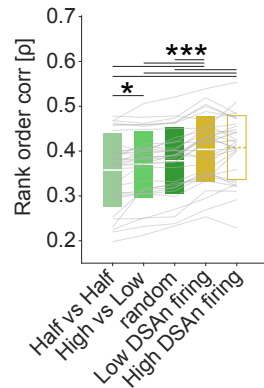
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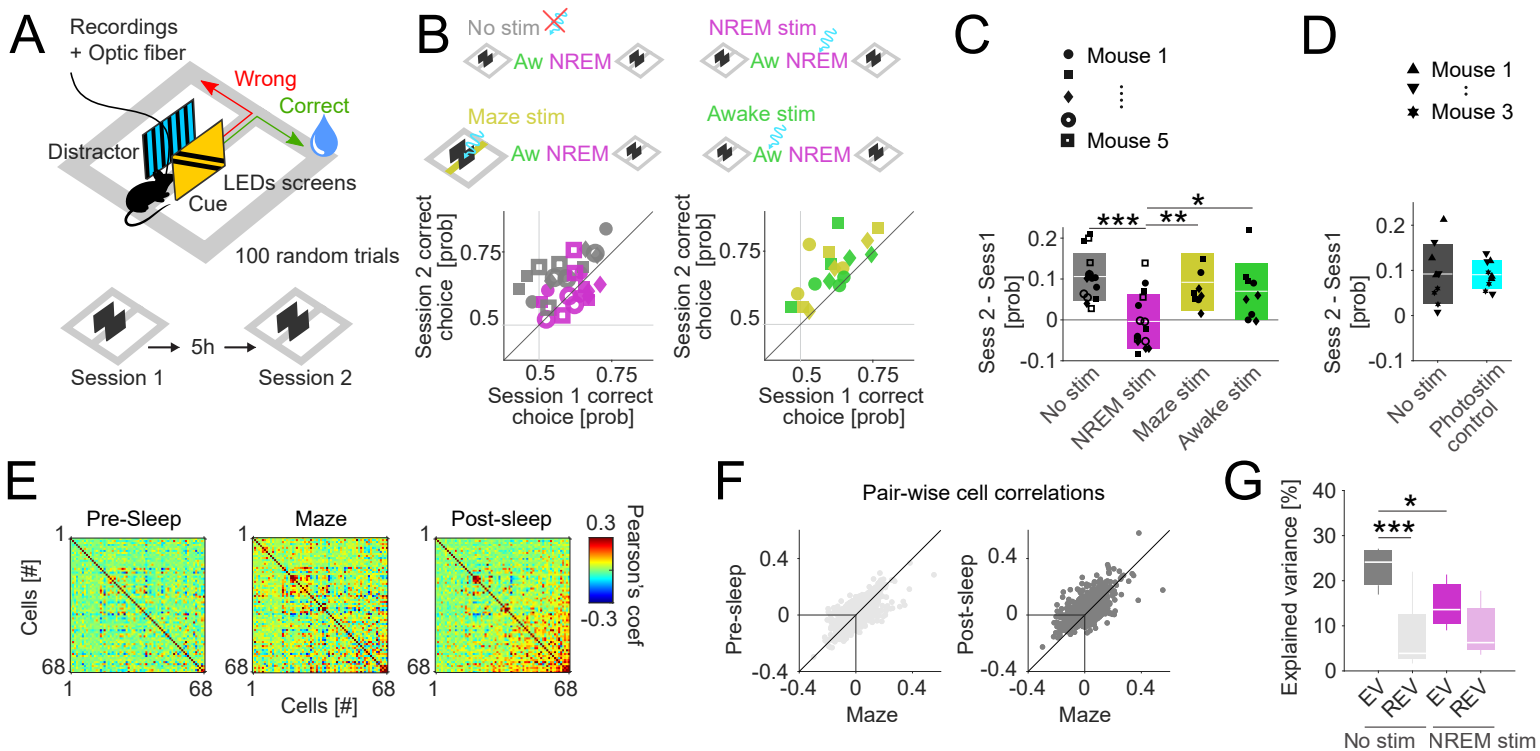


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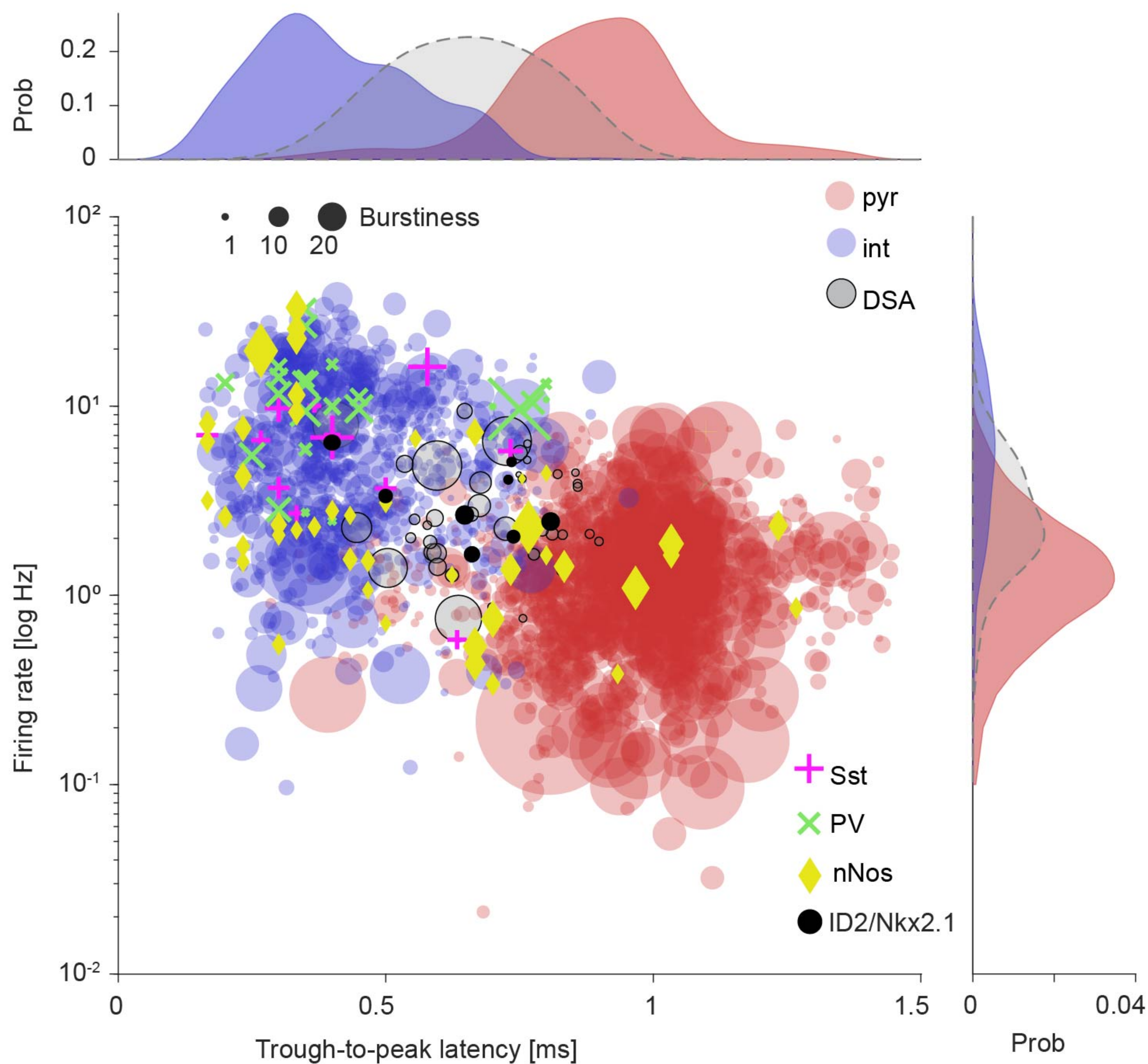
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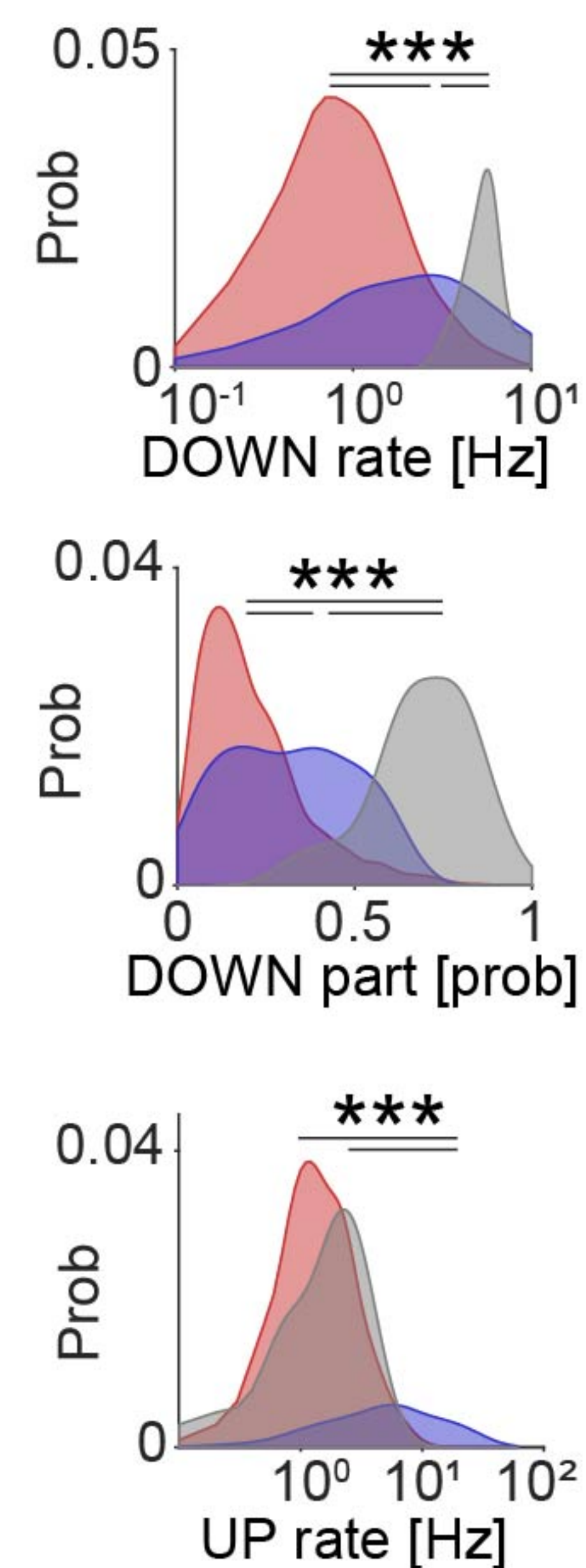
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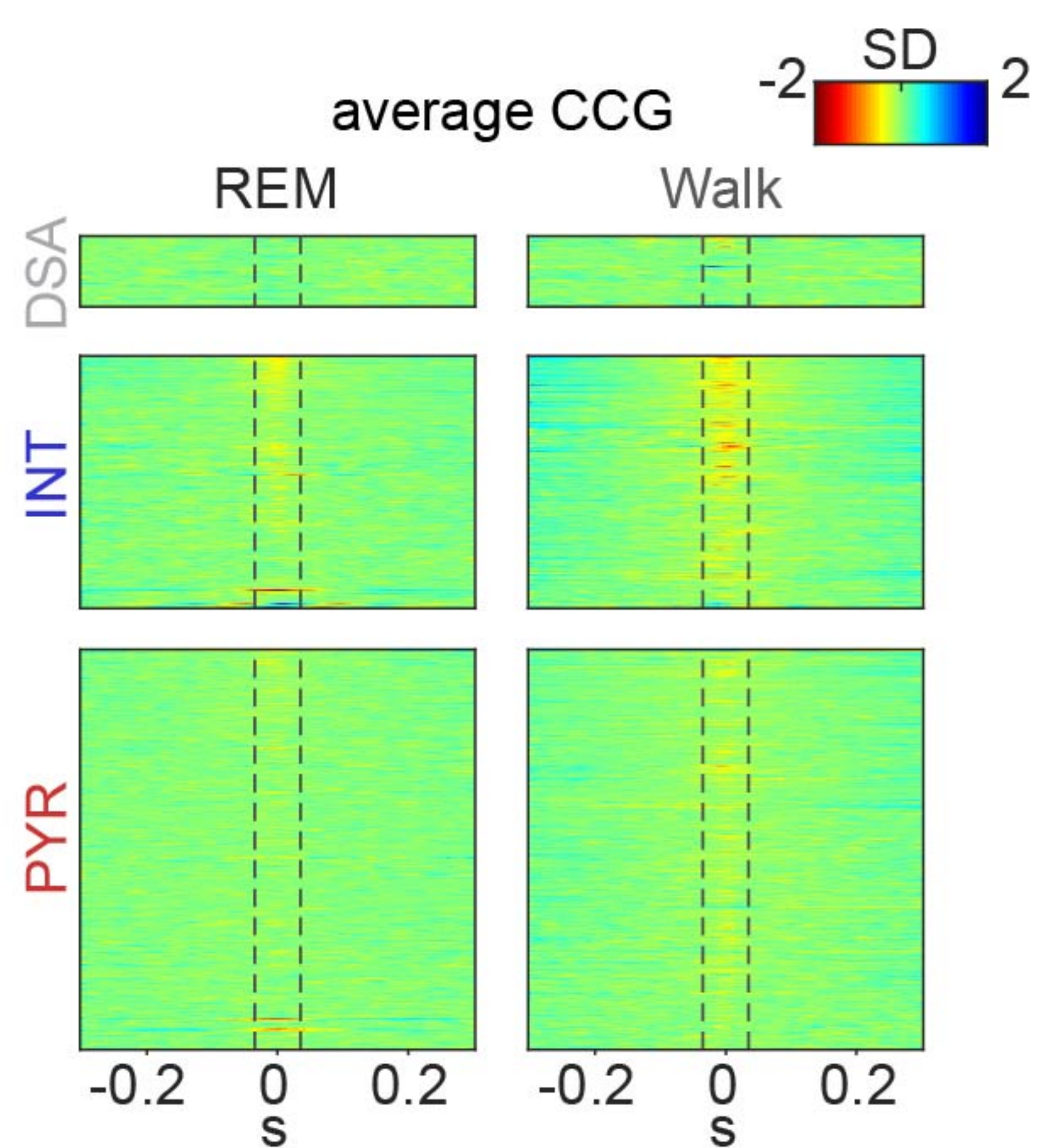
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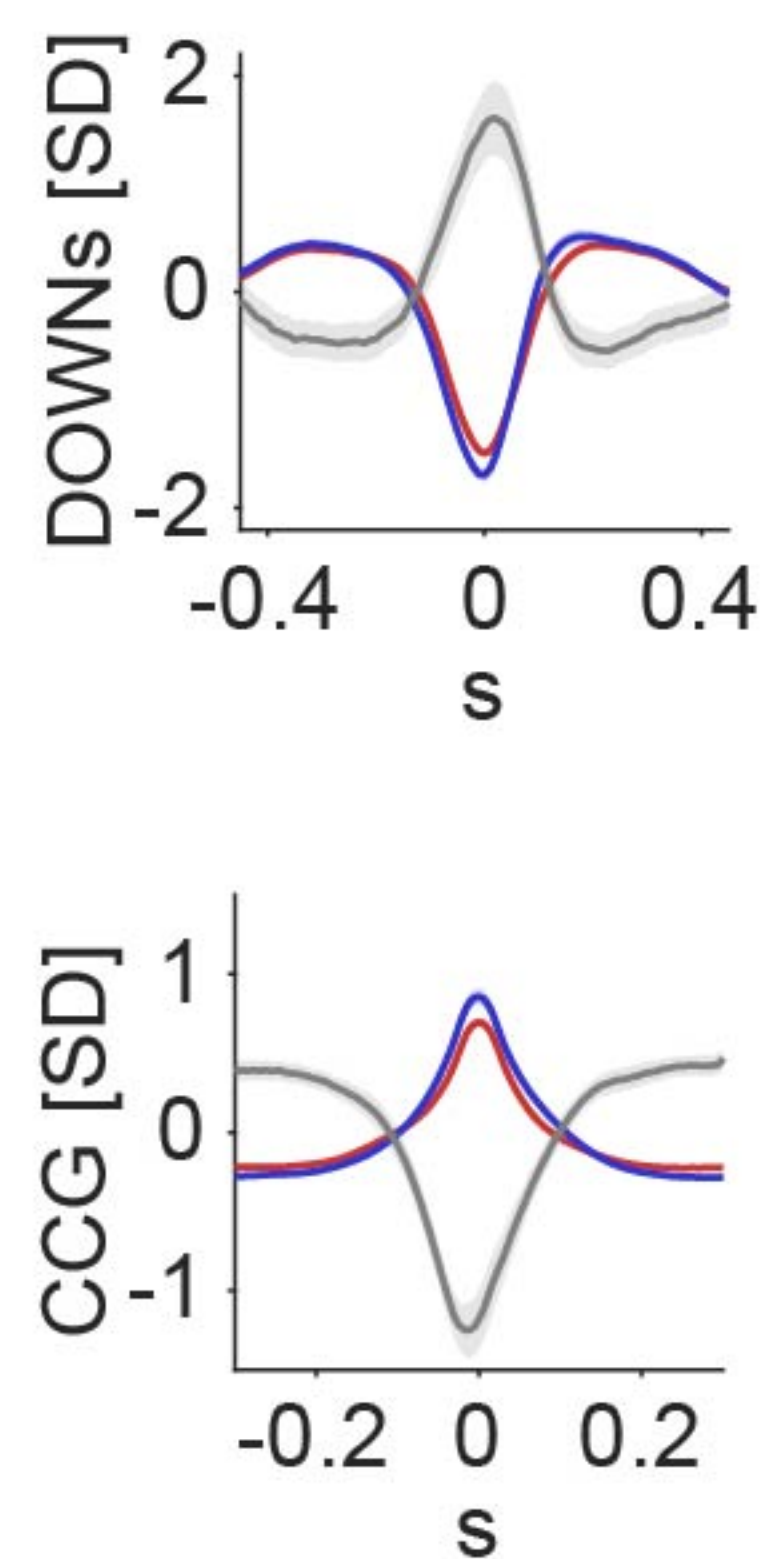
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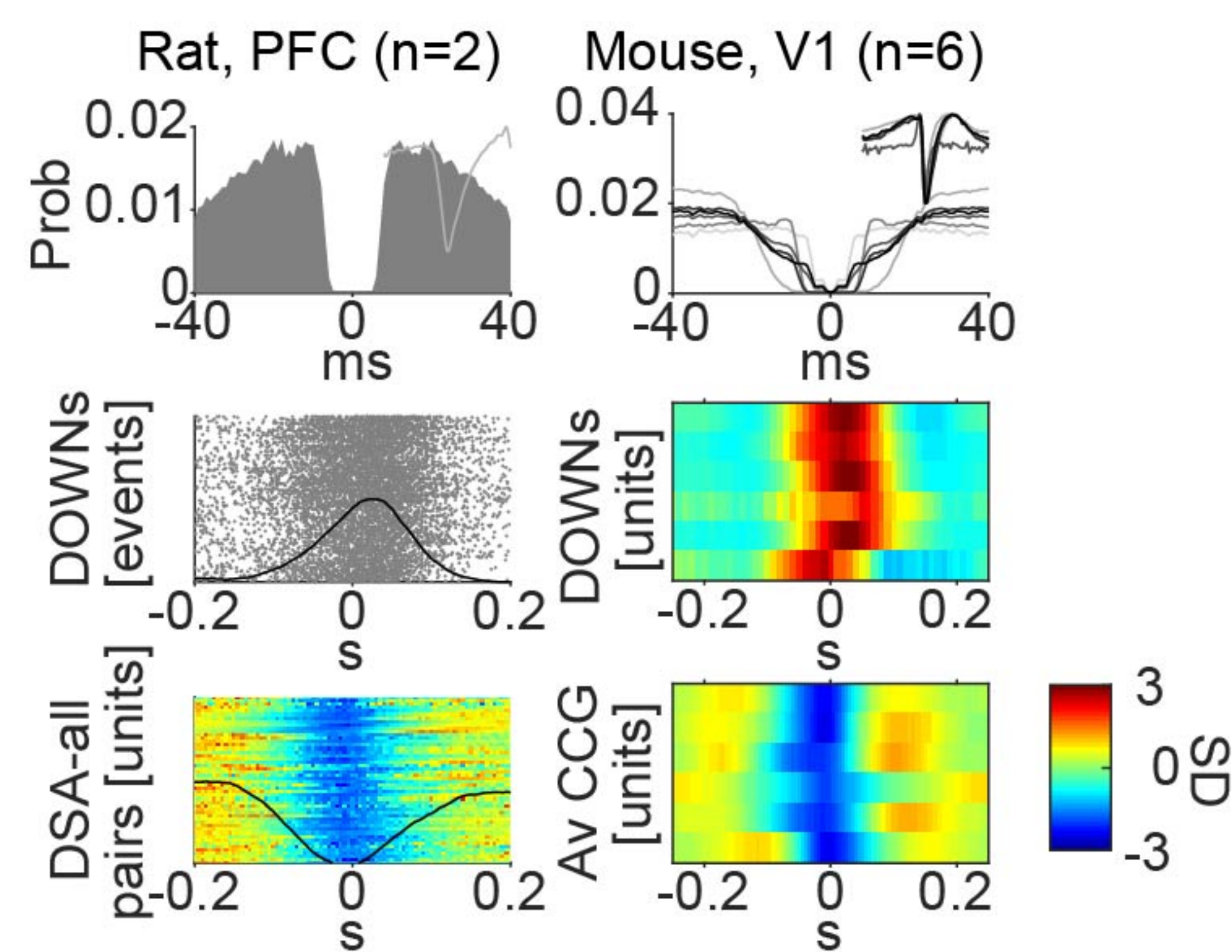
C



D

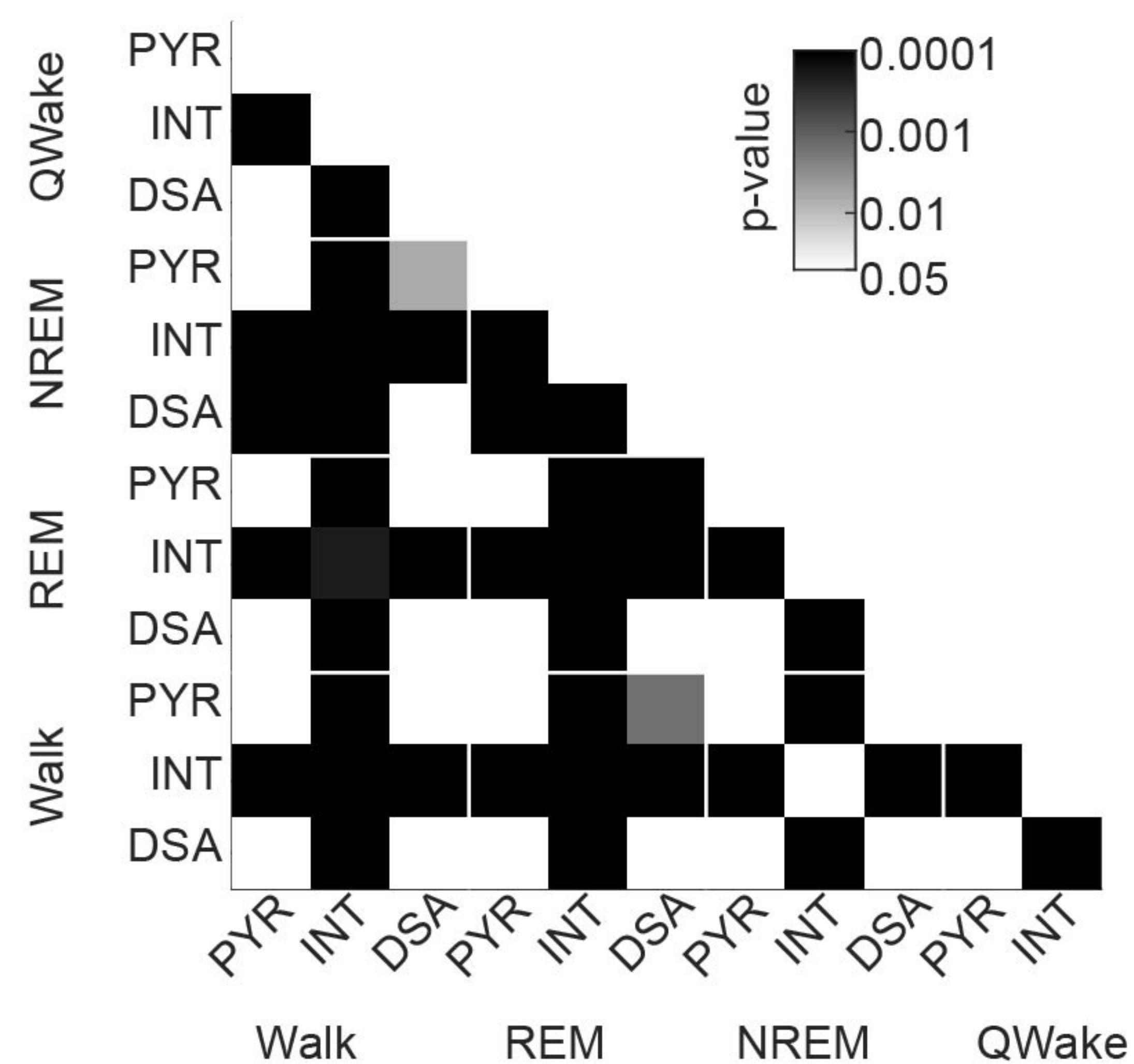


E

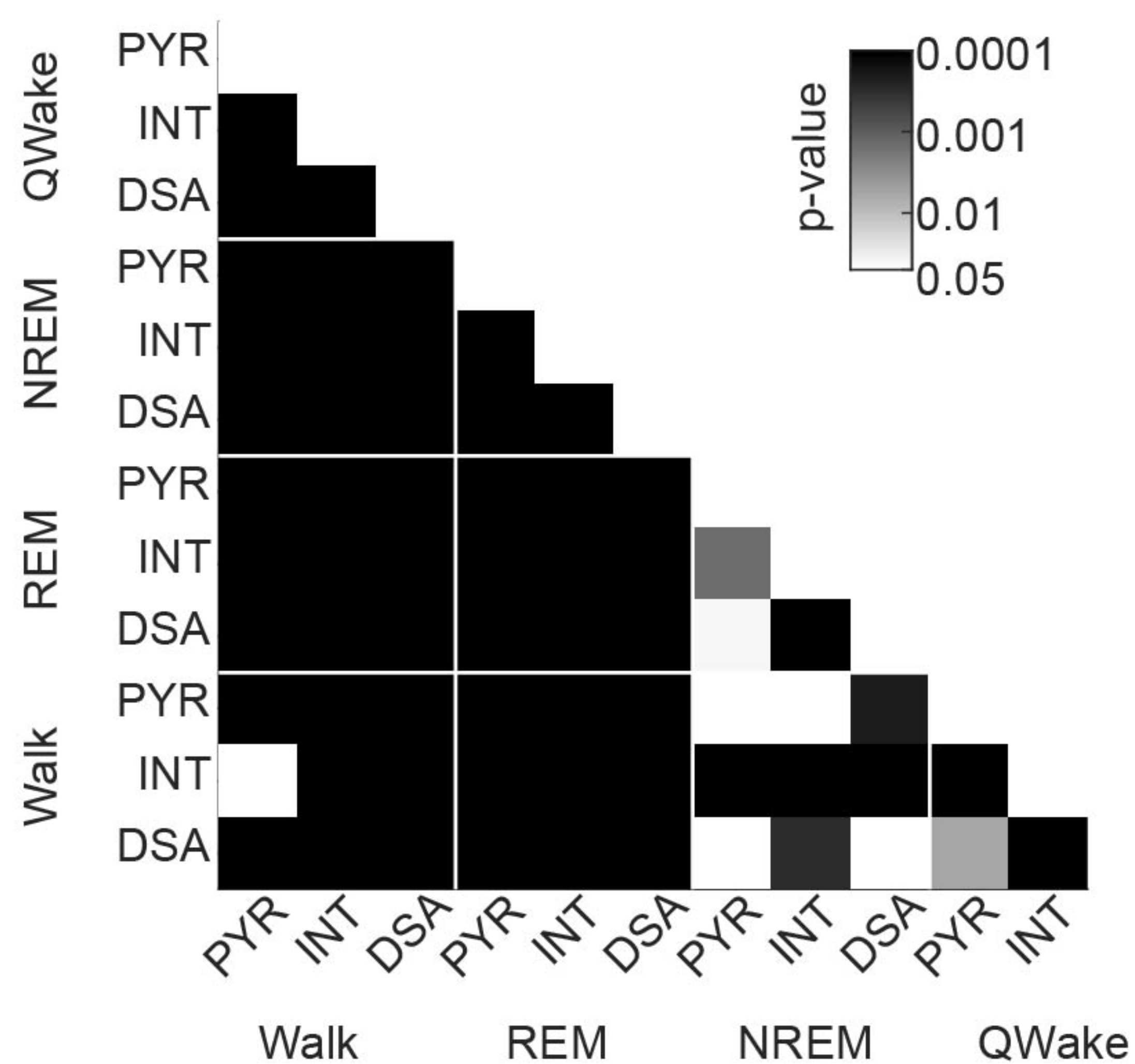
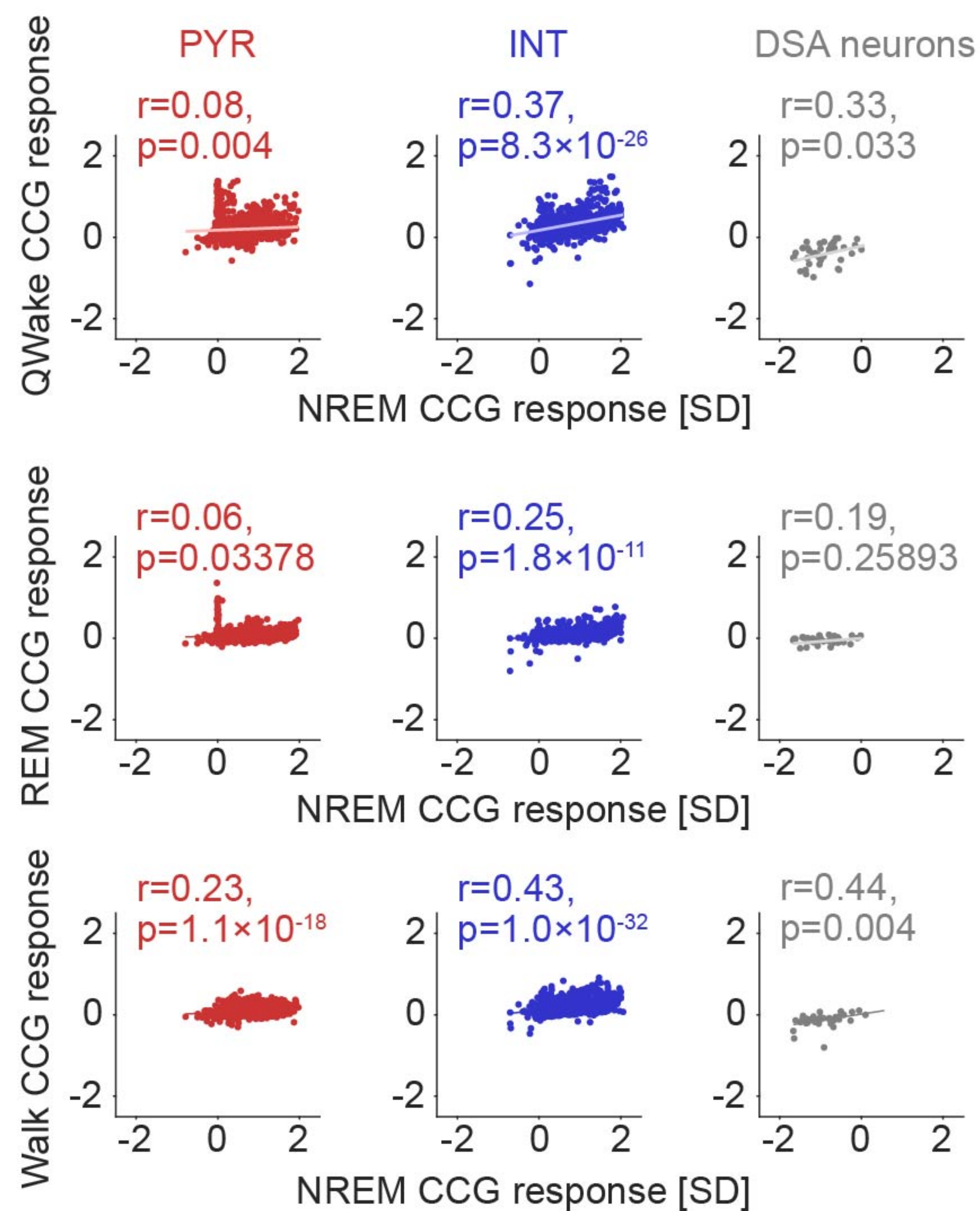
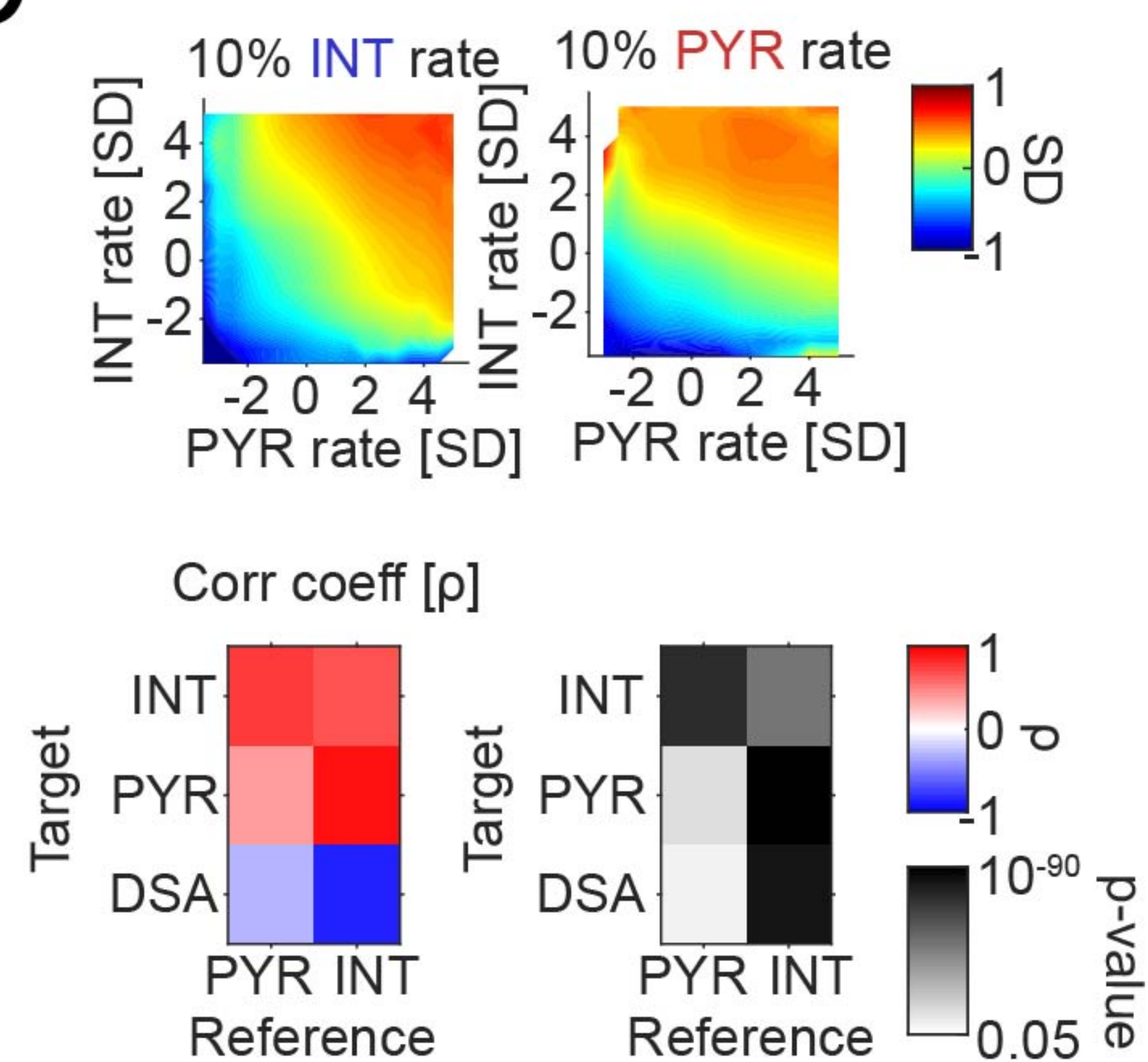


A

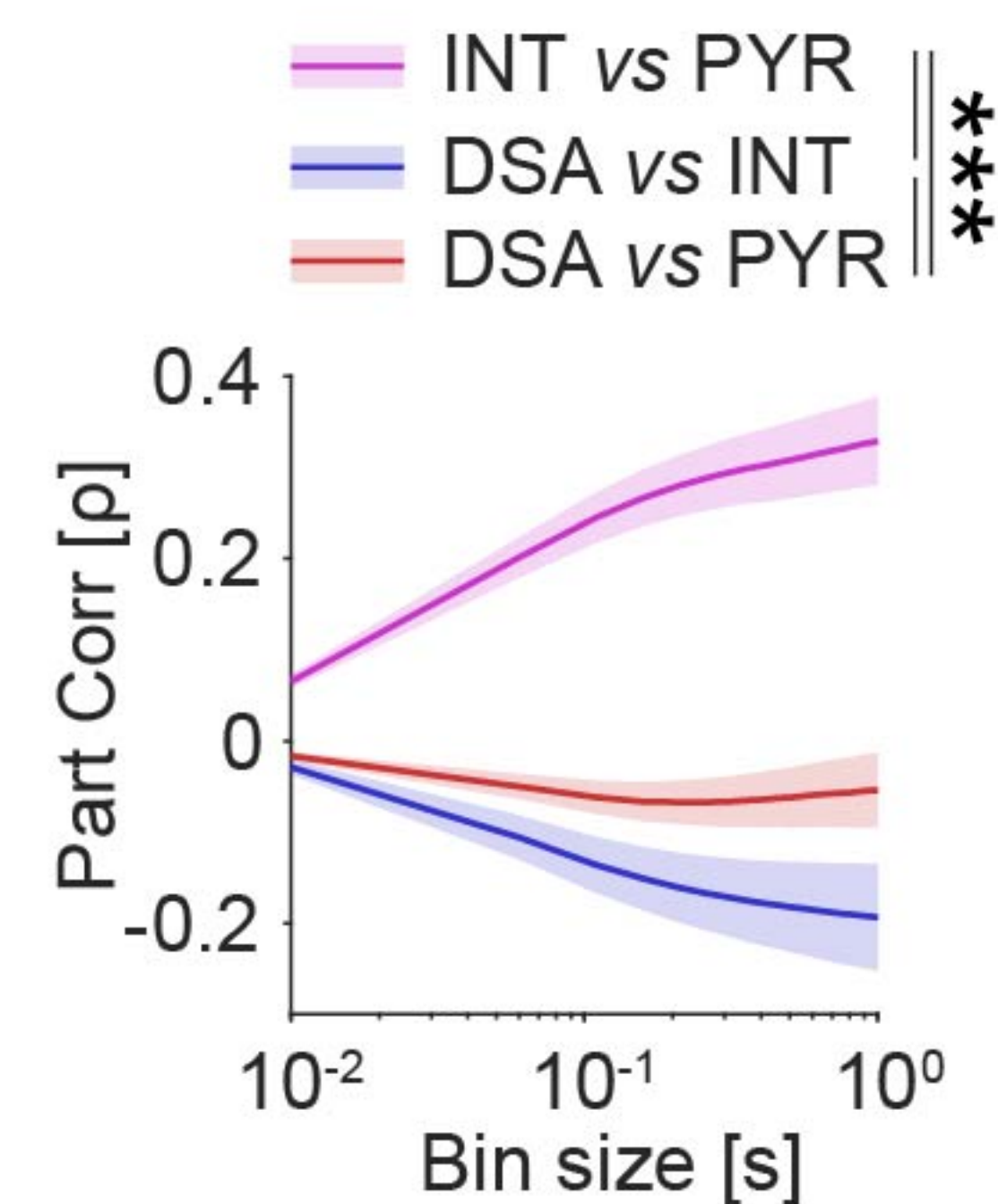
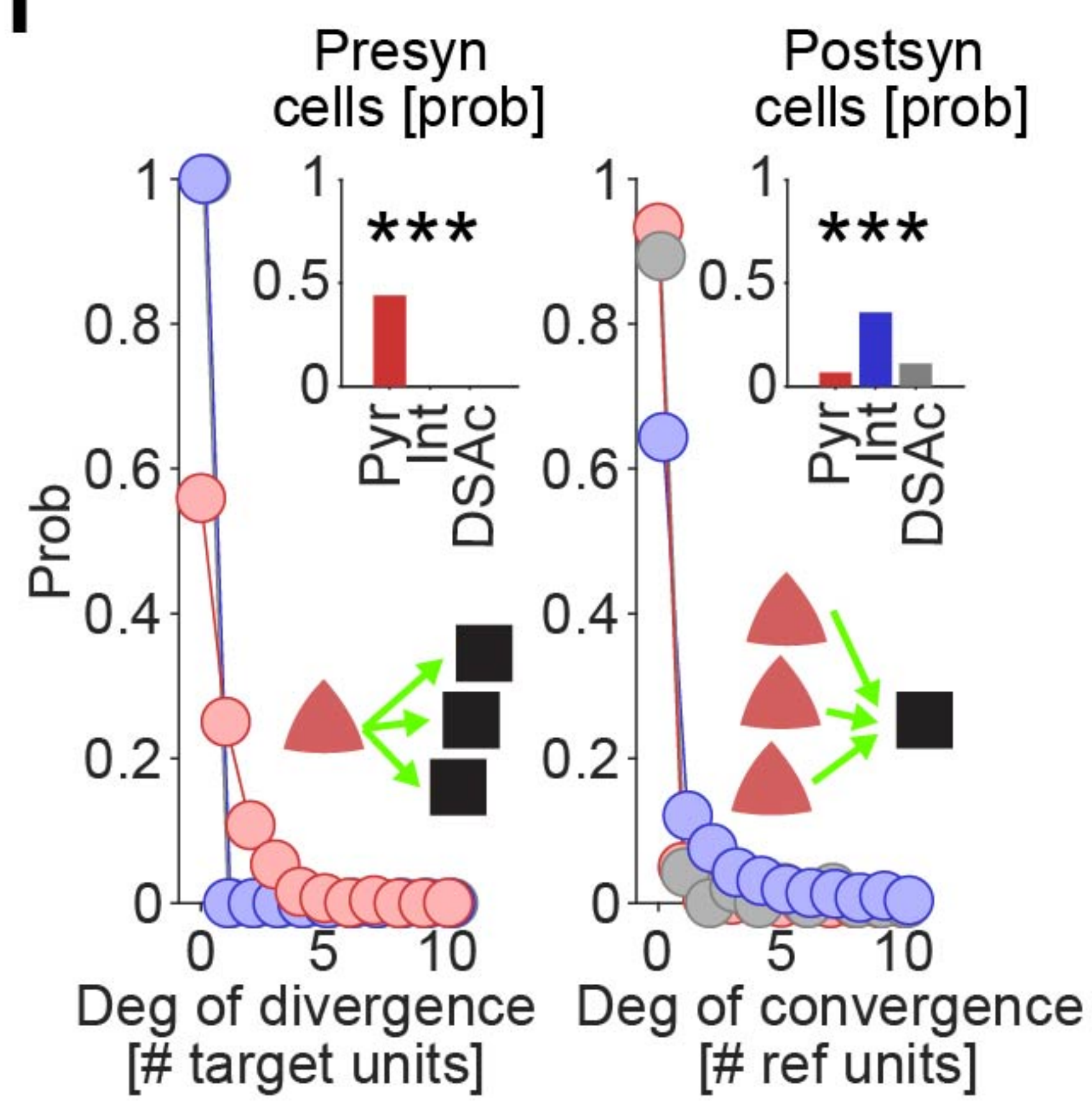
FIRING RATE

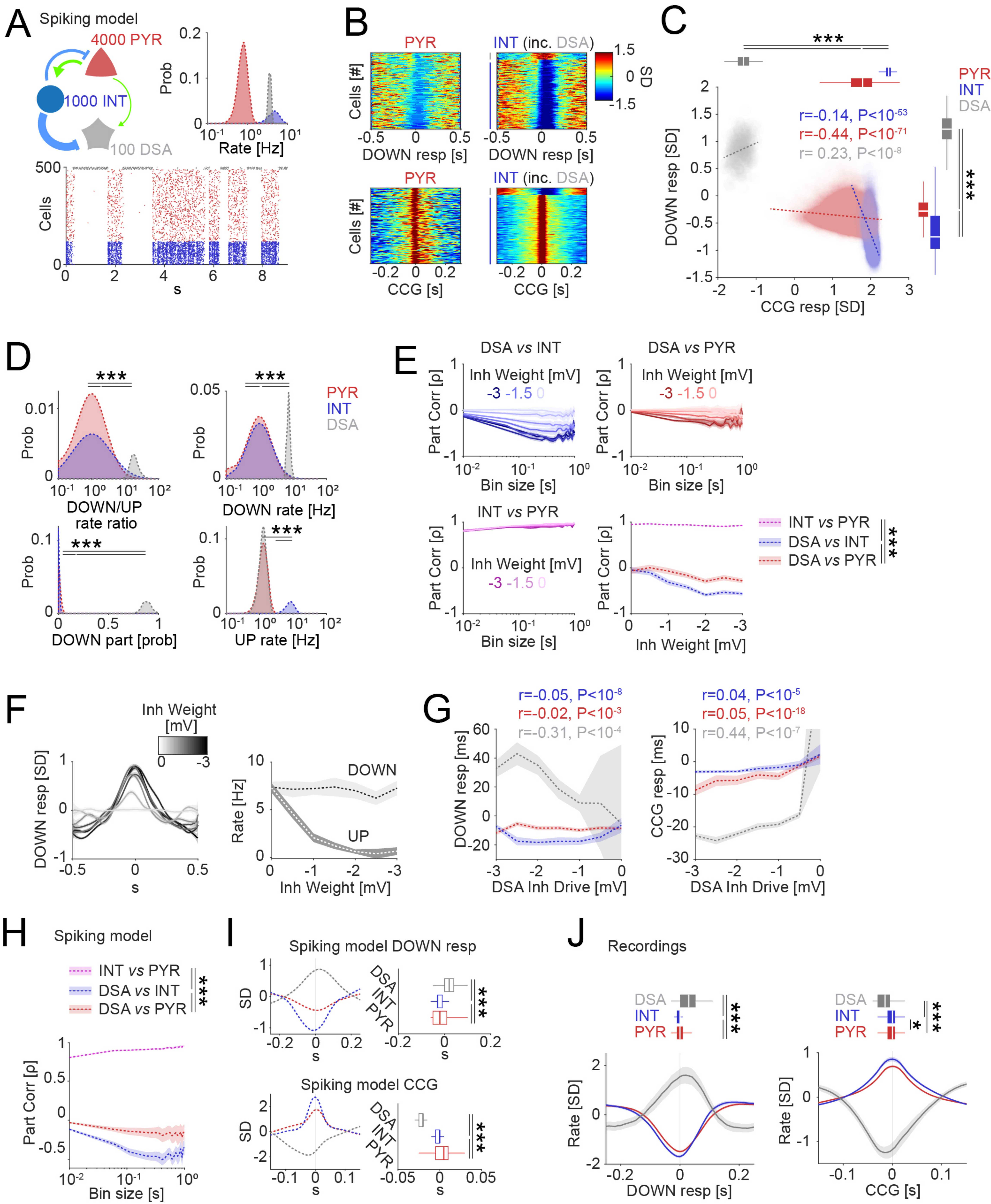
**B**

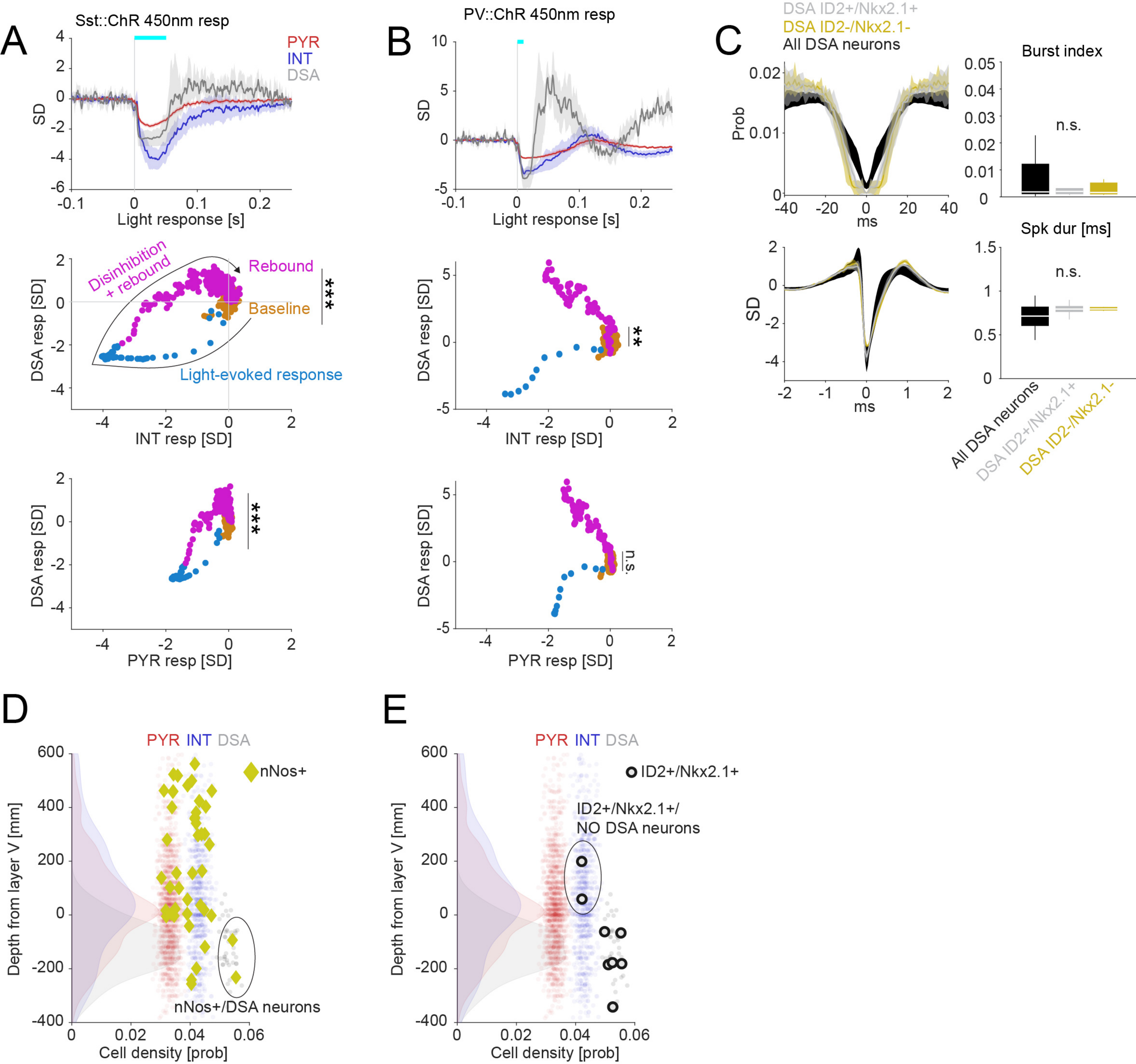
AVERAGE CCG

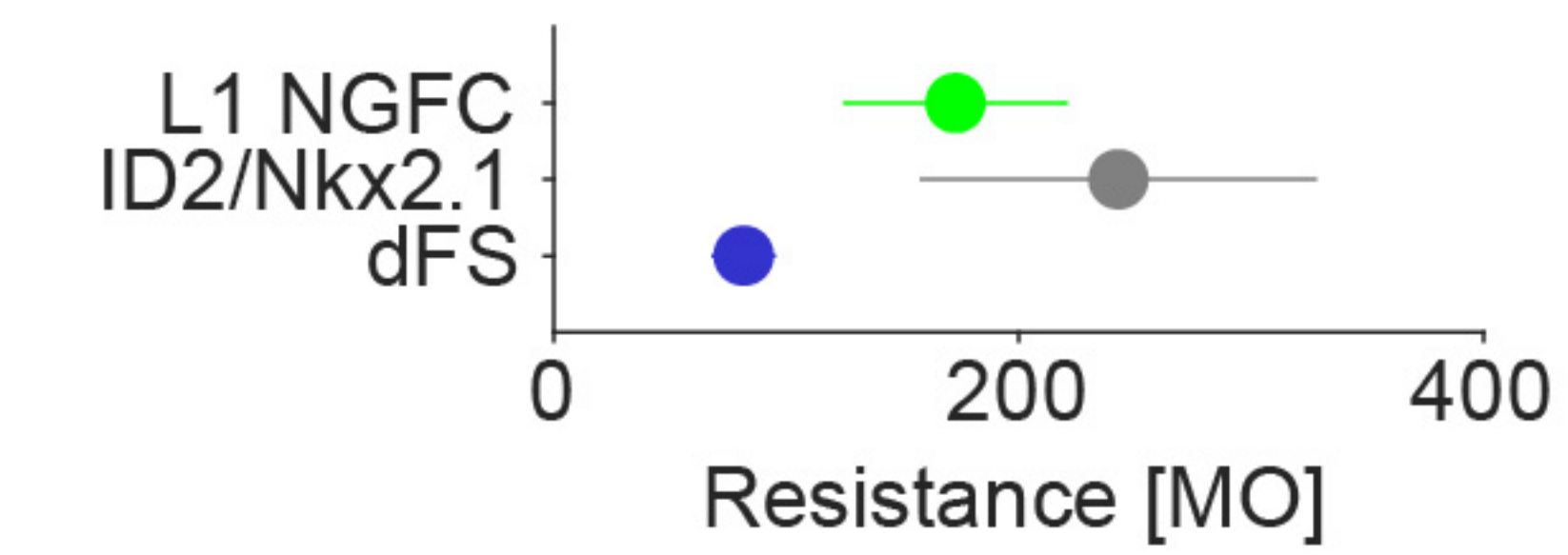
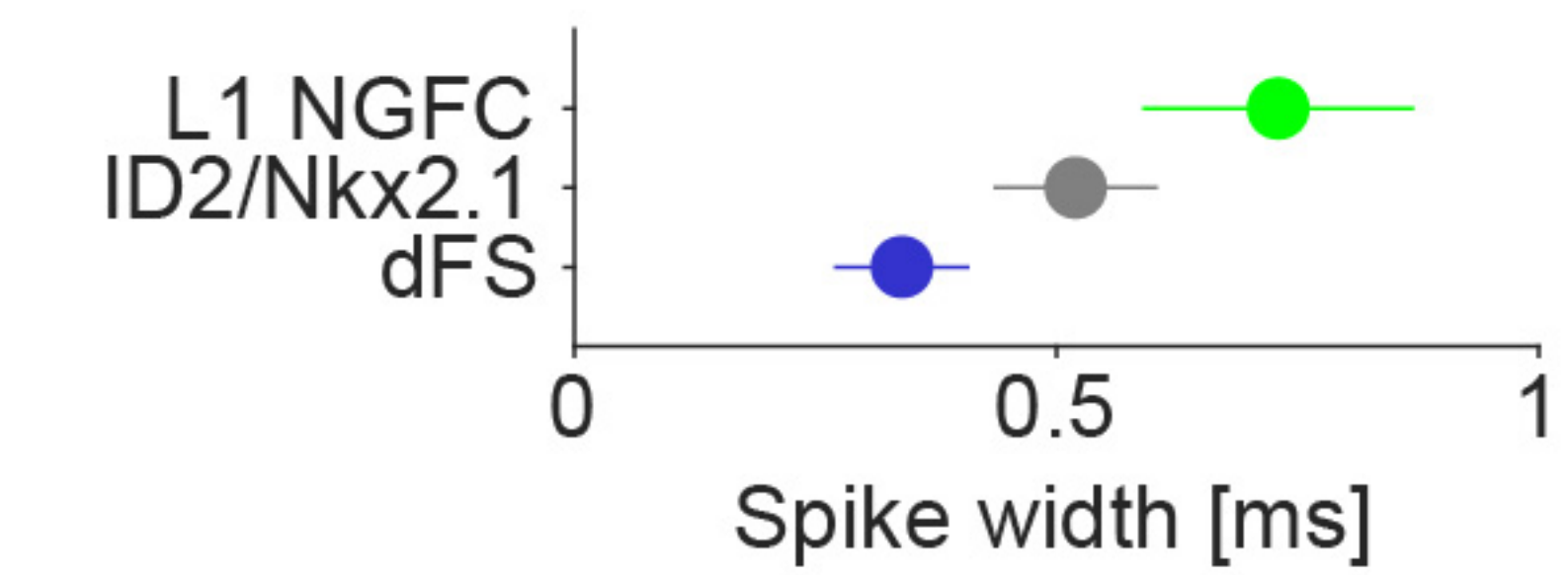
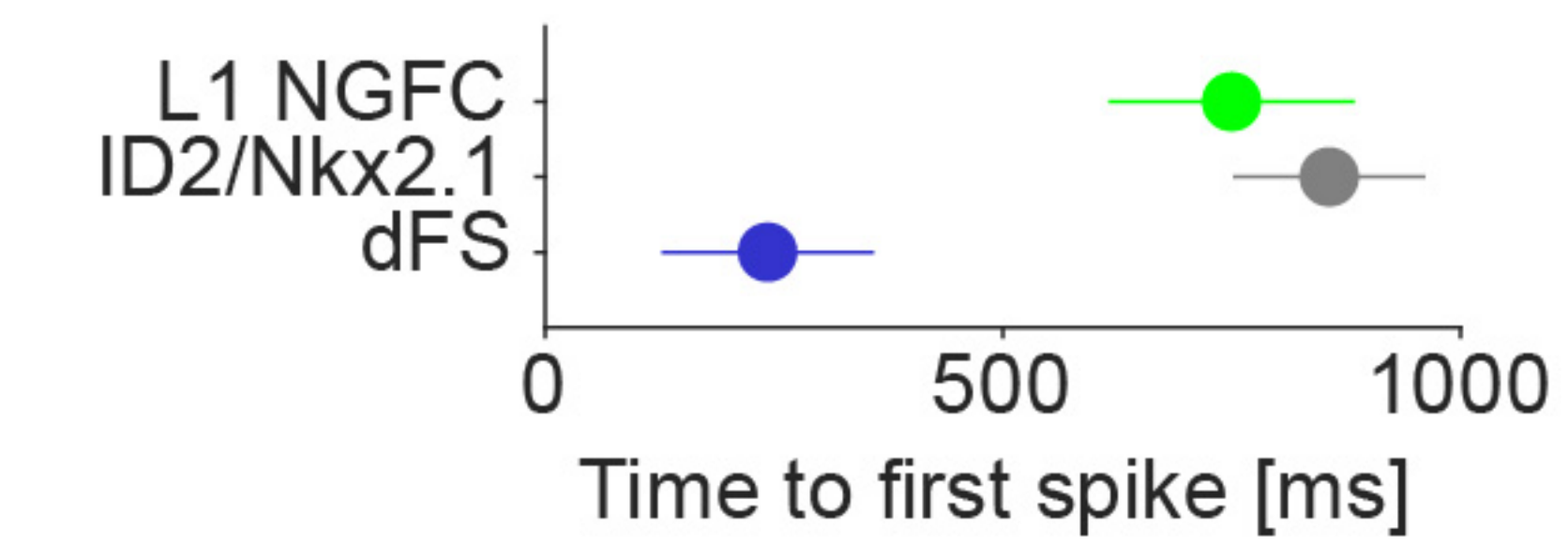
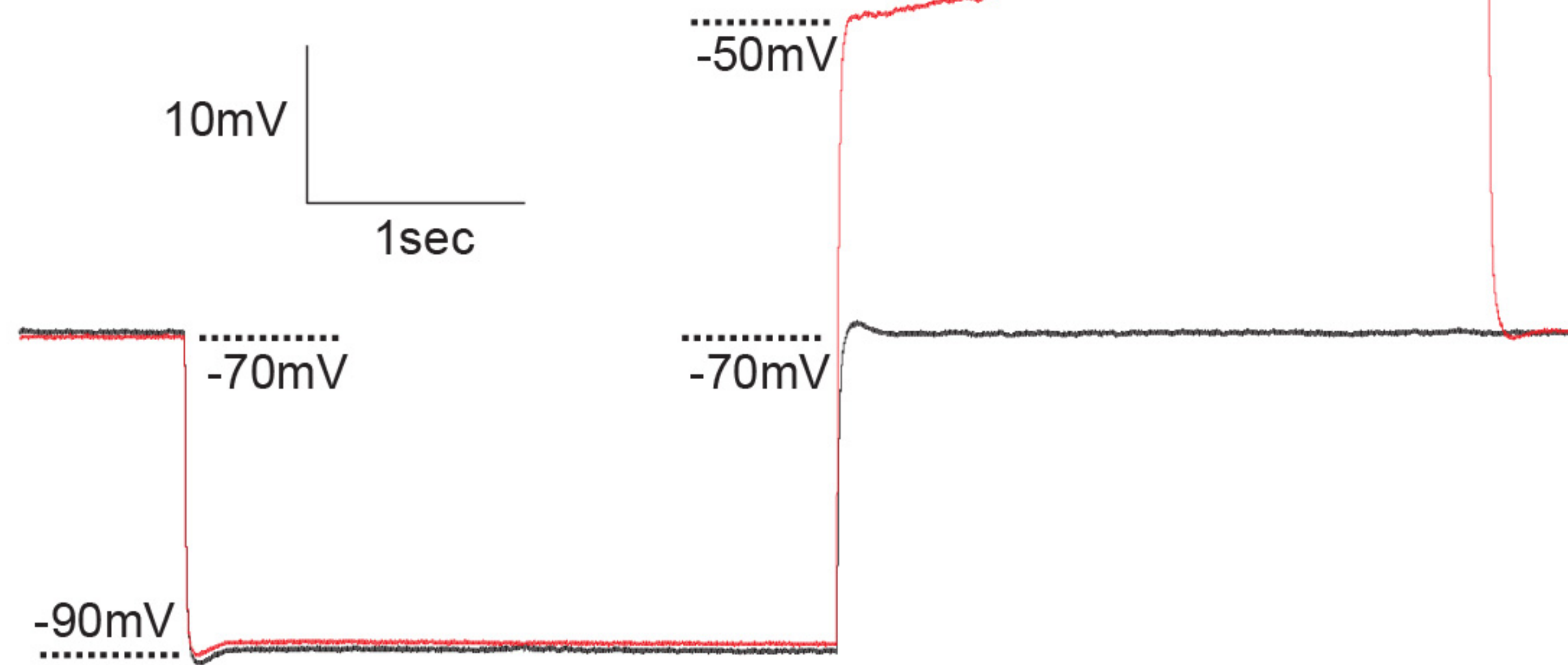
**C****D****E**

Same number of spikes across groups

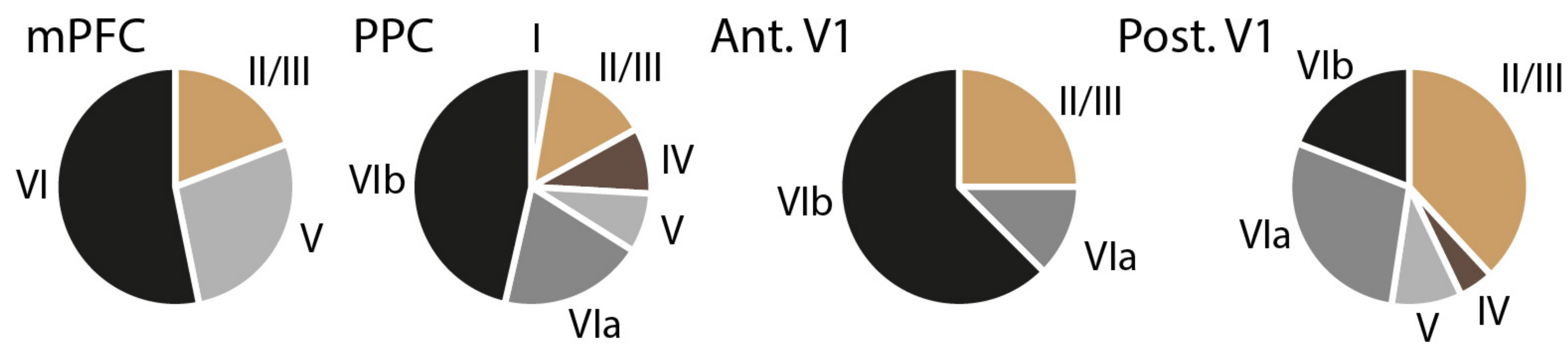
**F**



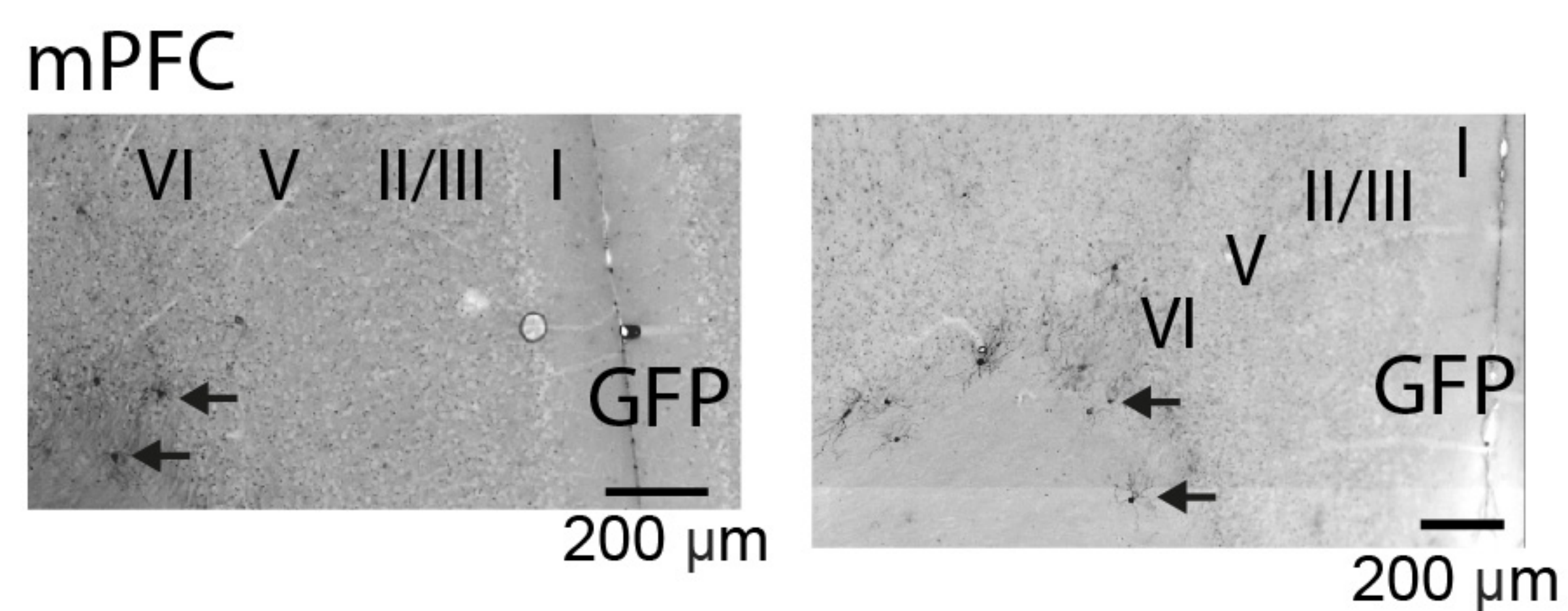


A**B**

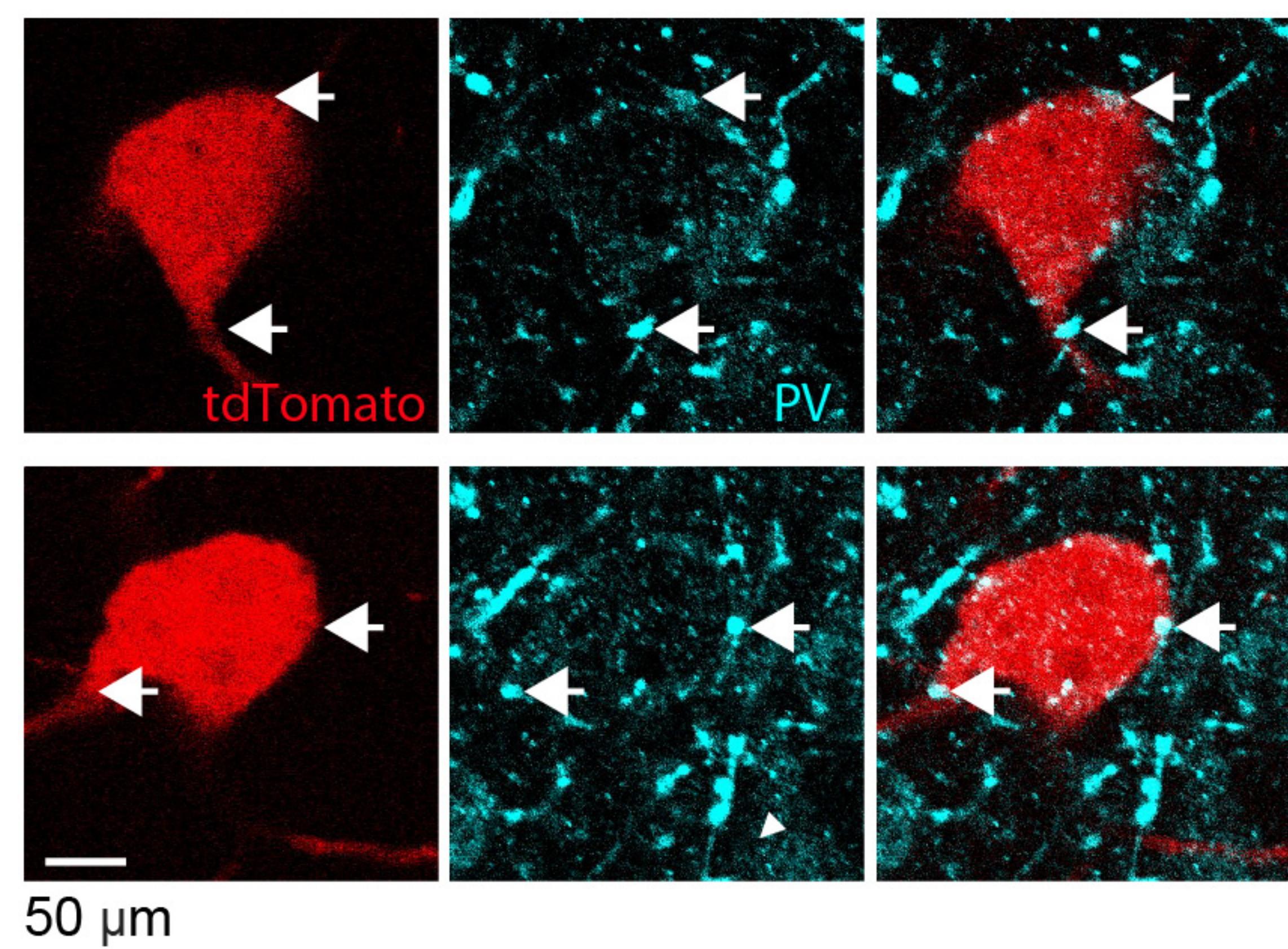
A ID2+/Nkx2.1+ layer distribution (%)



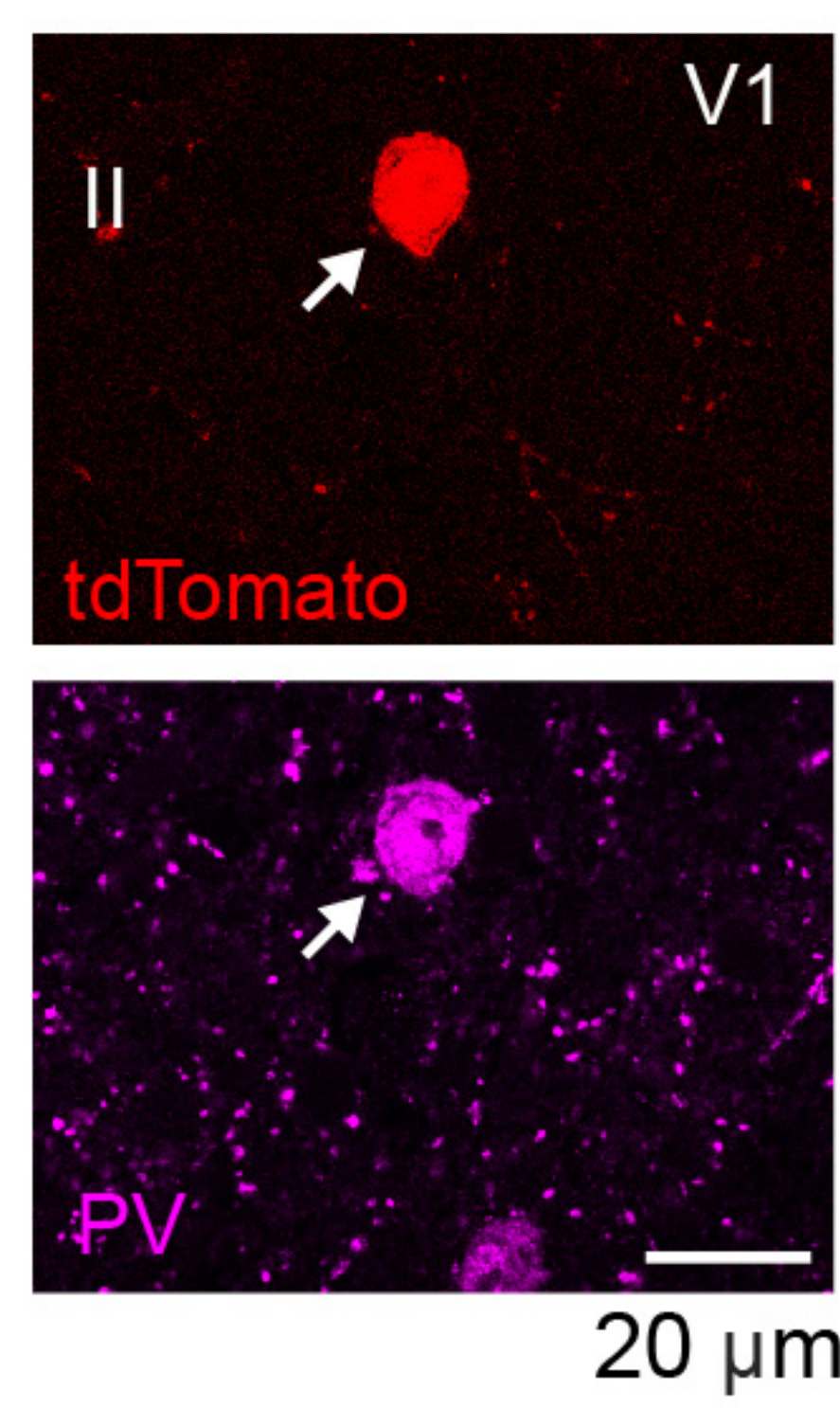
B



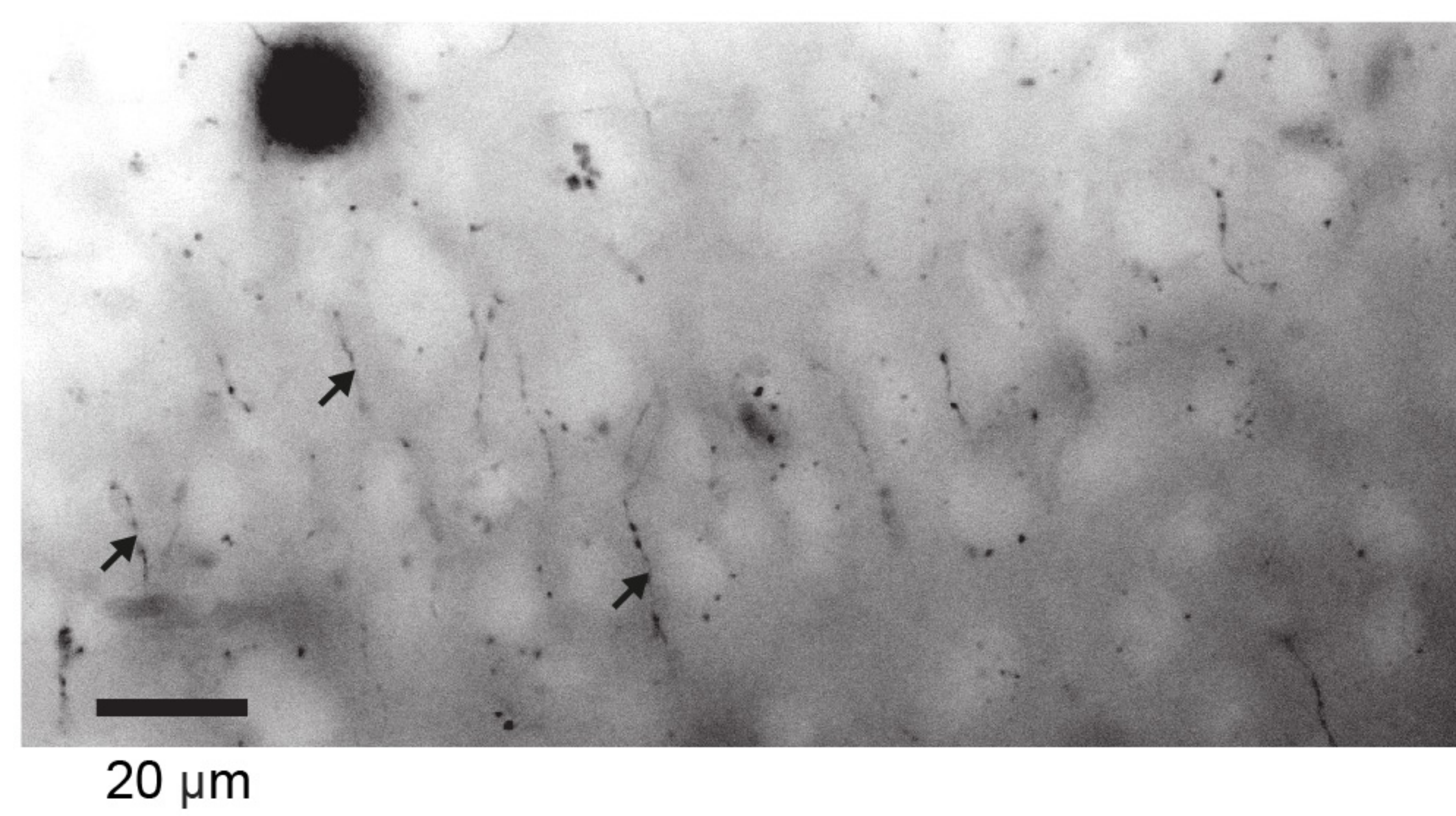
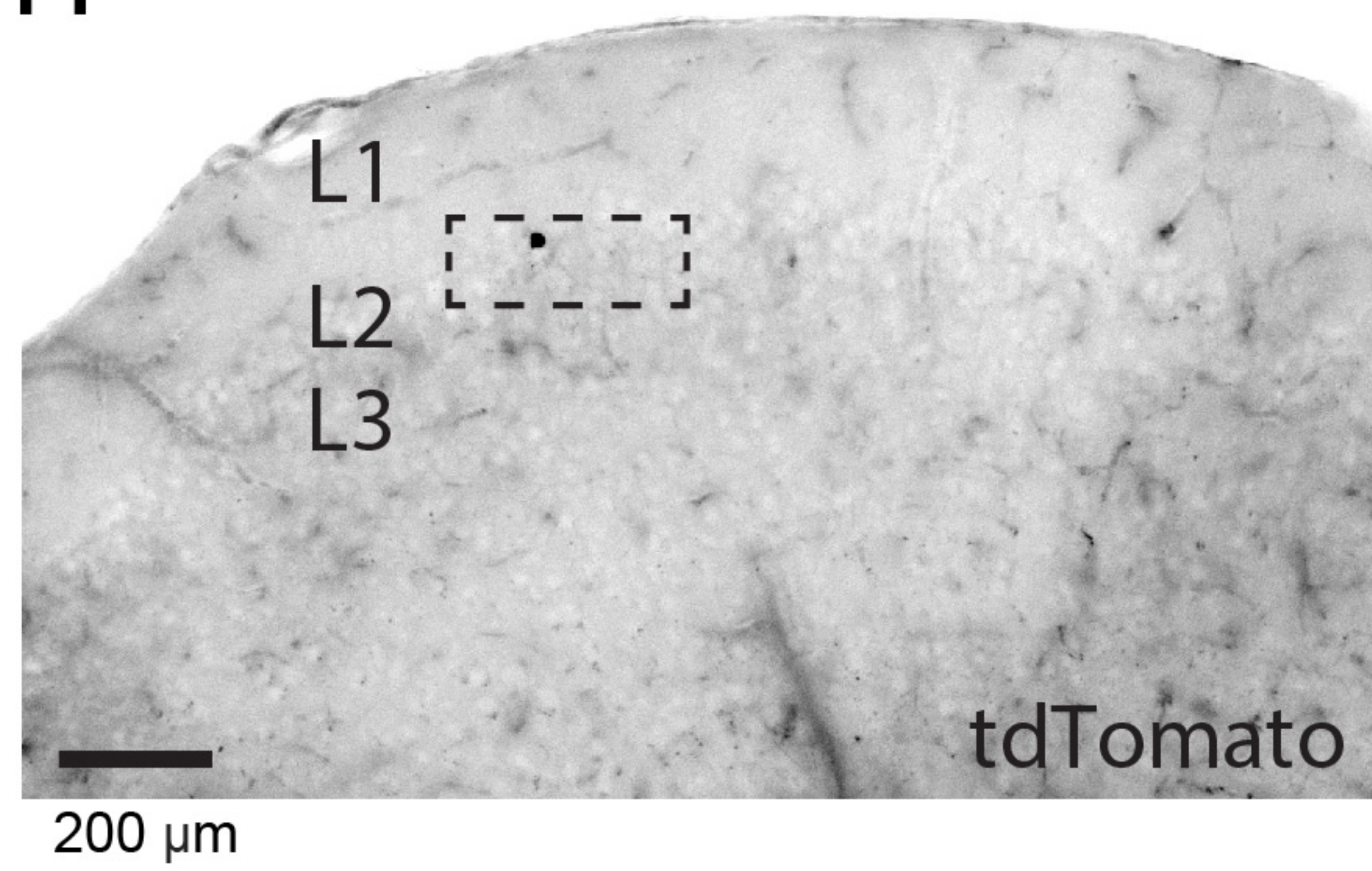
C



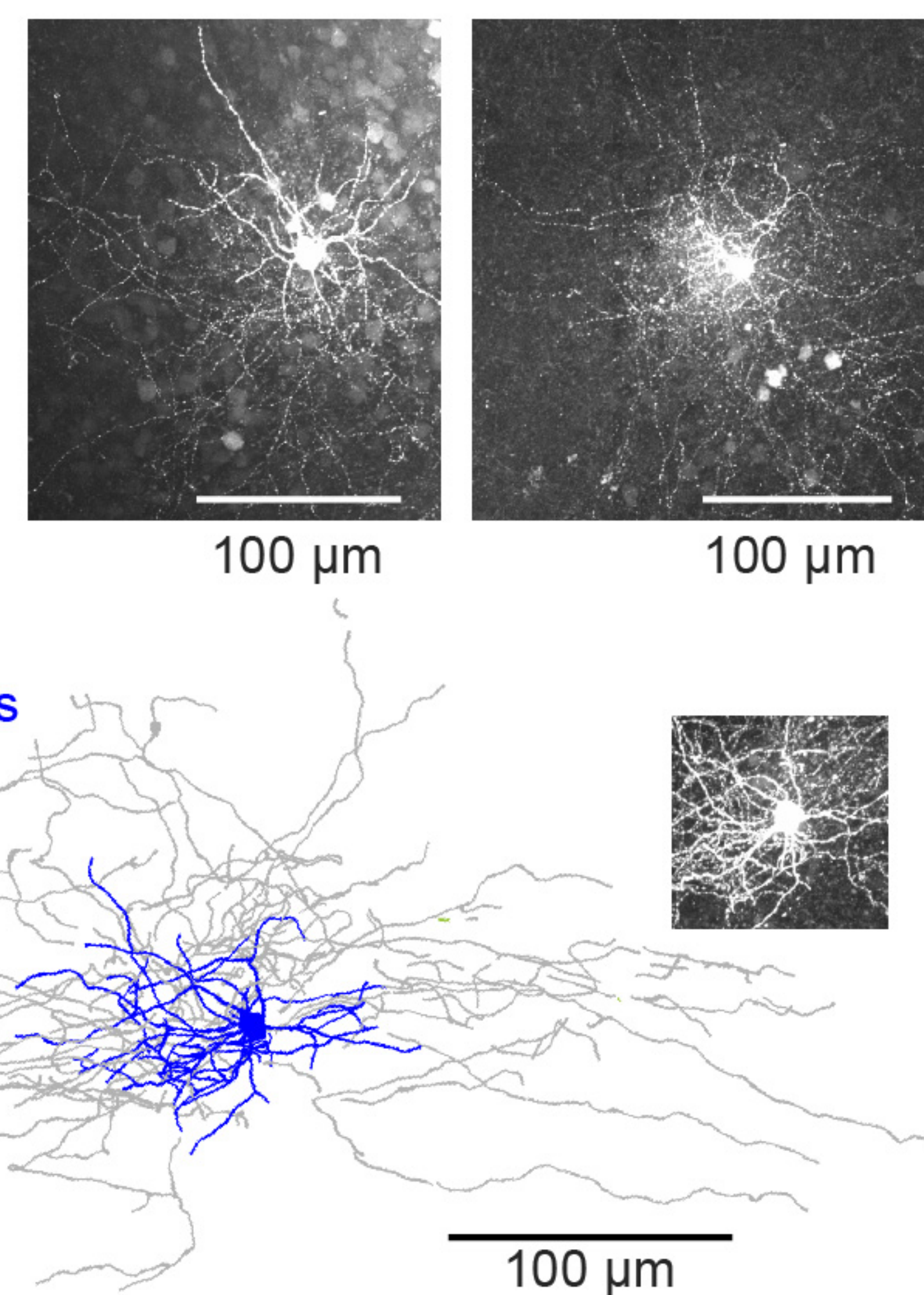
G



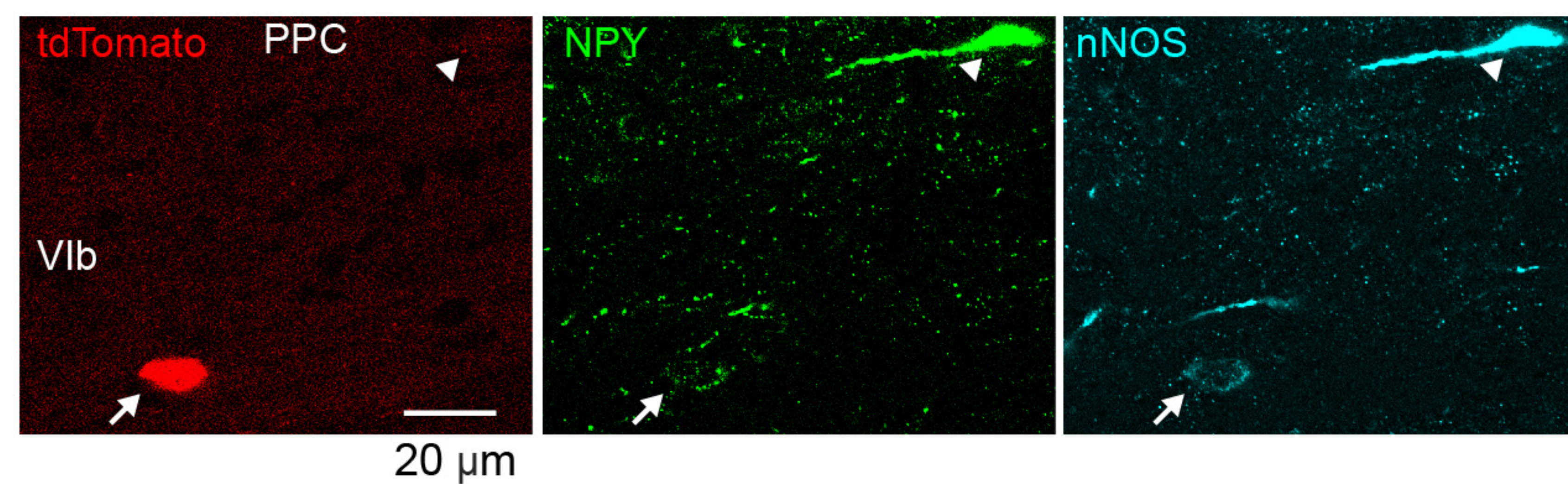
H



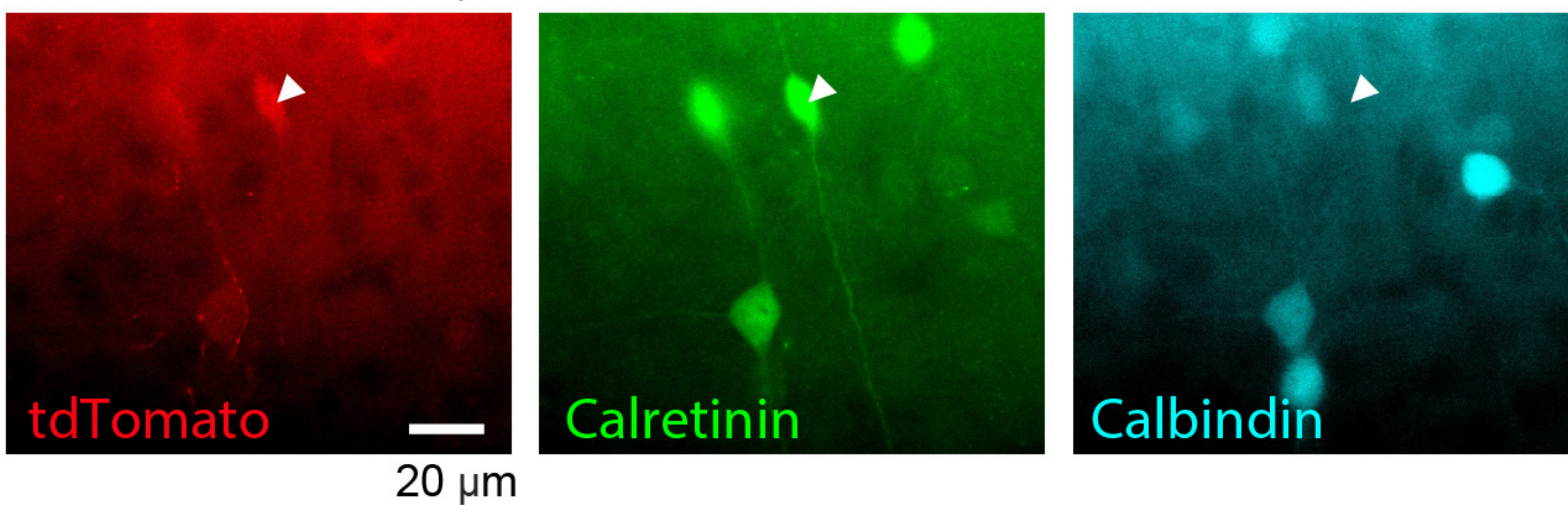
D

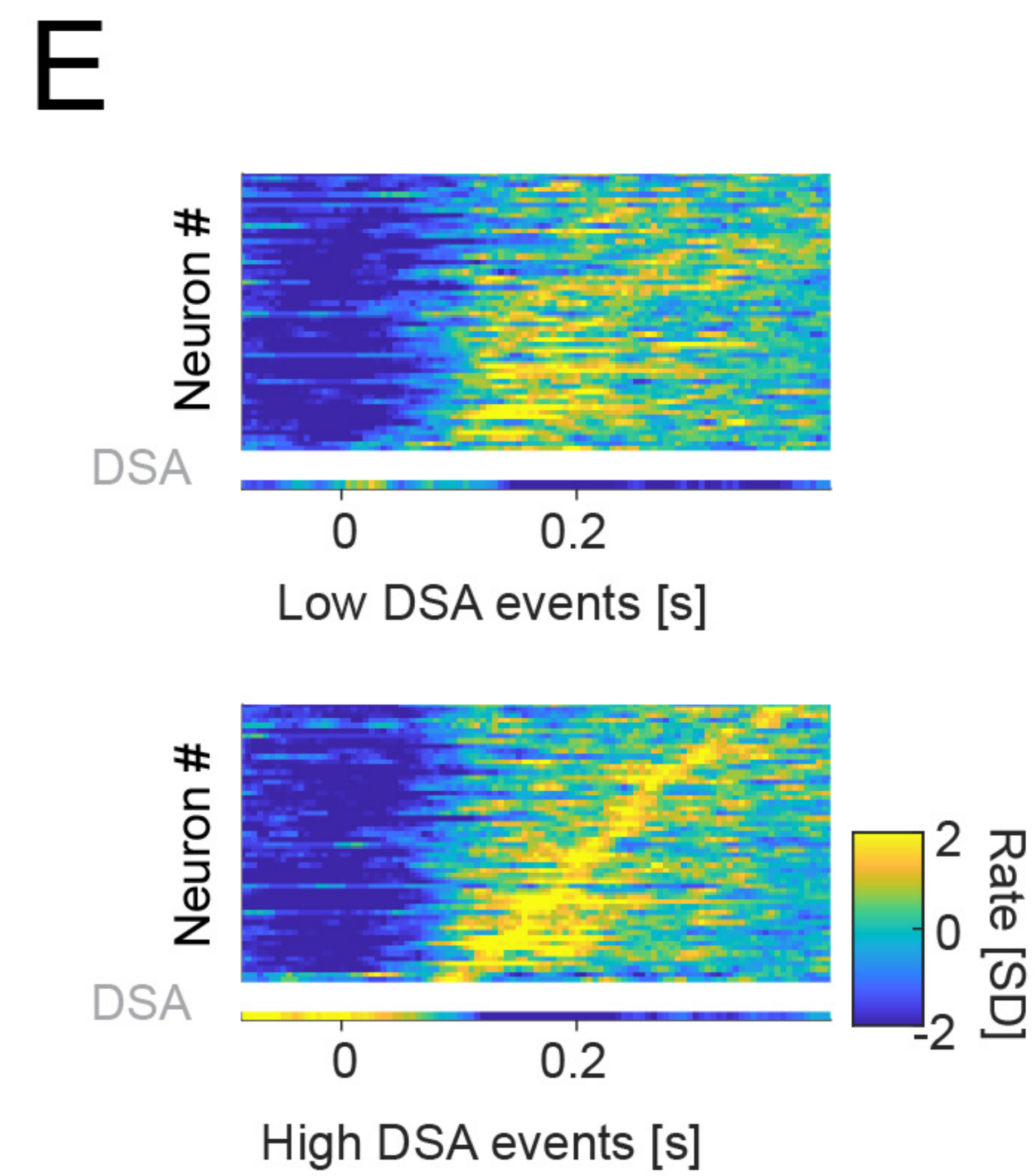
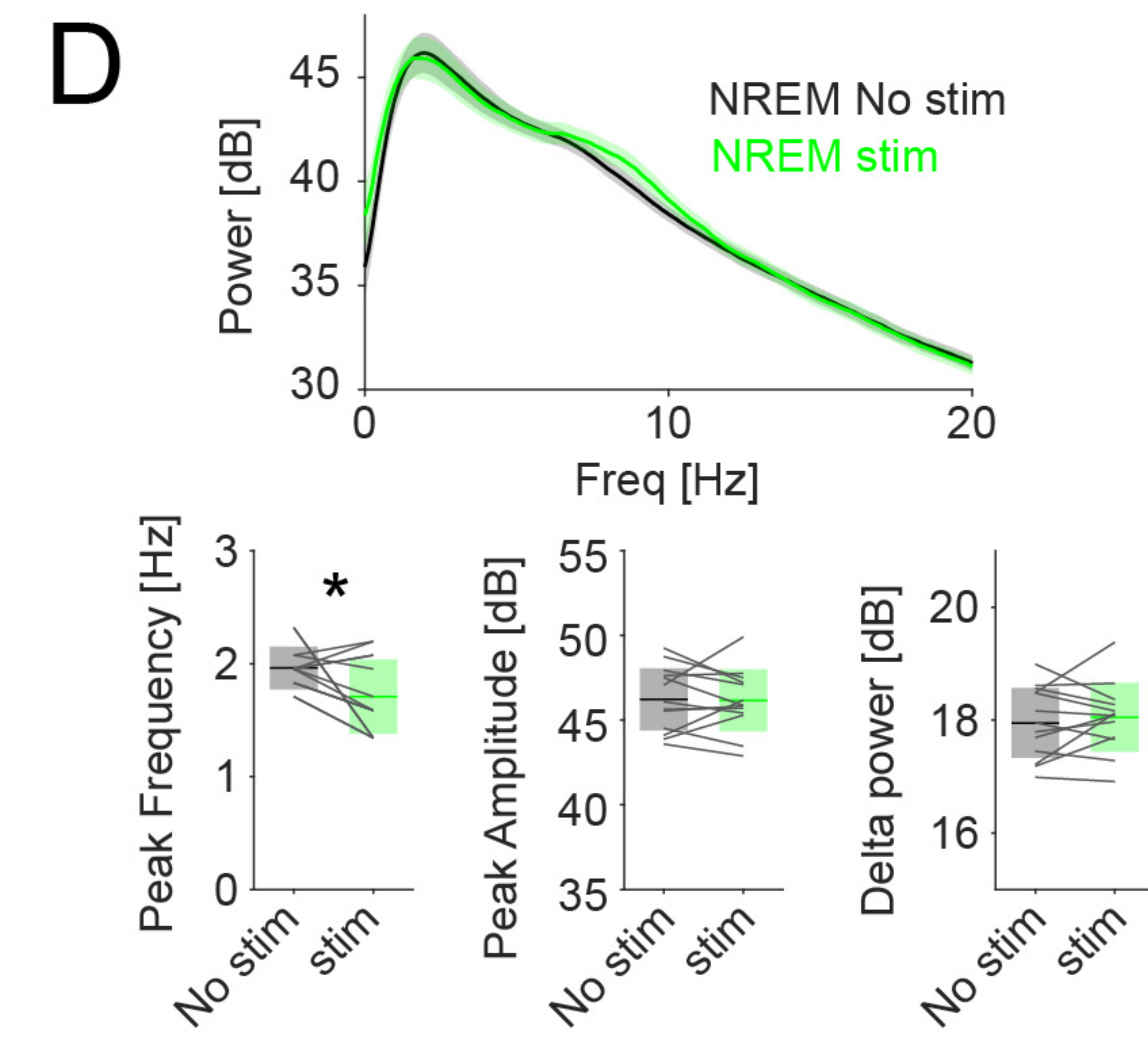
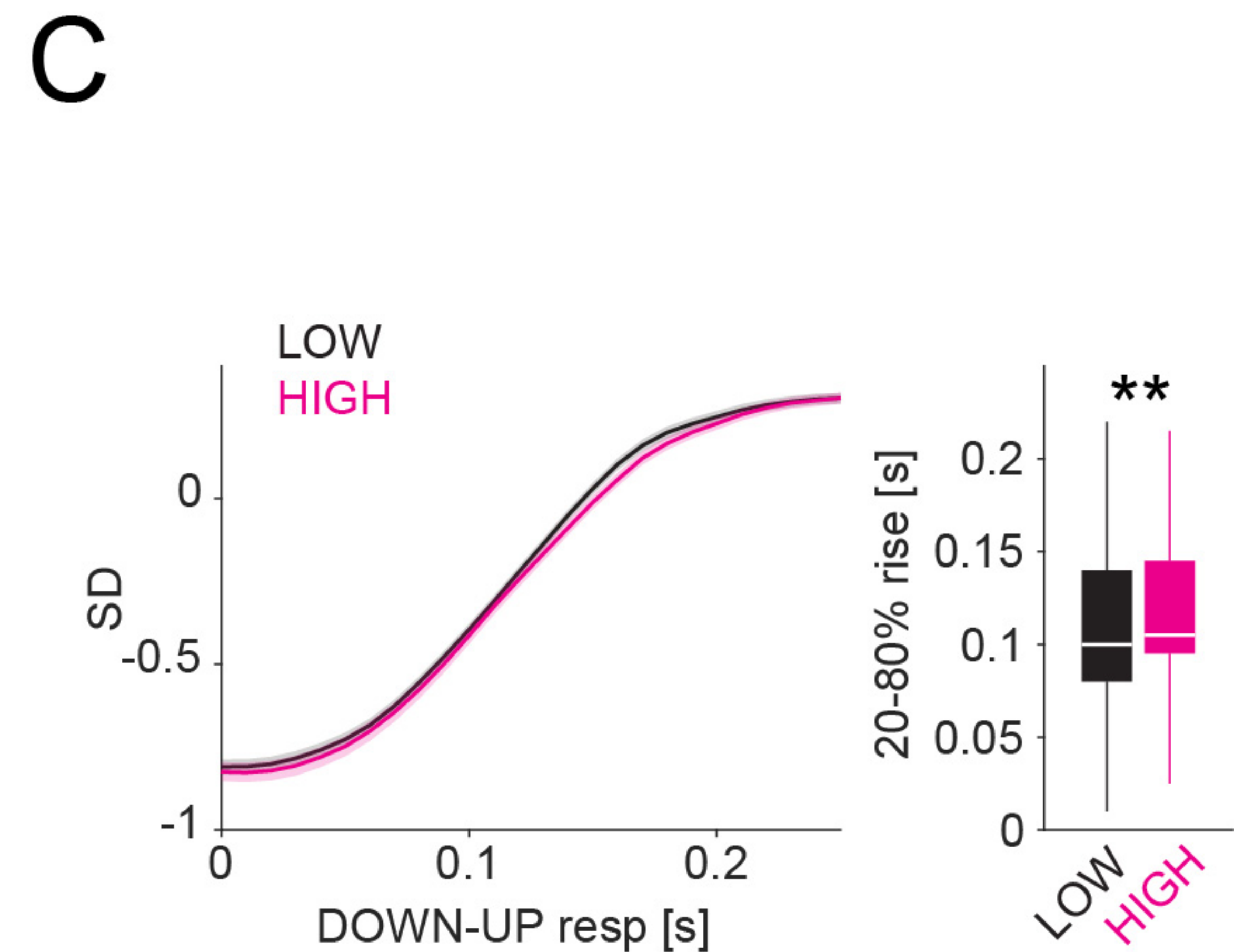
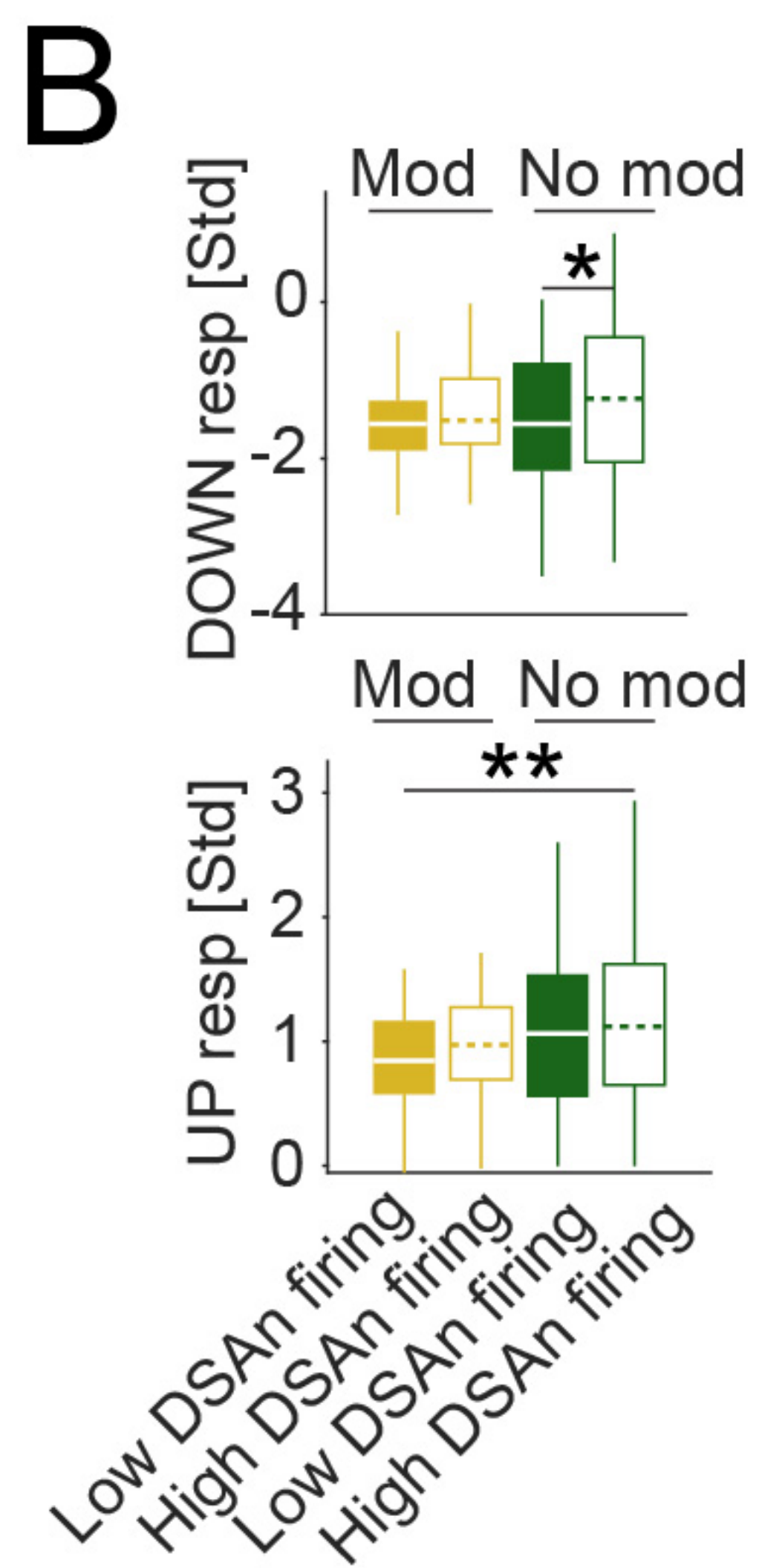
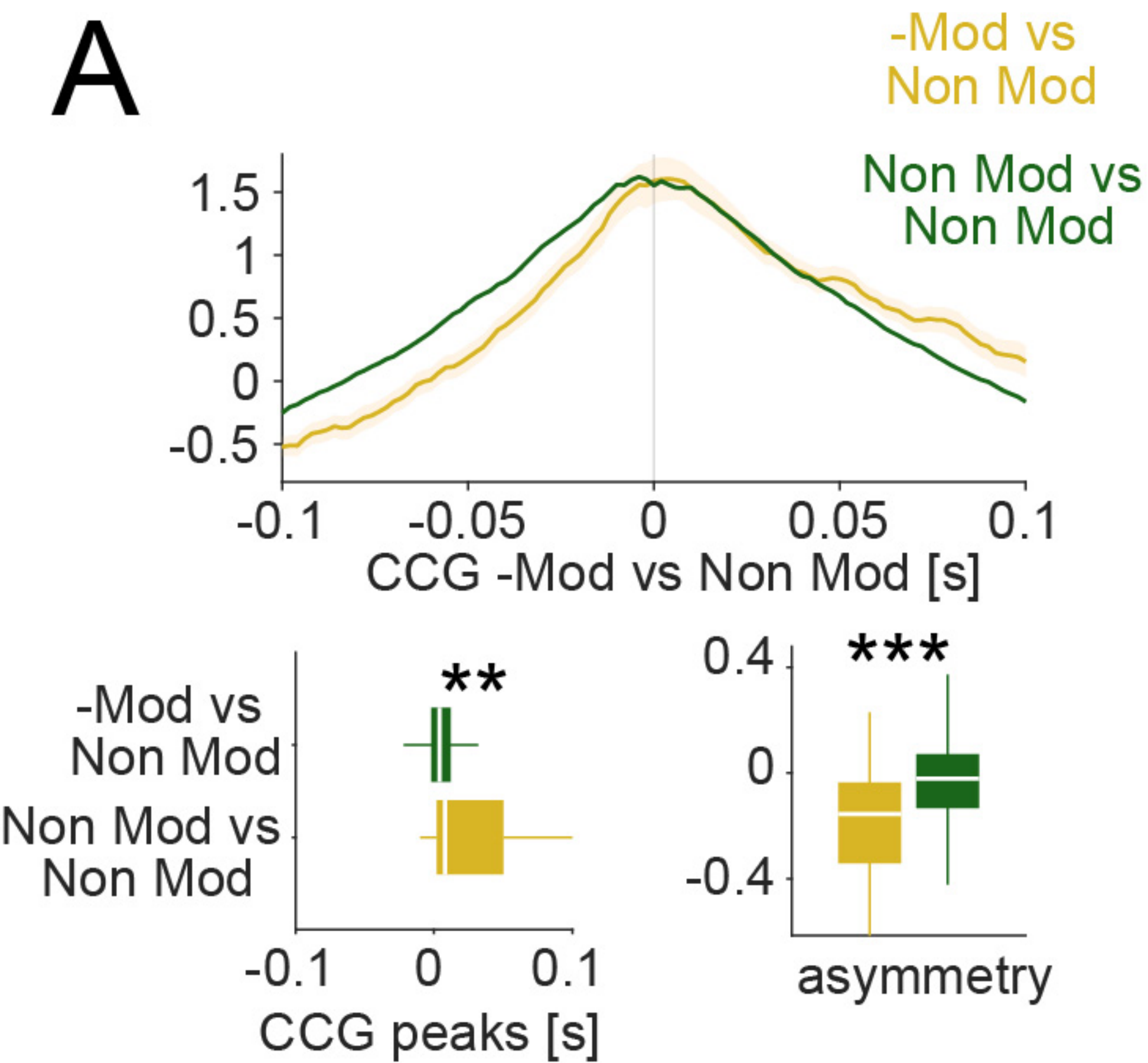


E

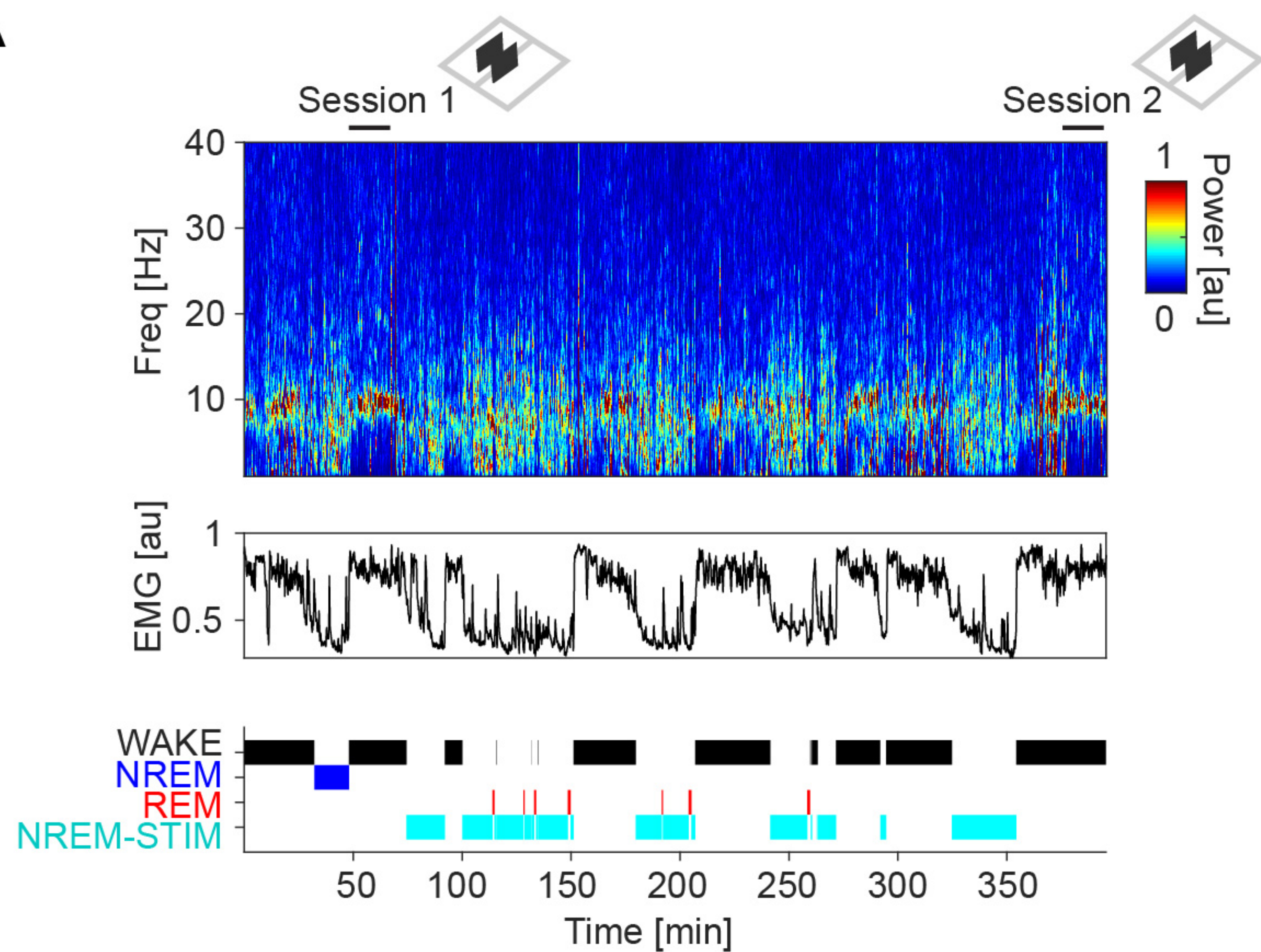


F

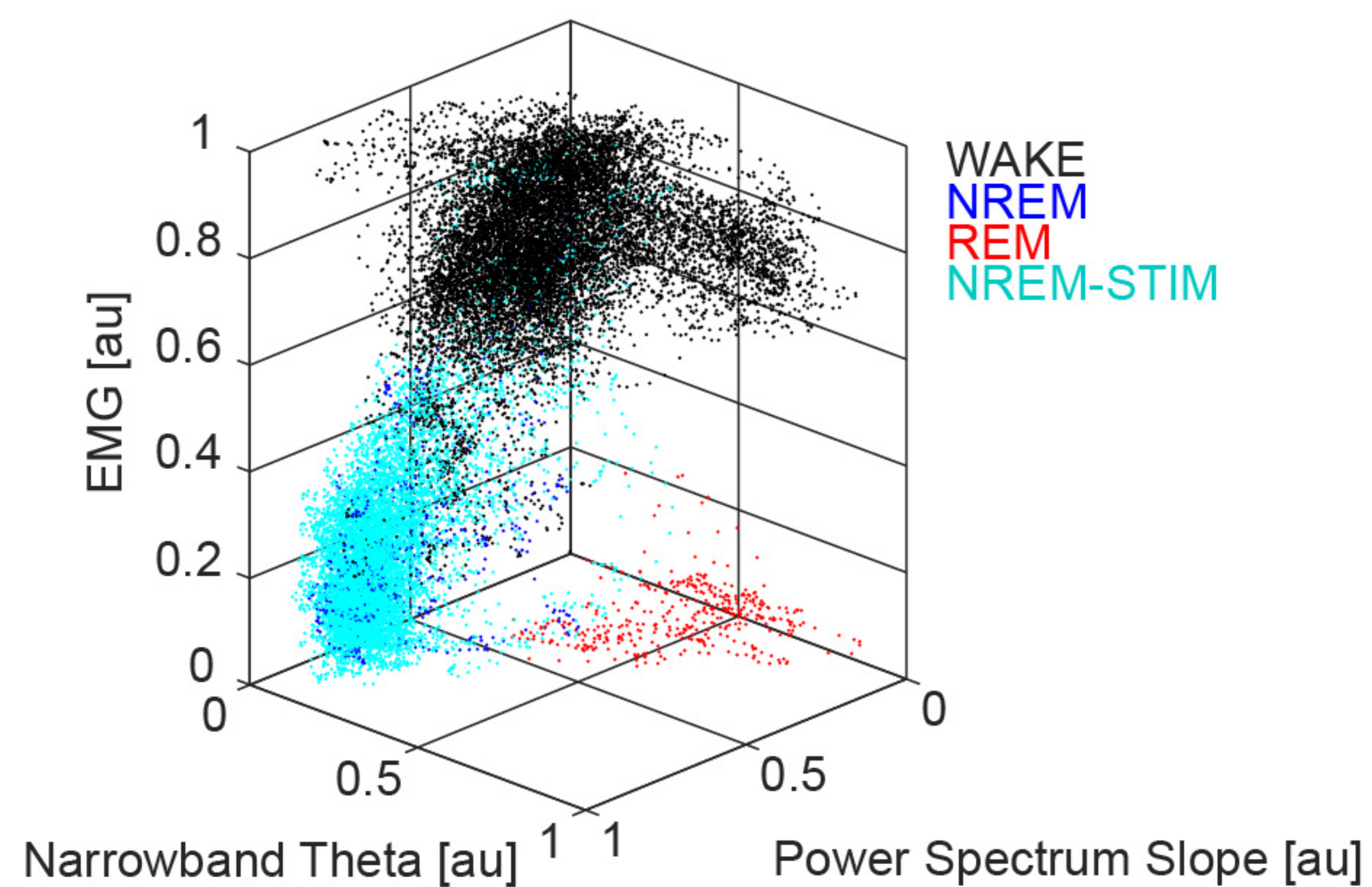




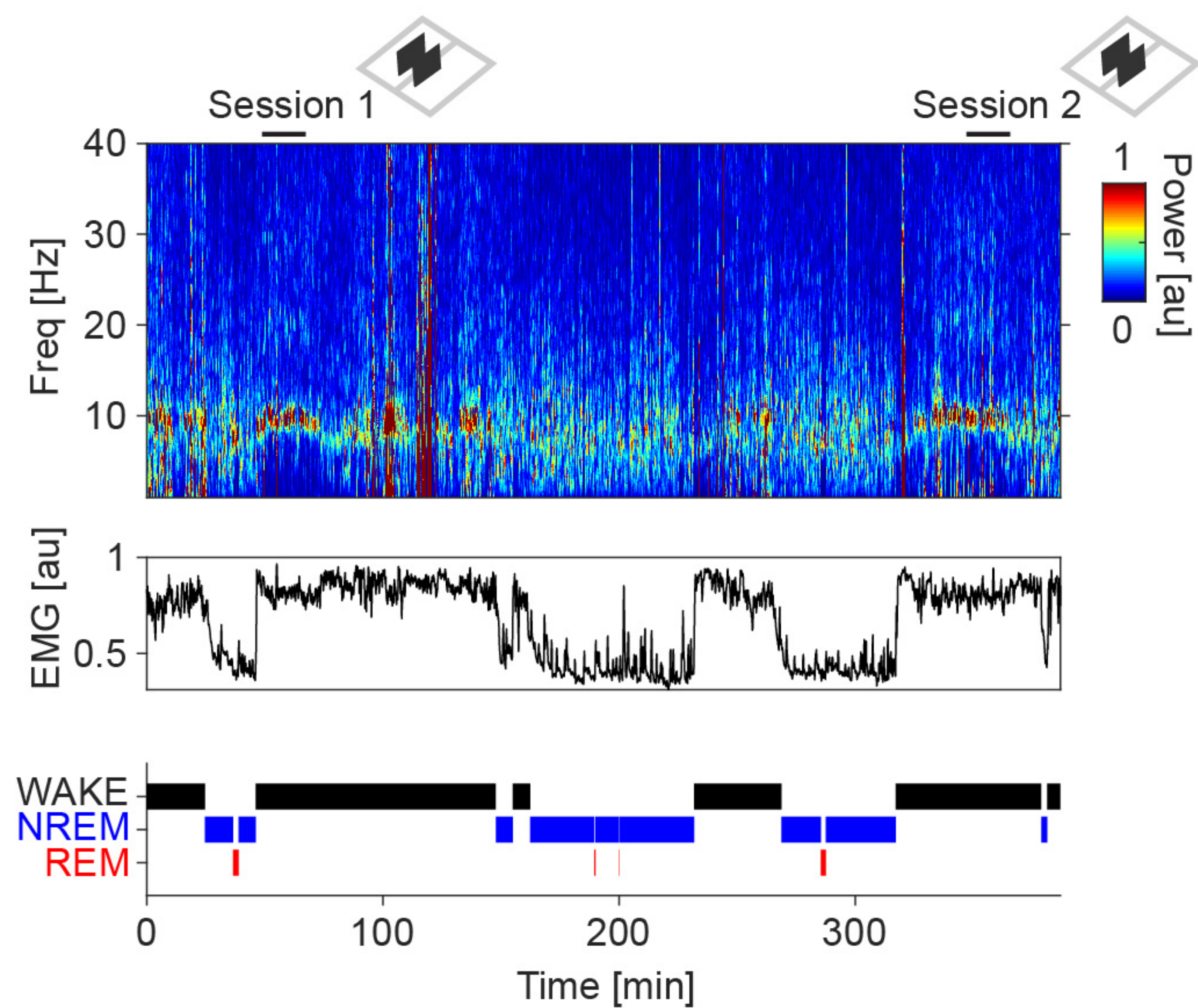
A



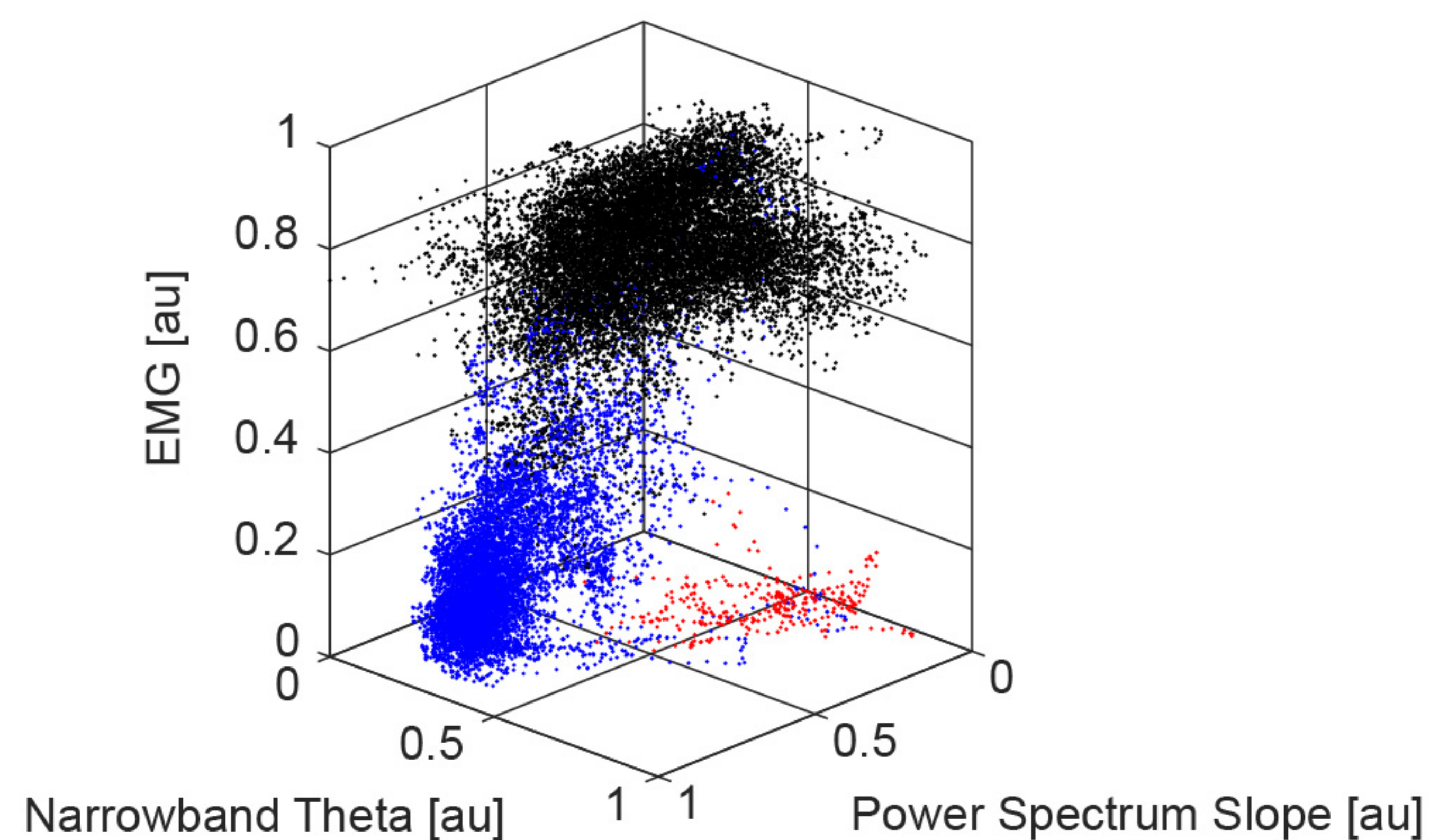
B



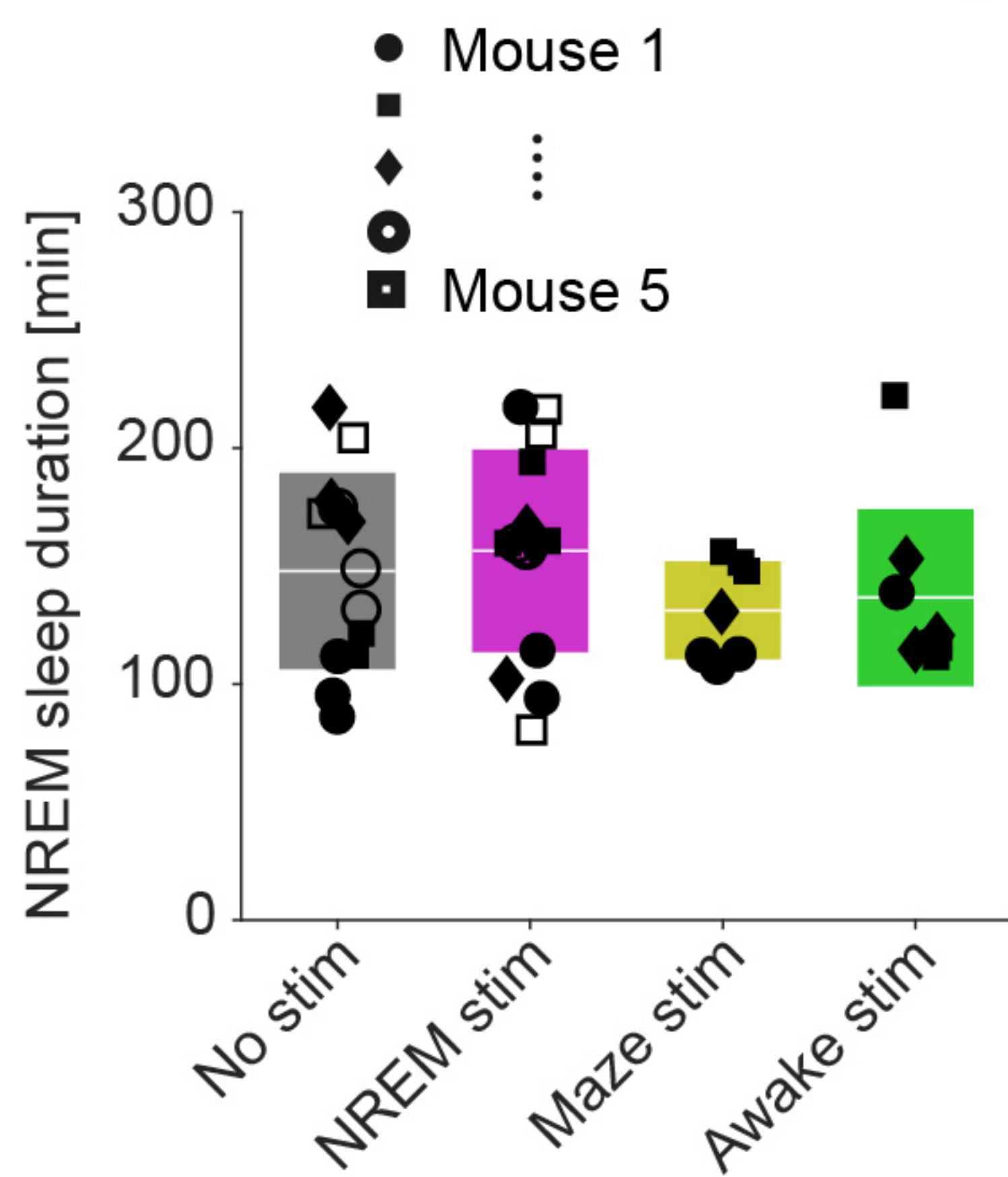
C



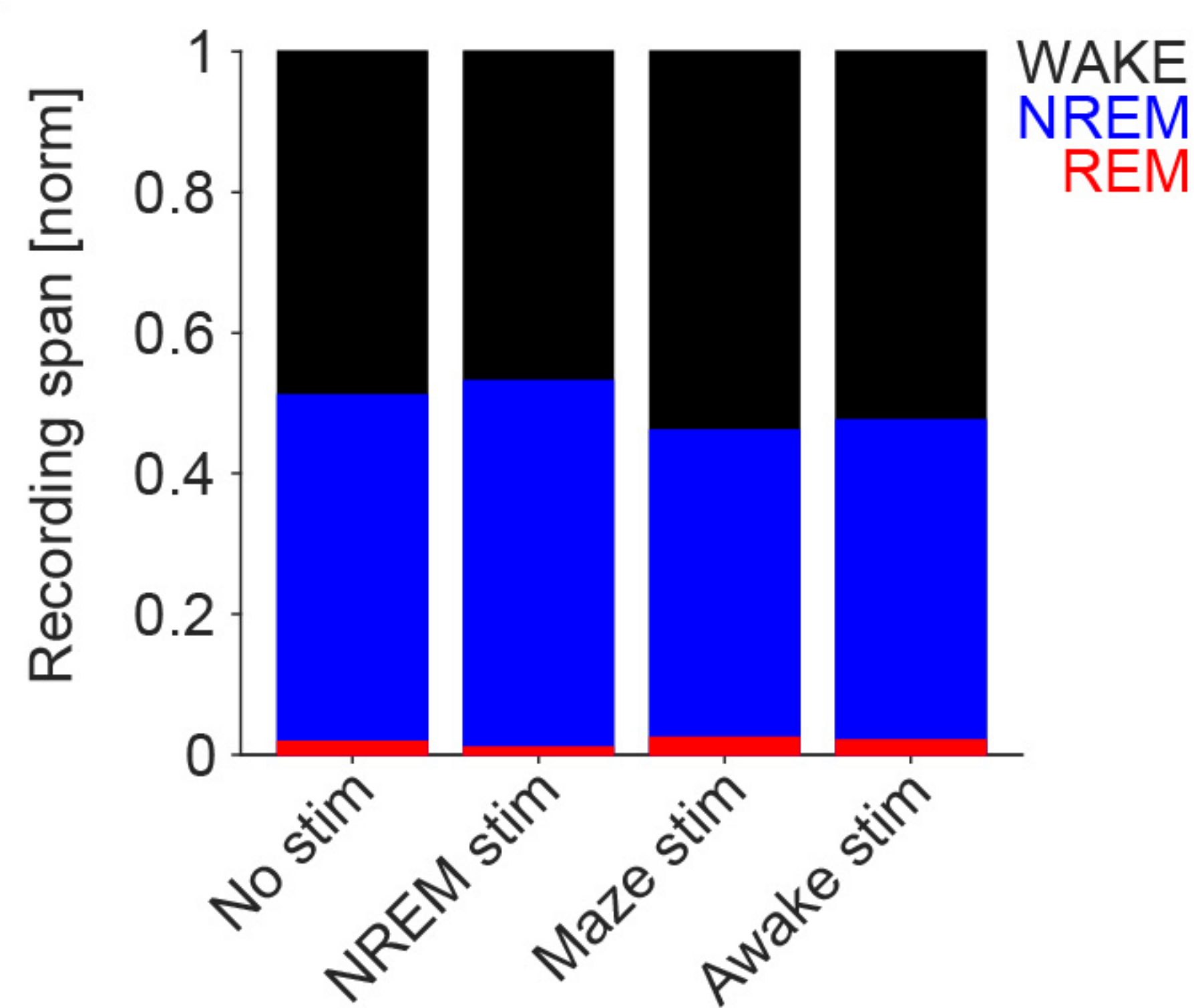
D



E



F



G

