WIDESPREAD RECEPTIVE FIELD REMAPPING IN EARLY VISUAL CORTEX

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ABSTRACT

Our eyes are in constant motion, yet we perceive the visual world as stable. Predictive remapping of receptive fields is thought to be one of the critical mechanisms for enforcing perceptual stability during eye movements. While receptive field remapping has been identified in several cortical areas, the spatiotemporal dynamics of remapping, and its consequences on the tuning properties of neurons, remain poorly understood. Here, we tracked remapping receptive fields in hundreds of neurons from visual Area V2 while subjects performed a cued saccade task. We found that remapping was far more widespread in Area V2 than previously reported and can be found in neurons from all recorded cortical layers and cell types. Surprisingly, neurons undergoing remapping exhibit sensitivity to two punctate locations in visual space. Furthermore, we found that feature selectivity is not only maintained during remapping but transiently increases due to untuned suppression. Taken together, these results shed light on the spatiotemporal dynamics of remapping

and its ubiquitous prevalence in the early visual cortex, and force us to revise current models of perceptual stability.

INTRODUCTION

Our early visual system is wired to store information in eye-centered (retinotopic) coordinates. With each movement of the eyes, the image falling onto the retina shifts rapidly, as does the visual information arriving at neurons in the cortex. Despite this, our perception of the world remains seamless and stable, implying that the visual system is able to compensate for self-generated movements. Previous research has suggested that receptive field (RF) remapping could contribute to this stability[1,2]. Remapping refers to the phenomenon in which neurons transiently shift their locus of spatial sensitivity (i.e. receptive field) before the onset of a saccadic eye movement towards their future, post-saccadic location. Remapping is considered to be a predictive mechanism because it both precedes and is temporally locked to eye movement initiation, and therefore requires advance information about both the timing and trajectory of an upcoming saccade. This information is thought to be conveyed through a corollary discharge signal originating in the brain regions responsible for initiating eye movements[1,3,4].

Receptive field remapping has been reported in many visual areas, including V1[5], V2[5], V3[5], V3A[5], V4[6,7], LIP[2], FEF[8,9], and SC[10-12]. In the cortex, early visual areas such as V1 and V2 are thought to have a low proportion of neurons that exhibit remapping, with higher order visual areas having a greater proportion[5]. Remapping has also been observed in humans using functional magnetic resonance imaging (fMRI)[13-15] and electroencephalography (EEG)[16-19], and an array of studies have reported on the behavioral consequences of remapping. Recently, it has also become clear that there may be multiple forms of remapping[6,7], with the exact mode of remapping

47 potentially depending on the brain region or cell type studied, or task demands. On one hand,
48 remapping to the future receptive field ('forward remapping'; **Figure 1A**) is thought to link pre-
49 and post-saccadic representations of visual space, thus helping to maintain perceptual continuity
50 and stability[2,5]. By contrast, remapping towards the saccade target ('convergent remapping';
51 **Figure 1B**) may serve to transiently enhance processing of visual information near that target[7,20,21].
52 More recent evidence has suggested that both forms of remapping may exist within the same
53 neurons[7]. There also remains concern that inconsistencies across some of these studies, which used
54 different experimental paradigms in different visual areas, could have contributed to these
55 divergent findings[22]. Furthermore, while work in human psychophysics has confirmed that
56 remapping preserves some[17,23] but perhaps not all[24] visual feature selectivity at the level of
57 perception, whether feature selectivity is preserved in individual neurons as they remap remains
58 an open question. Indeed, despite extensive study, much remains unknown about the properties
59 and extent of receptive field remapping, in large part due to limitations of the spatiotemporal
60 resolution at which the phenomenon was studied[25-28].

61 Here, we examined receptive field remapping in an early visual area (Area V2) with high-
62 density electrode arrays and a stimulation paradigm that provided significantly improved
63 spatiotemporal resolution. With this approach, we were able to track the time course of remapping
64 in discrete neural subpopulations in the laminar cortical circuit, allowing us to test whether
65 remapping is a global, trans-laminar phenomenon, or restricted to a particular cortical layer or cell
66 type. The use of oriented Gabor stimuli also allowed us to examine whether tuning for visual
67 features is altered during remapping. We found that remapping was far more prevalent in Area V2
68 than previously thought, and that it occurred in all recorded subpopulations. Neurons exhibit
69 transient sensitivity to two punctate locations in visual space during remapping, similar to activity

patterns found in the frontal cortex[29] but unlike that reported in Area LIP[30]. Furthermore, peri-saccadic firing rate suppression results in a transient increase in orientation selectivity during remapping.

RESULTS

We designed a cued saccade task, in which subjects held fixation for a variable delay period prior to initiating a saccade in response to a target point appearing in the periphery (**Figure 1C-D**). The simultaneous disappearance of the fixation point served as the go cue. After executing an accurate saccade, subjects then had to continue holding fixation at the target point to receive a reward. To prevent subjects from preemptively planning a saccade prior to the go cue, both the saccade target location and the delay period duration were pseudo-randomized. The target location was drawn from one of two possible locations, while the delay period duration was drawn from an exponential distribution. While the subjects executed these eye movements, oriented Gabor stimuli were continuously presented on a 13 x 13 grid spanning the visual region of interest at 60 Hz. On each frame of stimulus presentation, a single stimulus drawn from one of 6 random orientations was presented at a single grid location. Two rhesus macaques were trained to perform this task, and demonstrated consistent performance across trials and days (**Figure S1**).

While subjects performed the cued saccade task, we recorded neural activity from well-isolated single units in Area V2 using linear array electrodes (**Figure 1E**). The use of an artificial dura (**Figure S2A**) allowed us to clearly visualize the cortical vasculature, which provided landmarks for tracking successive probe insertion sites (**Figure S2B**). We confirmed that individual electrode penetrations were perpendicular to the cortical surface, and therefore in good alignment with individual cortical columns, by mapping receptive fields along the depth of the

cortex (**Figure S2C**). Laminar boundaries were identified with current source density (CSD) analysis[31,32] (**Figure S2D**), and units were classified as belonging to either the superficial (II/III), input (IV), or deep (V/VI) layers. Single units were also identified as either narrow- or broad-spiking based on their waveform duration (**Figure 1F-G**). In total, 923 single units were recorded, 822 of which were significantly visually responsive and included in our subsequent analyses.

Widespread Pre-Saccadic Remapping in Area V2

For each recorded single unit, we computed spatial sensitivity maps as a function of time relative to saccade onset (**Figure 2A; Figure S3A-C; Video S1 and S2**). We found that 73% of V2 units showed pre-saccadic remapping to the future receptive field (forward remapping) before the start of an eye movement. We found no evidence of widespread remapping towards the saccade target (convergent remapping), suggesting that this phenomenon may only arise in higher order visual areas. Using these spatial maps, we calculated the relative spatial distribution of sensitivity between the current and future receptive fields as a function of time (**Figure 2B; see Figure S3D for raw firing rate traces**). These time courses show that the handoff of spatial sensitivity begins well before saccade onset and has largely completed by the time the eyes begin to move. Surprisingly, our results suggest that for this brief period, these units are significantly responsive to two discrete locations in visual space. This pattern of spatial sensitivity transfer is not unique to any of the neural subpopulations that were recorded, and instead occurs in all three layers as well as in both broad- and narrow-spiking units (**Figure 2C**). Indeed, the timing of this transfer is also consistent across subpopulations, initiating approximately 40 ms before the saccade (**Figure 2D**), after spike binning normalizes for feed-forward signaling delays (**Figure S3C**).

Tracking Receptive Field Trajectories in Principal Component Space

To determine whether the transition in spatial activity patterns could be identified without pre-defined coordinates for the current and future fields, we performed unbiased dimensionality reduction analyses (principal component analysis, PCA) on our data. We vectorized the 13 x 13 stimulus grid into a 169-dimensional space with neural response values that changed over time for each unit; each timepoint was a single sample for the PCA. As all of the units from each session had largely overlapping receptive fields, and thus similar activity patterns in this space, they were fed into the PCA together. This approach produces a low-dimensional representation of receptive field trajectories during remapping (**Figure 3A; Figure S4A**) by, in effect, detecting the features of the spatial distribution of sensitivity (see Figure 2A for an example) that account for the greatest variance across time. Strikingly, the prominent features of this analysis are consistent across sessions, animals, and cortical layers, with a characteristic V-shaped trajectory (**Figure S4B; Figure S5; Figure S6**). These results can also be normalized and pooled for units across all sessions to produce the receptive field trajectory of an average V2 neuron during remapping (**Figure 3B**), which again illustrates this characteristic shape. To determine which features of the neural activity were being identified by the PCA, we compared values along the 1st PC dimension as a function of time with our previous current/future field sensitivity traces. We found that the 1st PC dimension closely tracks the relative distribution of spatial sensitivity between the current and future field (**Figure 3C**; Pearson's correlation = 0.996), remaining relatively stable until shortly before saccade onset and then shifting to values at the other extreme. Further examination of the 1st PC time courses revealed that this transition was consistent across all unit subpopulations (**Figure 3D-E**) and was initiated at a time that closely matches the results from our spatial map analysis (**Figure 3F**). The 2nd PC, on the other hand, appears to track the percentage of the total

response that is contained within the receptive field relative to the baseline response at non-receptive field positions (**Figure S4C**), which is an indirect measure of changes in peri-saccadic firing rate (**Figure S4D**) presumably caused by saccadic suppression. Therefore, it is the temporally overlapping effects of receptive field remapping and saccadic suppression that generate V-shaped trajectories in principal component space. Most notably, even with this approach that makes no assumptions about particular patterns of spatial sensitivity at particular moments in time, translational shifts to the future field remain the dominant feature of remapping in Area V2.

Firing Rate Suppression Drives a Transient Enhancement of Tuning During Remapping

To determine whether stimulus selectivity may be altered during remapping, we generated tuning curves for each unit during pre-saccadic, saccade planning, and post-saccadic periods (**Figure 4A**). The remapping period was defined as -75 to -25 ms relative to saccade onset to cover the transition between the current and future field, while the pre- and post-saccadic periods were well before and after saccade onset respectively. From these tuning curves, we computed the preferred orientation of each unit, as well as their orientation selectivity index (OSI; **Figure 4B-C**) and circular variance (**Figure 4D-E**). A higher OSI indicates a greater preference for one orientation over the orthogonal orientation. A circular variance of 1 would reflect equal responses to all orientations, while a circular variance of 0 would reflect responsiveness to only a single orientation. When comparing OSI and circular variance across the three conditions, we found that orientation tuning was transiently increased during saccade planning before returning to baseline levels. We next asked whether this increase in tuning could be the result of a firing rate change. Population averaged tuning curves revealed that the firing response to stimuli at both the preferred and non-preferred (orthogonal) orientations were suppressed during remapping (**Figure 4F-G**). Fitting the data from

each unit with a Gaussian tuning curve also showed the same pattern of suppression (**Figure 4H**). Quantifying the half width at half height (HWHH) from the fitted curves revealed no changes in the overall shape of the tuning curves (**Figure 4I**). Thus, this change in tuning is largely driven by untuned suppression, as firing in response to both preferred and non-preferred orientations is suppressed, and not by divisive changes to the shape of the tuning curve, as the HWHH of the tuning curves remains unchanged.

168 **DISCUSSION**

169 We used high density receptive field mapping during a cued saccade task in combination with
170 laminar electrophysiology to study receptive field remapping in Area V2. We found that
171 remapping was widespread in Area V2, and was present across all recorded cortical layers
172 (superficial, input, deep) and unit types (narrow- and broad-spiking). We identified forward
173 remapping towards the future receptive field location as the dominant mode of remapping in Area
174 V2. We demonstrated that these findings were not contingent on any assumptions about receptive
175 field structure or direction of remapping, as an unbiased dimensionality reduction approach arrives
176 at the same result. Further, we tested for changes in tuning during saccade planning and execution,
177 and found that a brief suppression of firing rates drives transiently increased tuning during saccade
178 planning.

179 Our data demonstrate that receptive field remapping is far more widespread in the early
180 visual cortex than was previously appreciated. Prior work has suggested that the proportion of
181 neurons that underwent remapping increased progressively along the visual stream, with early
182 visual cortex, including areas such as V1 and V2, having relatively few neurons that remapped[5].
183 It was thought that this low proportion may have reflected the prevalence of remapping in a
184 particular cell type, and the absence of remapping in others. However, our results indicate that
185 remapping is a global phenomenon that occurs in a majority of neurons in Area V2 and is not
186 restricted to a particular cell class or layer. This discrepancy can likely be explained by differences
187 in experimental approach. The only prior electrophysiological study to test for remapping in Area
188 V2[5] presented stimuli at just four timepoints relative to a go cue, resulting in a relatively course
189 temporal sampling. Given technical limitations at the time, the neural responses to these stimuli
190 were not aligned to saccade onset, and trial-to-trial variability in behavioral response times may

therefore have also limited the temporal clarity of the data. Flashing of single probes at discrete timepoints may also draw attention to the probe, resulting in attentional remapping that may affect visual responses[18]. In our study, we are able to generate continuous time courses of spatial sensitivity at both the current and future receptive field that are aligned to the onset of a saccade, and with this approach we observe a much higher proportion of neurons undergoing remapping.

The functional role of remapping remains a largely open question, although the prevailing hypothesis is that remapping plays a significant role in maintaining perceptual stability across eye movements[1,2]. Forward remapping towards the future receptive field may allow for visual processing before the saccade to occur in a post-saccadic frame of reference, thereby allowing the visual system to maintain a spatial frame of reference that would otherwise be disrupted by an eye movement. Given the existence of multiple modes of remapping[6,7] and presumed differences in their relative prevalence across cortical areas, it is also possible that the functional significance of remapping varies from region to region. Given the shifts in receptive field location, as well as the potential for spatial sensitivity at split locations in space, remapping may also be responsible for the mis-localization of stimuli around the time of saccades[33-35].

Given the prevalence of remapping in Area V2, it is also worth considering whether remapping may be important in other early visual cortical areas, such as Area V1. Remapping is often considered to be a result of, or at least a correlate to, the allocation of spatial attention[18,19,36,37]. However, V1 and V2 serve different roles in attentional allocation, with V1 generating a saliency map that guides attention[38-42], while V2 is not known to be significantly involved. Thus, remapping may be an undesirable property in the V1 neural population that conflicts with its role in attention guidance, resulting in very few V1 neurons showing remapping. Nonetheless, further studies are needed to identify possible differences in remapping along the visual hierarchy.

Corollary discharge signaling from the superior colliculus through the thalamus is thought to be responsible for initiating receptive field remapping in the cortex. Pharmacological interventions have demonstrated that thalamic inactivation impairs performance in a double-step saccade task[4] and limits cortical remapping[29], confirming the role of the thalamus as a relay station for this signal. Recent evidence suggests that thalamic projections to the visual cortex may also be used for distinguishing between self-generated and saccade-generated visual motion in both primates[43] and rodents[44]. Interestingly, there may also be a high degree of redundancy in pathways for the updating of peri-saccadic spatial information, as behavioral performance in a double-step saccade task was found to be impaired in split-brain monkeys, but could be recovered substantially with training[45]. Our results are consistent with a thalamic origin for the signal initiating receptive field remapping. Our timing analysis, which normalizes for feedforward timing delays as a byproduct of spike binning, found no significant differences in the onset of receptive field remapping across layers. This suggests that the feedforward transfer of visual information across layers, starting with the input layer and then spreading to the superficial and deep layers, also carries the remapping signal. Thus, the remapping signal appears to first arrive in the input layer of V2, which is where thalamic inputs to V2 terminate in both macaques and other primates[46-48]. Recent evidence from an intermediate visual region, Area V4, suggests that thalamic projections to the input layer may also be responsible for initiating saccadic suppression[43], raising the possibility that a common signaling pathway from the thalamus may be responsible for both phenomena. An alternative possibility is that receptive field remapping in V2 is the result of feedforward input from V1. However, we consider this to be the less likely option given the importance of V1 in maintaining an attentional saliency map, as discussed above, and the much

236 stronger input that V2 receives from the pulvinar[47], the thalamic nucleus thought to relay saccade-
237 related signals.

238 One understudied aspect of remapping has been the question of whether feature selectivity
239 remaps alongside the spatial receptive field. This question is of critical behavioral importance, as
240 perceptual stability requires the ability to identify stimuli as well as to locate them. Thus far,
241 research on this topic has been both sparse and conflicting. In LIP, it is thought that stimulus tuning
242 is preserved during remapping[49], while in MT, there is no evidence of tuning in remapped fields[50].
243 Here, we show that one prominent aspect of feature encoding in Area V2, orientation tuning,
244 persists during remapping. Indeed, we find that orientation selectivity transiently increases due to
245 the suppression of overall firing during the saccade planning period.

246 Together, our results reveal the widespread nature of receptive field remapping in the early
247 visual cortex and suggest that the fundamental computations underlying perceptual stability are
248 enacted from these early stages. Furthermore, we demonstrate that remapping overlaps and
249 interacts with changes in feature tuning that are driven by saccadic suppression of neural firing.
250 The cortical column-wide nature of the changes suggests that remapping is conveyed as a global
251 signal to the early visual cortex. The fact that neurons exhibit transient split sensitivity to two
252 punctate locations in space necessitates a rethinking of the nature and functional role of remapping.
253 Further experiments are needed to both fully characterize sensitivity and tuning changes at a sub-
254 receptive field level and to elucidate the neural circuits that enable these phenomena.

255

256 **Limitations of the study**

257 This study has several limitations that should be considered alongside our findings. For one, our
258 approach treats the receptive field of V2 neurons as homogenous and is unable to resolve potential

differences in remapping across subfields. Second, it is unknown whether neural response latencies may change during remapping, and we assume that they are constant when settings our binning windows (Figure S3A-B). And lastly, despite our approach providing us with cell type- and layer-specific insights into remapping, we remain limited by the tools available to us and are unable to causally link these effects to a specific signaling pathway.

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AUTHOR CONTRIBUTIONS

SD & ASN conceptualized the project. SD, MPM, and NVH collected the data. SD analyzed the data. ASN supervised the project. SD & ASN wrote the manuscript.

DECLARATION OF INTERESTS

The authors declare no competing interests.

INCLUSION AND ETHICS

We support inclusive, diverse, and equitable conduct of research.

FIGURE LEGENDS

Figure 1. Task Design and Single Unit Recordings

(A) Forward remapping shifts the current receptive field by the vector of the upcoming saccade to form a forward field.

(B) Convergent remapping shifts the current receptive field towards the saccade target to form a convergent field.

(C) Progression of a trial during the cued saccade task as the subject holds fixation, executes a saccade to the target, and then fixates on the target. The period in blue indicates probe presentation. A reward is delivered after fixating on the target for 500 ms.

(D) Fixation point, saccade targets, and stimulus grid layout during the cued saccade task.

(E) Snippet of data from one probe shank during a trial of the cued saccade task. LFP traces are color by cortical layer, and spikes are overlaid on their channel of origin as vertical lines. Red arrows indicate synchronized spiking and local field potential deflections along the depth of the cortex in response to a stimulus being flashed in the receptive field of the recording site.

(F) Action potential waveforms of all 923 single units. Units were classified as either narrow-spiking (blue) or broad-spiking (orange) on the basis of their peak-to-trough waveform duration.

(G) Distribution of waveform durations for single units from (F). The distribution shows bimodality for the two unit types (Hartigan's dip test; $p = 1.15*10^{-4}$).

Figure 2. Receptive Field Remapping is Widespread in Area V2

(A) Receptive field location for an example single unit during remapping. Times are relative to saccade onset. Each row shows remapping during saccades to one of the two saccade targets.

Fixation point and saccade target are overlaid in green and blue, respectively. Heatmaps at each timepoint are individually normalized to account for possible changes in firing rate.

(B) Normalized sensitivity at the current and future field locations for all single units (n = 822), as a function of time relative to saccade onset. Error bars indicate standard error of the mean.

(C) Normalized sensitivity at the current and future field locations as a function of time relative to saccade onset for each of the recorded neural subpopulations. Error bars indicate standard error of the mean.

(D) The time relative to saccade onset at which remapping initiates for each of the recorded neural subpopulations. Error bars indicate bootstrapped 95% confidence intervals.

Figure 3. Tracking Remapping Trajectories in Principal Component Space

(A) Left, principle component trajectories of all single units from an example session. Right, principle component trajectory of an example single unit from the same session. Top, target 1. Bottom, target 2. Each point represents a single timepoint from a single unit. On average across sessions and targets, PC1 explains 11.3% of the variance, and PC2 explains 3.3% of the variance.

(B) Averaged remapping trajectory in principal component space across all single units for both targets.

(C) Correlation between values along the 1st principle component axis and sensitivity at the future field (Figure 2B). Pearson's correlation = 0.996.

(D) Time course of values along the 1st principle component axis for all recorded layers. Error bars indicate standard error of the mean.

(E) Time course of values along the 1st principle component axis for both recorded unit types. Error bars indicate standard error of the mean.

(F) The time relative to saccade onset at which remapping initiates for each of the recorded neural subpopulations based on the 1st PC timecourses. Error bars indicate bootstrapped 95% confidence intervals.

Figure 4. Orientation Selectivity is Transiently Increased During Saccade Planning

(A) Orientation tuning curves from four example single units during the pre-saccadic (-250 to -200 ms), saccade planning (-75 to -25 ms), and post-saccadic (150 to 200 ms) time periods. Times are relative to saccade onset. FR = firing rate.

(B) Distribution of orientation selectivity index (OSI) for all single units during the pre-saccadic period.

(C) Change in OSI during the saccade planning and post-saccadic periods, as compared to the pre-saccadic period. Error bars indicate bootstrapped 95% confidence intervals.

(D) Distribution of circular variance for all single units during the pre-saccadic period.

(E) Change in circular variance during the saccade planning and post-saccadic periods, as compared to the pre-saccadic period. Error bars indicate bootstrapped 95% confidence intervals.

(F) Change in firing rate in response to presentation of a stimulus at the preferred orientation during the saccade planning and post-saccadic periods, as compared to the pre-saccadic period. Error bars indicate bootstrapped 95% confidence intervals.

(G) Change in firing rate in response to presentation of a stimulus at the non-preferred (orthogonal) orientation during the saccade planning and post-saccadic periods, as compared to the pre-saccadic period. Error bars indicate bootstrapped 95% confidence intervals.

(H) Average of Gaussian tuning curve fits across all single units during the pre-saccadic, saccade planning, and post-saccadic time periods. Error bars indicate standard error of the mean.

(I) Change in half width at half height (HWHH) of the fitted tunning curves during the saccade planning and post-saccadic periods, as compared to the pre-saccadic period. Error bars indicate bootstrapped 95% confidence intervals.

Figure S1. Behavioral Performance

(A) Receptive field and saccade target locations for all sessions (left) and one sample session (right).

(B) Distribution of saccade landing errors for both subjects. Landing error was defined as the distance between the saccade target and the terminal point of the saccade.

(C) Distribution of reaction times for both subjects.

(D) Relationship between the trial-by-trial pre-saccadic fixation time (pseudorandomly determined for each trial) and reaction time. The dashed red line depicts a linear least-squares fit. The flat relationship between fixation time and reaction time suggests that subjects are not able to anticipate the timing of the go cue.

Figure S2. Recording Methodology

(A) Schematic of the artifical dura and recording chamber (see Methods).

(B) Microscope image inside the recording chamber of one subject. Each 'X' indicates a cortical recording site. The inset shows a close-up image of the linear array probe penetration at the site marked in red. The black line is the estimated boundary between Area V1 and Area V2 based on the distinct change in surface vasculature density between the two areas.

(C) Receptive fields along the depth of one shank as computed from local field potential deflections. dva = degrees of visual angle

(D) Current source density (CSD) analysis of the shank from (C). Blue indicates a current source, while red indicates a current sink. Dashed lines indicates laminar boundaries, as determined from the CSD. White bar indicates duration of stimulus presentation.

(E) Kernel density distribution estimates for several metrics[51-53] that quantify recording stability and single unit quality. Presence ratio reflects the proportion of a session during which a unit was present and firing action potentials. Maximum drift is the distance between the highest and lowest channels on which a unit was detected during a session. Isolation distance and d-prime quantify the separation of spike waveform clusters in principal component space.

Figure S3. Stimulus-evoked firing rates at current and future fields

(A) Average firing response to stimulus flashes in the current receptive field during a baseline period before presentation of the go cue (n = 822 single units). Dashed vertical lines indicate the start and end of the binning window. Error bars indicate 95% bootstrapped confidence intervals.

(B) Same as in (A), but separated by cortical layer (superficial, n = 303 single units; input, n = 321 single units; deep, n = 198 single units). Dashed vertical lines indicate the start and end of the binning window. Error bars indicate 95% bootstrapped confidence intervals.

(C) Time of peak firing relative to stimulus onset for data in (B). Error bars represent bootstrapped 95% confidence intervals (superficial, n = 303 single units; input, n = 321 single units; deep, n = 198 single units).

(D) Firing rate responses at current and future field locations (n = 822 single units). Contrast with sensitivity plots, which are normalized at each timepoint (Figure 2B). Error bars indicate standard error of the mean.

Figure S4. Remapping Trajectories are Tracked in Principal Component Space

(A) Variance explained by each principal component, averaged across sessions and targets. Error bars indicate bootstrapped 95% confidence intervals.

(B) Averaged remapping trajectory in principal component space across all single units for both targets when PC analysis is performed on units from a given layer.

(C) Correlation between values along the 2nd principal component axis and the percent of total response contained within the current and future fields (as opposed to firing evoked by stimuli landing outside the receptive field). Pearson's correlation = -0.890.

(D) Correlation between values along the 2nd principal component axis and firing rate. Pearson's correlation = 0.512.

Figure S5. Principle Component Trajectories of All Sessions

Receptive field remapping trajectories in principal component space of all sessions. Across sessions, the data consistently shows a V-shaped trajectory in this space as remapping occurs.

Figure S6. Average Principle Component Trajectories of All Sessions

Average receptive field remapping trajectories in principal component space of all sessions.

Video S1. Continuous Tracking of Receptive Field Remapping

Spatial location of an example unit's receptive field sensitivity as a function of time (bottom left). Times are relative to saccade onset.

Video S2. Continuous Tracking of Receptive Field Remapping

Same as Video S1, but for another example unit from a different session.

STAR METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Experimental models: Organisms/strains		
Rhesus Macaques (Macaca mulatta)	Worldwide Primates	n/a
Chemicals		
PEDOT:PSS	Sigma-Aldrich	655201
Software and algorithms		
MATLAB	Mathworks	R2019a
Python	python.org	n/a
Kilosort2	Pachitariu M & MouseLand	https://github.com/MouseLand/Kilosort
phy (spike curation GUI)	Rossant C & cortex-lab	https://github.com/cortex-lab/phy
Custom code and analyses	This paper	https://doi.org/10.5281/zenodo.11044507
Other		
Silicon probes	NeuroNexus	a2x32_6mm35_200_177
RHD 512 channel recording controller	Intan	C3004
64 channel recording headstages	Intan	C3315
nanoZ	White Matter LLC	n/a

RESOURCE AVAILABILITY

Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Sachira Denagamage (sachira.denagamage@yale.edu).

Materials availability

This study did not generate new unique reagents or materials.

Data and code availability

All data reported in this paper will be shared by the lead contact upon request. All original code has been deposited at Zenodo and are publicly available as of the date of publication. DOIs are listed in the key resources table. Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Two male rhesus macaques (*Macaca mulatta,* D: age 6, M: age 8) were used as subjects in this study. All experimental procedures were approved by the Institutional Animal Care and Use Committee at Yale University, and conformed to NIH guidelines.

METHOD DETAILS

Experimental design

The study did not involve randomization or blinding, and we did not estimate sample-size before carrying out the study. No subjects or data were excluded from the study.

Surgical procedures

Surgical procedures were similar to those described previously [32,54,55]. Low-profile titanium recording chambers were implanted in two rhesus macaques. Using preoperative structural MRI and sulcal reconstruction, the chambers were targeted over the lunate sulcus, allowing access to Area V2 (left hemisphere in monkey M, right hemisphere in monkey D). The native dura mater overlying this region was removed and replaced with a transparent silicone artificial dura (AD). The AD allowed for visualization of area V2 and facilitated the targeting of electrode arrays.

Electrophysiology

453 Prior to recording, 64-channel electrode arrays ('laminar probes'; NeuroNexus Technologies, Inc.;
454 2 shanks, 32 channels/shank, 70 μm site spacing, 200 μm shank spacing) were electroplated
455 (nanoZ, White Matter LLC) in a solution of poly(3,4-ethylenedioxythiophene) polystyrene
456 sulfonate (*PEDOT*:PSS). At the start of each recording session, a laminar probe was lowered into
457 Area V2 through the use of electronic micromanipulator (Narishige Inc.). Visual inspection of the
458 cortical surface through a surgical microscope (Leica Microsystems) allowed for precise targeting
459 of these probes to desired locations, as well as continuous monitoring of electrode entry. The initial
460 penetration through the AD, arachnoid, and pia was done at a higher speed (>100 μm/s), after
461 which the penetration continued as slow speeds (2 μm/s). Following complete insertion, the probe
462 was retracted slowly (2 μm/s) to relieve pressure without shifting the position of the probe relative
463 to the cortex.

464 Electrical signals from the probe were collected at 30 kHz, digitized on a 64-channel
465 headstage, and send to the recording controller (RHD Recording System, Intan Technologies).
466 Action potential waveforms were extracted offline with Kilosort2[56,57] with default settings
467 (threshold = [10, 4], lambda = 10, AUC for splitting = 0.9) and manually sorted into single- and
468 multi-unit clusters. To quantify the stability of our single unit recordings, we computed kernel
469 density distribution estimates for several metrics[51-53] (Figure S2E). Presence ratio reflects the
470 proportion of a session during which a unit was present and firing action potentials. Maximum
471 drift is the distance between the highest and lowest channels on which a unit was detected during
472 a session. Isolation distance and d-prime quantify the separation of spike waveform clusters in
473 principal component space. Single-unit clusters were further classified into broad- and narrow-
474 spiking units based on their trough-to-peak waveform duration[32,58]. Units with waveform durations
475 less than 350 μs were labelled as narrow-spiking, while units with waveform durations greater

than 350 μs were labelled as broad-spiking. Units with a maximum waveform amplitude preceding the trough were classified as axonal spikes and excluded. Recordings were collected over the course of 17 sessions (8 in monkey M, 9 in monkey D). In total, 923 single units were recorded (461 in monkey M, 462 in monkey D). Only single units with a significant spatial receptive field (89.06%), as determined by a chi-squared test, were considered for subsequent analysis.

Behavioral Control and Eye Tracking

Behavioral experiments were controlled with NIMH Monkeylogic[59] in MATLAB. Eye position and pupil diameter were continuously sampled at 120 Hz (ETL-200, ISCAN Inc.) and sent to the behavioral control system. Stimuli were presented on a monitor (BenQ XL2411; 60 Hz refresh rate) positioned 57 cm from the monkey. Tolerance windows for fixation control were one degree of visual angle.

Receptive Field Mapping

Receptive fields (RFs) were mapped with Gabor patch stimuli (2-4 cycles/deg, 0.5-1.5 deg Gaussian half-width, 100% luminance contrast) on a square grid spanning the visual quadrant of interest (lower right in monkey M, lower left in monkey D) while the subject maintained fixation. Grid spacing parameters were optimized for each session based on receptive field eccentricity and ranged from 0.25 – 1.0 degrees of visual angle (dva). A single Gabor was presented at one of six orientations (0, 30, 60, 90, 120, 150°) and at a grid location, both chosen at random, on each frame of stimulus presentation (60 Hz). Stimulus-evoked local field potential (LFP) power at each grid location on each recording channel was smoothed with a Gaussian kernel, and the peak location was defined as the RF center. Spatial RF maps for each channel were plotted as stacked contours for each shank to aid in visualization.

Current Source Density Mapping

499 Current source density (CSD) mapping[31,32] was used to identify laminar boundaries. While
500 subjects held fixation, 100% contrast annular stimuli were flashed for 32ms, positioned over the
501 center of the RF. The CSD was calculated as the second spatial derivative of the LFP. CSD traces
502 were spatially smoothed with a Gaussian kernel (sigma = 140μm). The input layer was identified
503 by an early current sink, representing feedforward input into layer IV. Channels above and below
504 this sink were classified as superficial and deep respectively.

505 **Cued Saccade Task**

506 During the task, subjects acquired and held fixation for a variable delay period (500-900 ms) prior
507 to initiating a saccade in response to a target point appearing in the periphery. The simultaneous
508 disappearance of the fixation point served as the go cue. After executing an accurate saccade,
509 subjects then had to continue holding fixation at the target point for 500 ms to receive a reward.
510 To prevent subjects from preemptively planning a saccade prior to the go cue, both the saccade
511 target location and the delay period duration were pseudo-randomized. The target location was
512 drawn from one of two possible locations, while the delay period duration was drawn from an
513 exponential distribution. Targets were located 2.8 dva from the initial fixation point. Target
514 locations were orthogonal to one another, and were each oriented 45 degrees to and equidistant
515 from the fixation to receptive field axis. Only eye movements originating from < 0.75 dva of the
516 initial fixation point, and terminating < 0.75 dva from the target were considered successful trials.
517 While the subjects executed these eye movements, oriented Gabor stimuli were continuously
518 presented on a 13 x 13 grid spanning the visual region of interest at 60 Hz. The grid was centered
519 on a point 4 dva from fixation along the fixation-receptive field axis. On each frame of stimulus
520 presentation, a single stimulus drawn from one of 6 random orientations (0, 30, 60, 90, 120, 150
521 degrees) was presented at a single grid location. Saccades were identified from eye-tracker data

522 with a velocity-thresholding algorithm.[60,61] On average, subjects performed 895 trials of the cued
523 saccade task (minimum of 729 trials, maximum of 1029 trials).

524 **Continuous Receptive Field Mapping**

525 Receptive fields were mapped for each single unit as a function of time. Spikes were binned for
526 each unit in response to each stimulus flash in a time window 50 to 100ms after flash onset.
527 Stimulus flashes were then binned (51 ms centered window slid from -400 to 400ms relative to
528 saccade onset) and their corresponding spike counts were averaged. This procedure generated a 13
529 x 13 grid of spike counts at each location for each timepoint relative to saccade onset. Each
530 timepoint was normalized such that the sum of all grid positions was equal to one to control for
531 changes in firing rate. For visualization (Figure 2A only), this spatial grid was smoothed with a
532 Gaussian kernel. The location of the current and future fields was determined by finding the
533 stimulus position that elicited the maximum firing in the pre- and post-saccadic time periods,
534 respectively. To compute sensitivity, the spike counts at the current and future fields were
535 normalized to sum to one at each time point, such that the sensitivity reflects the relative proportion
536 of firing in response to a stimulus presentation at the given field. Sensitivity analyses were done
537 on the unsmoothed, spike count data. To determine the time at which each neural subpopulation
538 first began to show remapping, we computed a bootstrapped 95% interval for the baseline
539 sensitivity (-100 to -50 ms relative to saccade onset) at the future field. The first increase in
540 sensitivity beyond these bounds was marked as the start of remapping for each bootstrapped
541 population mean. Lastly, we determined the proportion of single units that showed clear
542 remapping. Single units were considered to be remapping pre-saccadically if future field sensitivity
543 exceeded (and remained above) current field sensitivity beginning at a timepoint before saccade

544 onset. Only units with firing rates greater than 5 Hz were included in this proportion analysis (175
545 total units).

546 **Principal Component Analysis**

547 A 13 x 13 grid of sensitivity was generated for each timepoint, as detailed above. Each timepoint
548 was then vectorized to produce a 169-dimensional sample for each timepoint. All units from a
549 given session on trials towards one of the two targets were fed into a PCA together, as they had
550 overlapping current and future field locations in that condition. Each sample for a PCA thus reflects
551 the 169-dimensional stimulus response space from one single unit at one timepoint. To average
552 PCA results across sessions, values along the 1st and 2nd principal component axes were range
553 normalized between 0 and 1 for each session. The time at which each subpopulation shows a
554 significant change along the 1st principal component axis was computed with bootstrapping, as
555 described above for the sensitivity analysis.

556 **Tuning**

557 To compute the tuning curves for each unit, spikes were binned in response to stimulus flashes of
558 a given orientation at all positions within one of three epochs: pre-saccadic (-250 to -200 ms before
559 saccade onset), saccade planning (-75 to -25 ms before saccade onset), and post-saccadic (150 to
560 200 ms after saccade onset). An orientation selectivity index and circular variance were calculated
561 for each unit in each of the three epochs[62]. Each unit was fit with a Gaussian plus constant model
562 function using the gaussfitn toolbox in MATLAB. Changes in orientation selectivity index,
563 circular variance, firing rate, and half width at half height were quantified with the estimation stats
564 toolbox[63]. Only units with a session-wide firing rate greater than 1 Hz were included in the tuning
565 analysis.

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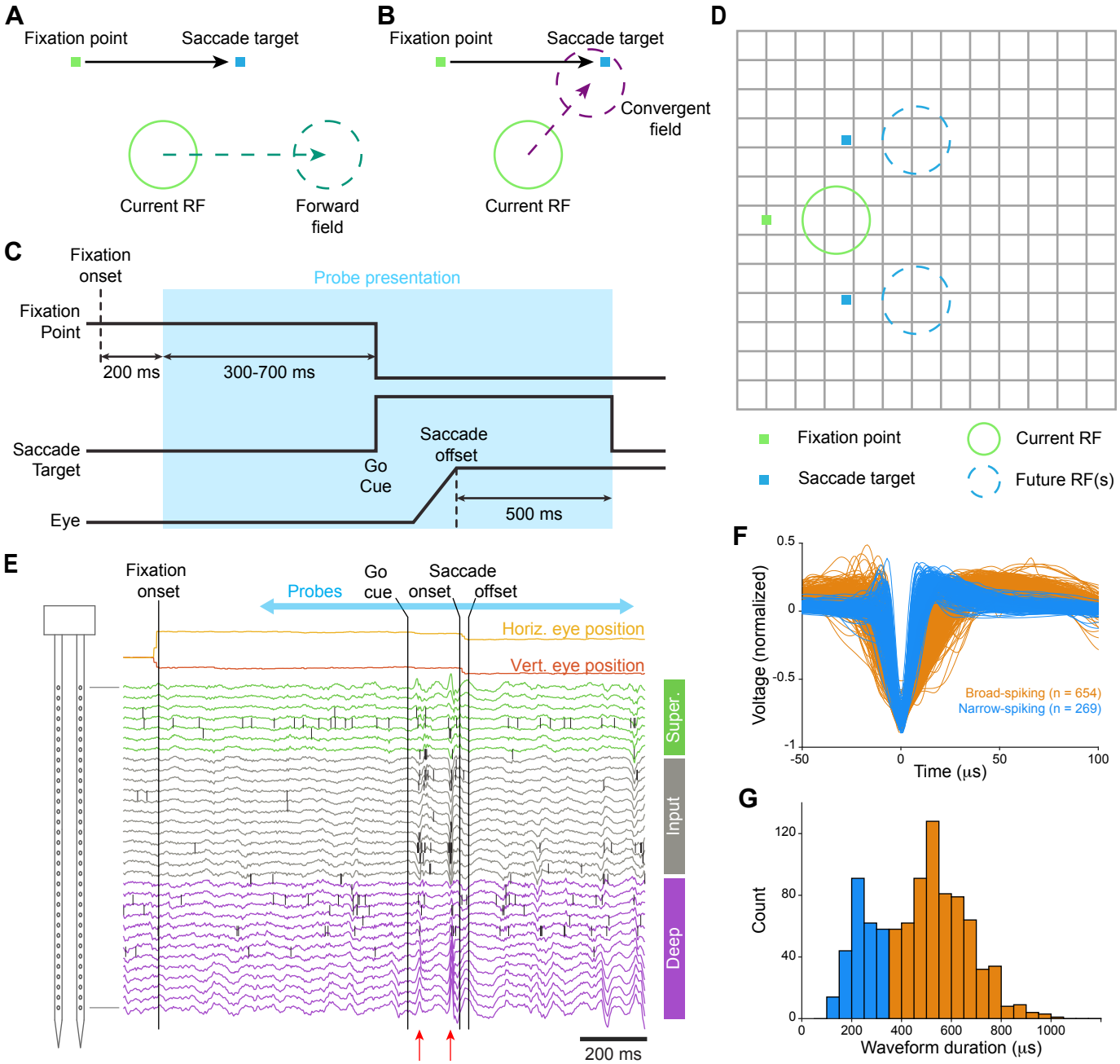

Figure 1. Task Design and Single Unit Recordings

(A) Forward remapping shifts the current receptive field by the vector of the upcoming saccade to form a forward field.

(B) Convergent remapping shifts the current receptive field towards the saccade target to form a convergent field.

(C) Progression of a trial during the cued saccade task as the subject holds fixation, executes a saccade to the target, and then fixates on the target. The period in blue indicates probe presentation. A reward is delivered after fixating on the target for 500 ms.

(D) Fixation point, saccade targets, and stimulus grid layout during the cued saccade task.

(E) Snippet of data from one probe shank during a trial of the cued saccade task. LFP traces are color by cortical layer, and spikes are overlaid on their channel of origin as vertical lines. Red arrows indicate synchronized spiking and local field potential deflections along the depth of the cortex in response to a stimulus being flashed in the receptive field of the recording site.

(F) Action potential waveforms of all 923 single units. Units were classified as either narrow-spiking (blue) or broad-spiking (orange) on the basis of their peak-to-trough waveform duration.

(G) Distribution of waveform durations for single units from (F). The distribution shows bimodality for the two unit types (Hartigan's dip test; $p = 1.15*10^{-4}$).

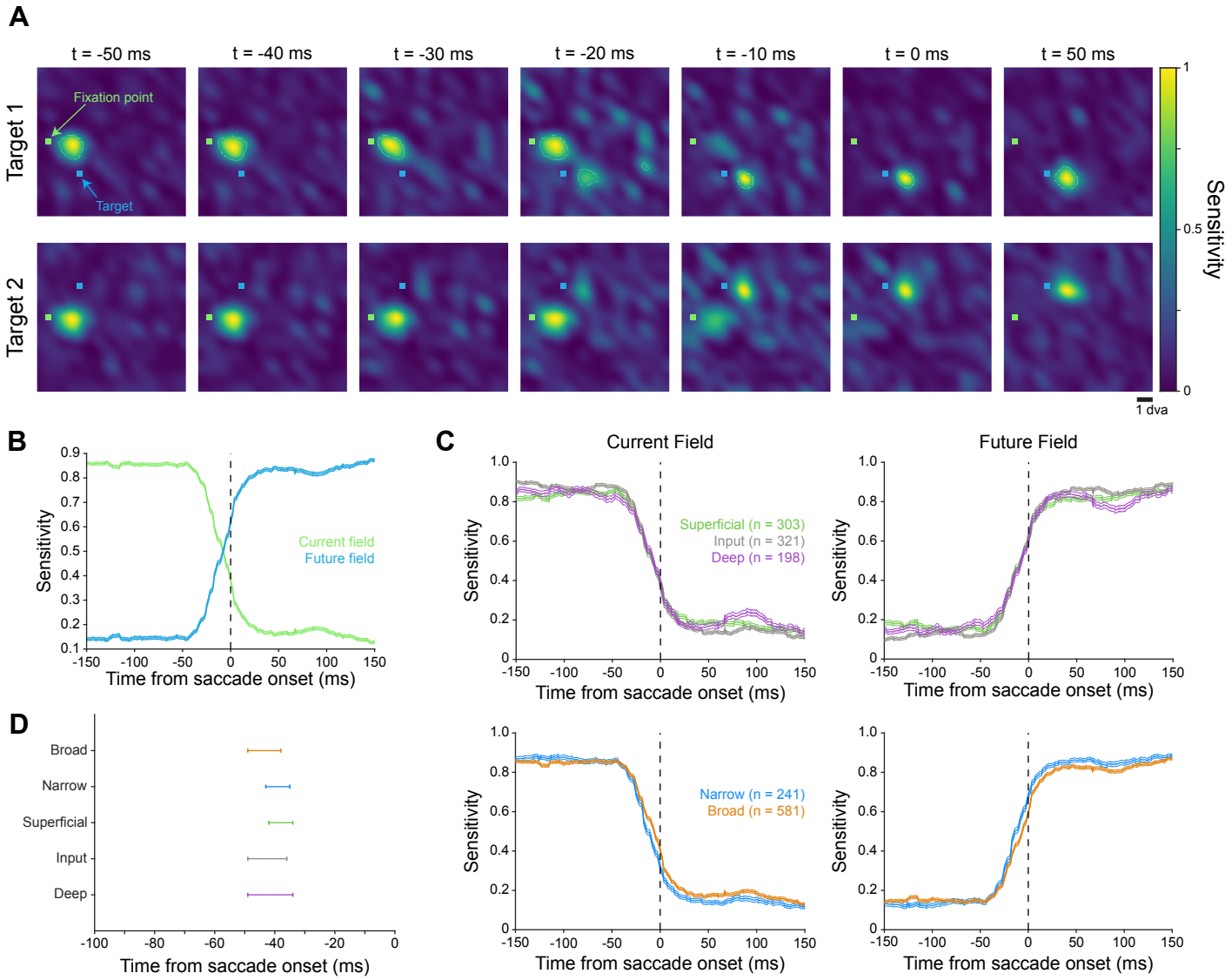

Figure 2. Receptive Field Remapping is Widespread in Area V2

(A) Receptive field location for an example single unit during remapping. Times are relative to saccade onset. Each row shows remapping during saccades to one of the two saccade targets. Fixation point and saccade target are overlaid in green and blue, respectively. Heatmaps at each timepoint are individually normalized to account for possible changes in firing rate.

(B) Normalized sensitivity at the current and future field locations for all single units (n = 822), as a function of time relative to saccade onset. Error bars indicate standard error of the mean.

(C) Normalized sensitivity at the current and future field locations as a function of time relative to saccade onset for each of the recorded neural subpopulations. Error bars indicate standard error of the mean.

(D) The time relative to saccade onset at which remapping initiates for each of the recorded neural subpopulations. Error bars indicate bootstrapped 95% confidence intervals.

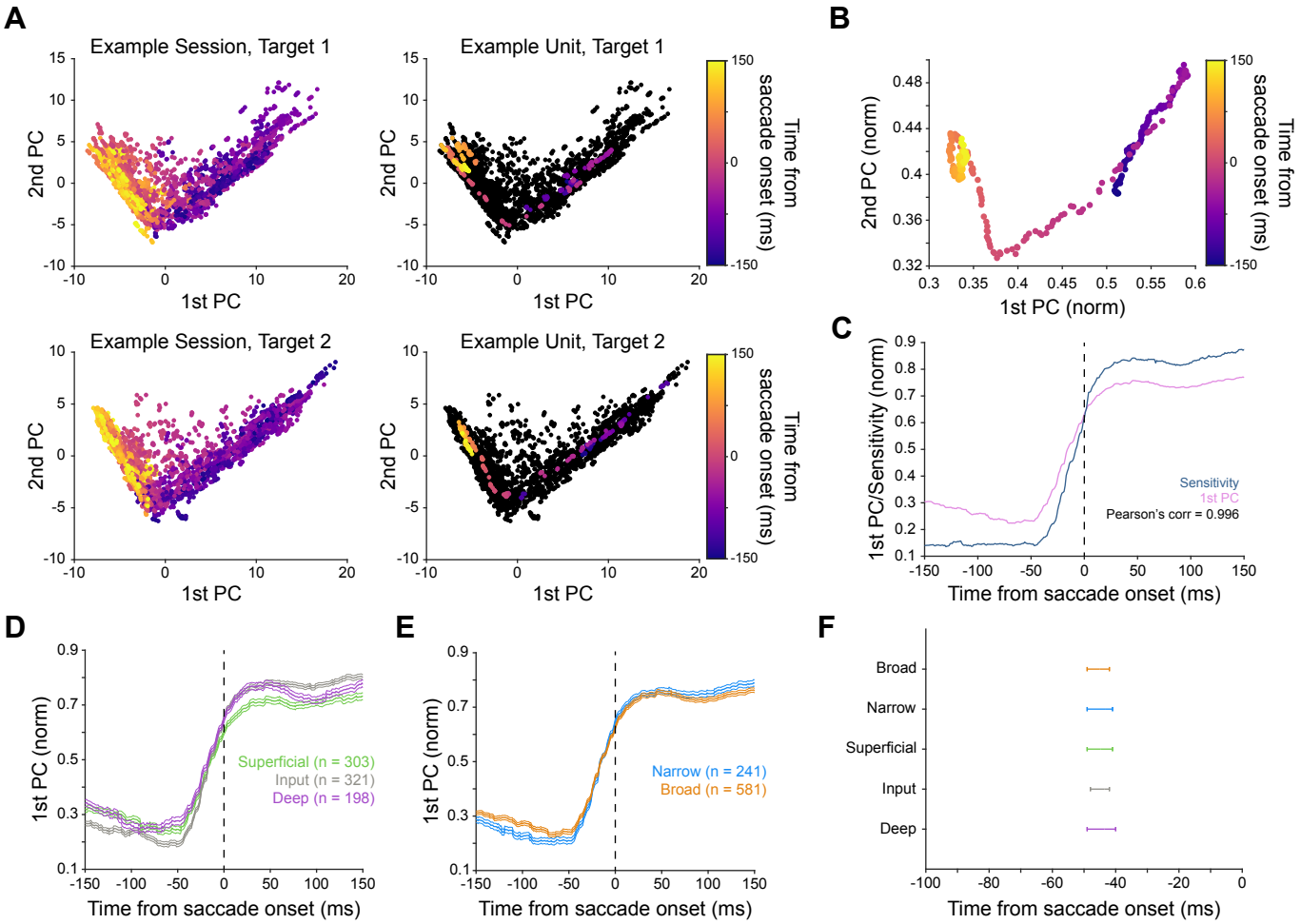

Figure 3. Tracking Remapping Trajectories in Principal Component Space

(A) Left, principle component trajectories of all single units from an example session. Right, principle component trajectory of an example single unit from the same session. Top, target 1. Bottom, target 2. Each point represents a single timepoint from a single unit. On average across sessions and targets, PC1 explains 11.3% of the variance, and PC2 explains 3.3% of the variance.

(B) Averaged remapping trajectory in principal component space across all single units for both targets.

(C) Correlation between values along the 1st principle component axis and sensitivity at the future field (Figure 2B). Pearson's correlation = 0.996.

(D) Time course of values along the 1st principle component axis for all recorded layers. Error bars indicate standard error of the mean.

(E) Time course of values along the 1st principle component axis for both recorded unit types. Error bars indicate standard error of the mean.

(F) The time relative to saccade onset at which remapping initiates for each of the recorded neural subpopulations based on the 1st PC timecourses. Error bars indicate bootstrapped 95% confidence intervals.

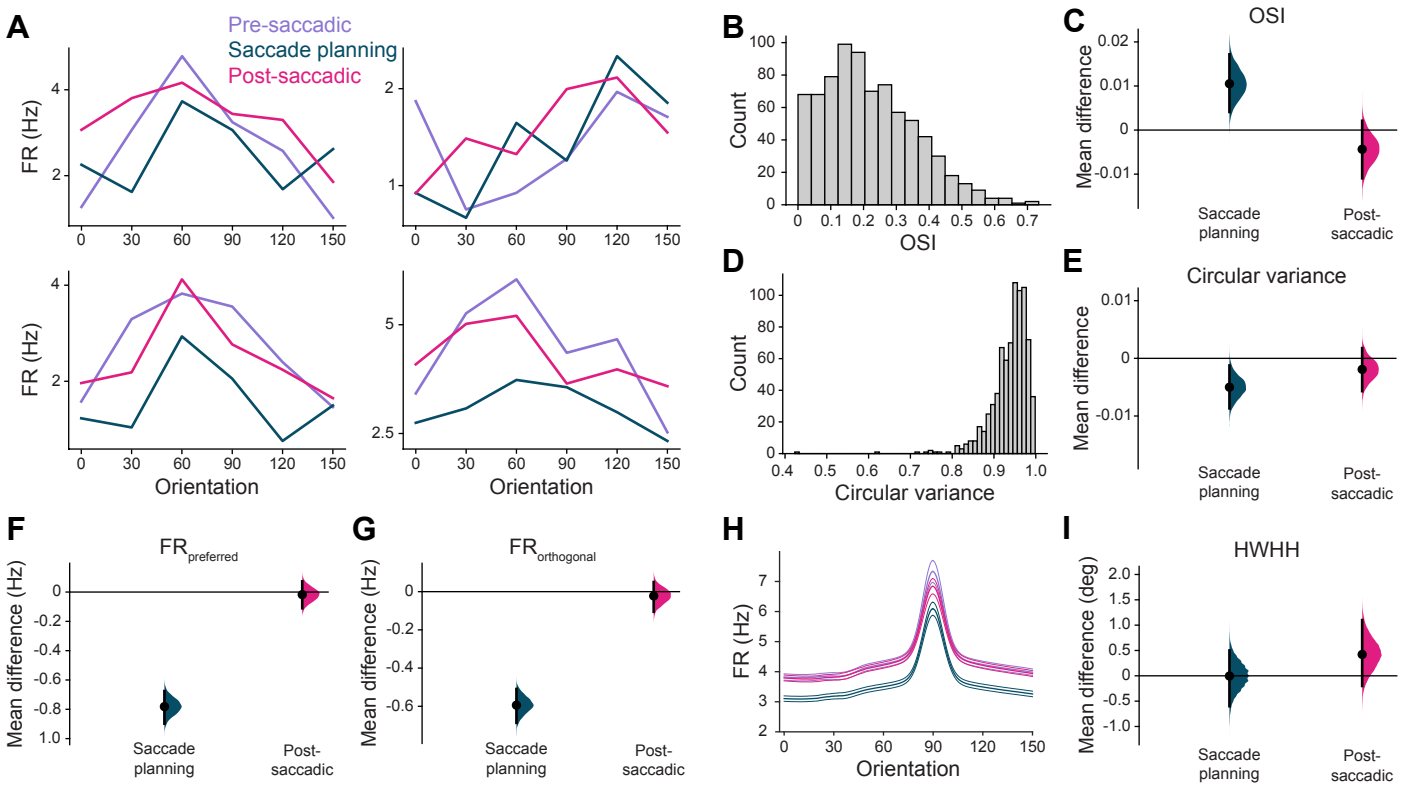

Figure 4. Orientation Selectivity is Transiently Increased During Saccade Planning

(A) Orientation tuning curves from four example single units during the pre-saccadic (-250 to -200 ms), saccade planning (-75 to -25 ms), and post-saccadic (150 to 200 ms) time periods. Times are relative to saccade onset. FR = firing rate.

(B) Distribution of orientation selectivity index (OSI) for all single units during the pre-saccadic period.

(C) Change in OSI during the saccade planning and post-saccadic periods, as compared to the pre-saccadic period. Error bars indicate bootstrapped 95% confidence intervals.

(D) Distribution of circular variance for all single units during the pre-saccadic period.

(E) Change in circular variance during the saccade planning and post-saccadic periods, as compared to the pre-saccadic period. Error bars indicate bootstrapped 95% confidence intervals.

(F) Change in firing rate in response to presentation of a stimulus at the preferred orientation during the saccade planning and post-saccadic periods, as compared to the pre-saccadic period. Error bars indicate bootstrapped 95% confidence intervals.

(G) Change in firing rate in response to presentation of a stimulus at the non-preferred (orthogonal) orientation during the saccade planning and post-saccadic periods, as compared to the pre-saccadic period. Error bars indicate bootstrapped 95% confidence intervals.

(H) Average of Gaussian tuning curve fits across all single units during the pre-saccadic, saccade planning, and post-saccadic time periods. Error bars indicate standard error of the mean.

(I) Change in half width at half height (HWHH) of the fitted tunning curves during the saccade planning and post-saccadic periods, as compared to the pre-saccadic period. Error bars indicate bootstrapped 95% confidence intervals.

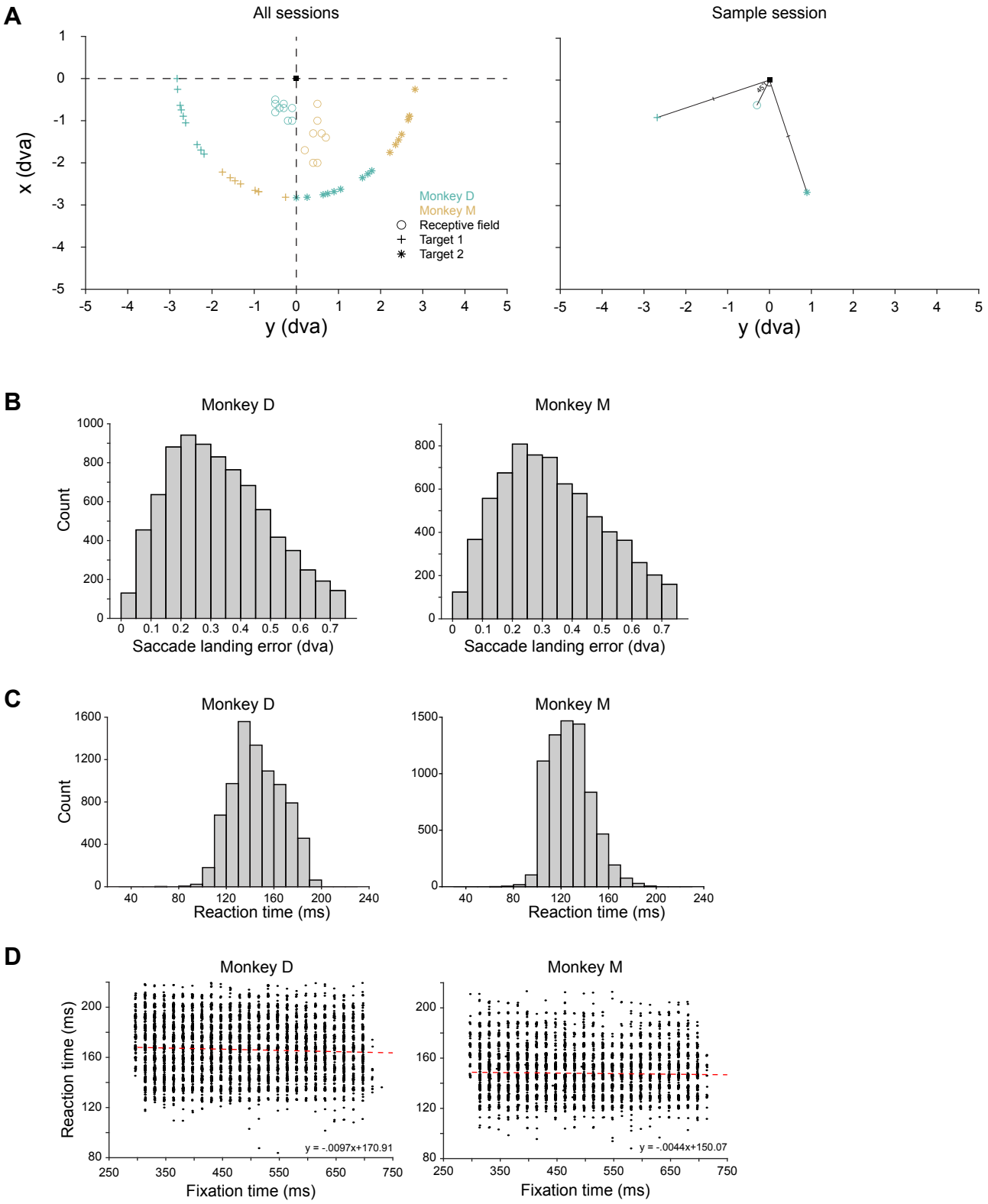

Figure S1. Behavioral Performance

(A) Receptive field and saccade target locations for all sessions (left) and one sample session (right).

(B) Distribution of saccade landing errors for both subjects. Landing error was defined as the distance between the saccade target and the terminal point of the saccade.

(C) Distribution of reaction times for both subjects.

(D) Relationship between the trial-by-trial pre-saccadic fixation time (pseudorandomly determined for each trial) and reaction time. The dashed red line depicts a linear least-squares fit. The flat relationship between fixation time and reaction time suggests that subjects are not able to anticipate the timing of the go cue.

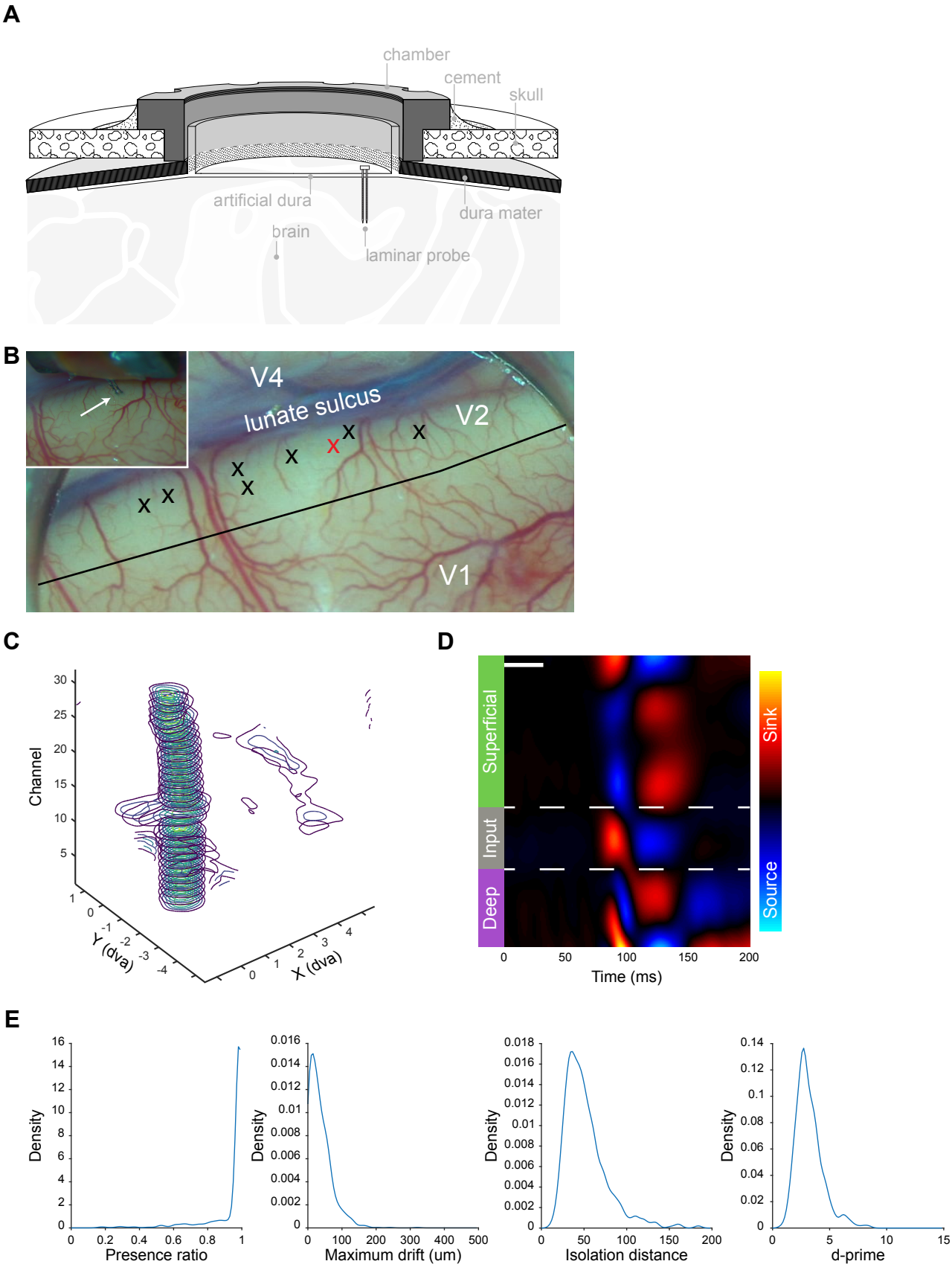
A
chamber
cement
skull
artificial dura
dura mater
brain
laminar probe
B
V4
lunate sulcus
V2
V1
C
Channel
Y (dva)
X (dva)
D
Superficial
Input
Deep
Sink
Source
Time (ms)
E
Density
Presence ratio
Maximum drift (um)
Isolation distance
d-prime

Figure S2. Recording Methodology

(A) Schematic of the artifical dura and recording chamber (see Methods).

(B) Microscope image inside the recording chamber of one subject. Each ‘X’ indicates a cortical recording site. The inset shows a close-up image of the linear array probe penetration at the site marked in red. The black line is the estimated boundary between Area V1 and Area V2 based on the distinct change in surface vasculature density between the two areas.

(C) Receptive fields along the depth of one shank as computed from local field potential deflections. dva = degrees of visual angle

(D) Current source density (CSD) analysis of the shank from (C). Blue indicates a current source, while red indicates a current sink. Dashed lines indicates laminar boundaries, as determined from the CSD. White bar indicates duration of stimulus presentation.

(E) Kernel density distribution estimates for several metrics29-31 that quantify recording stability and single unit quality. Presence ratio reflects the proportion of a session during which a unit was present and firing action potentials. Maximum drift is the distance between the highest and lowest channels on which a unit was detected during a session. Isolation distance and d-prime quantify the separation of spike waveform clusters in principal component space.

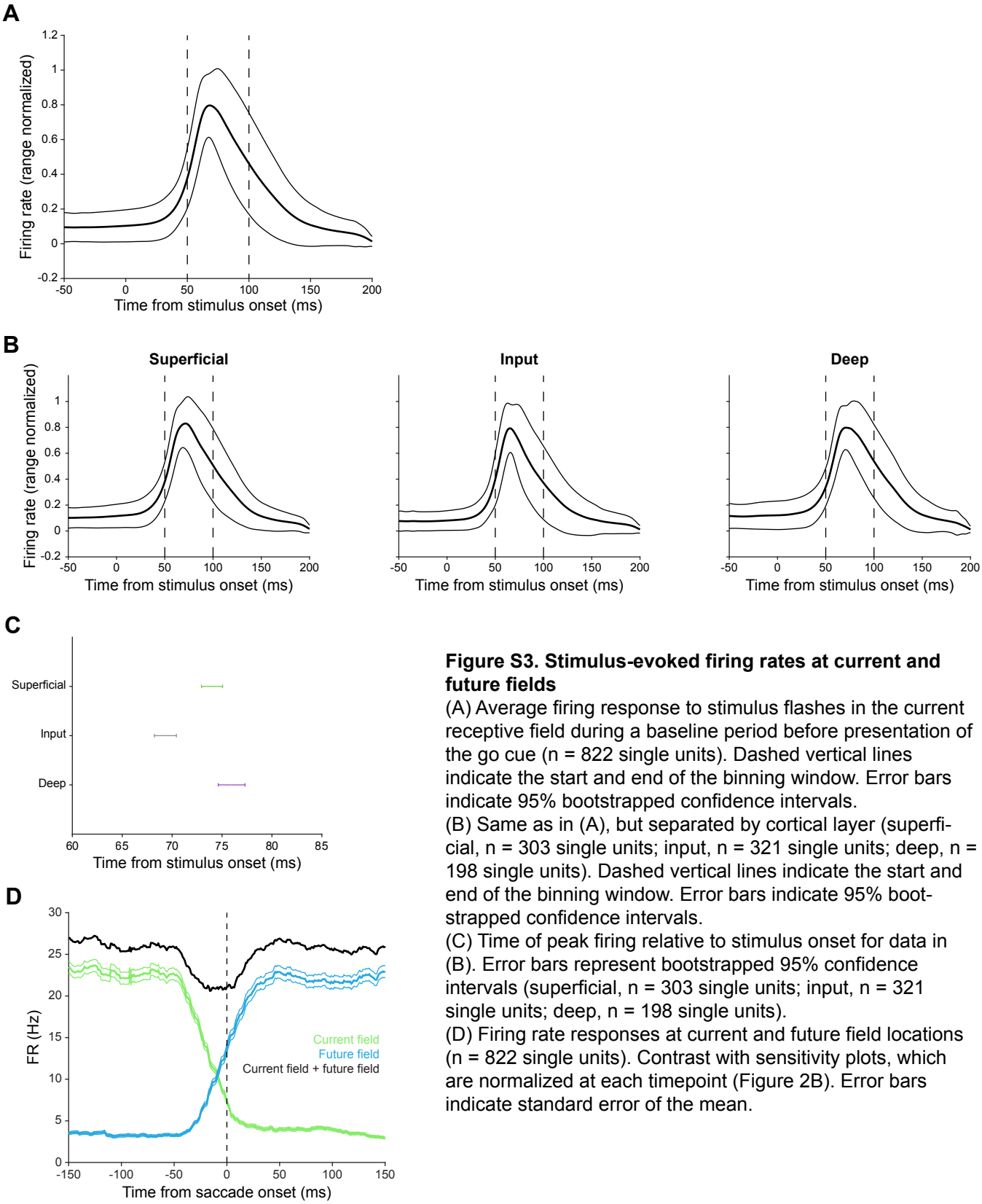

Figure S3. Stimulus-evoked firing rates at current and future fields

(A) Average firing response to stimulus flashes in the current receptive field during a baseline period before presentation of the go cue (n = 822 single units). Dashed vertical lines indicate the start and end of the binning window. Error bars indicate 95% bootstrapped confidence intervals.

(B) Same as in (A), but separated by cortical layer (superficial, n = 303 single units; input, n = 321 single units; deep, n = 198 single units). Dashed vertical lines indicate the start and end of the binning window. Error bars indicate 95% bootstrapped confidence intervals.

(C) Time of peak firing relative to stimulus onset for data in (B). Error bars represent bootstrapped 95% confidence intervals (superficial, n = 303 single units; input, n = 321 single units; deep, n = 198 single units).

(D) Firing rate responses at current and future field locations (n = 822 single units). Contrast with sensitivity plots, which are normalized at each timepoint (Figure 2B). Error bars indicate standard error of the mean.

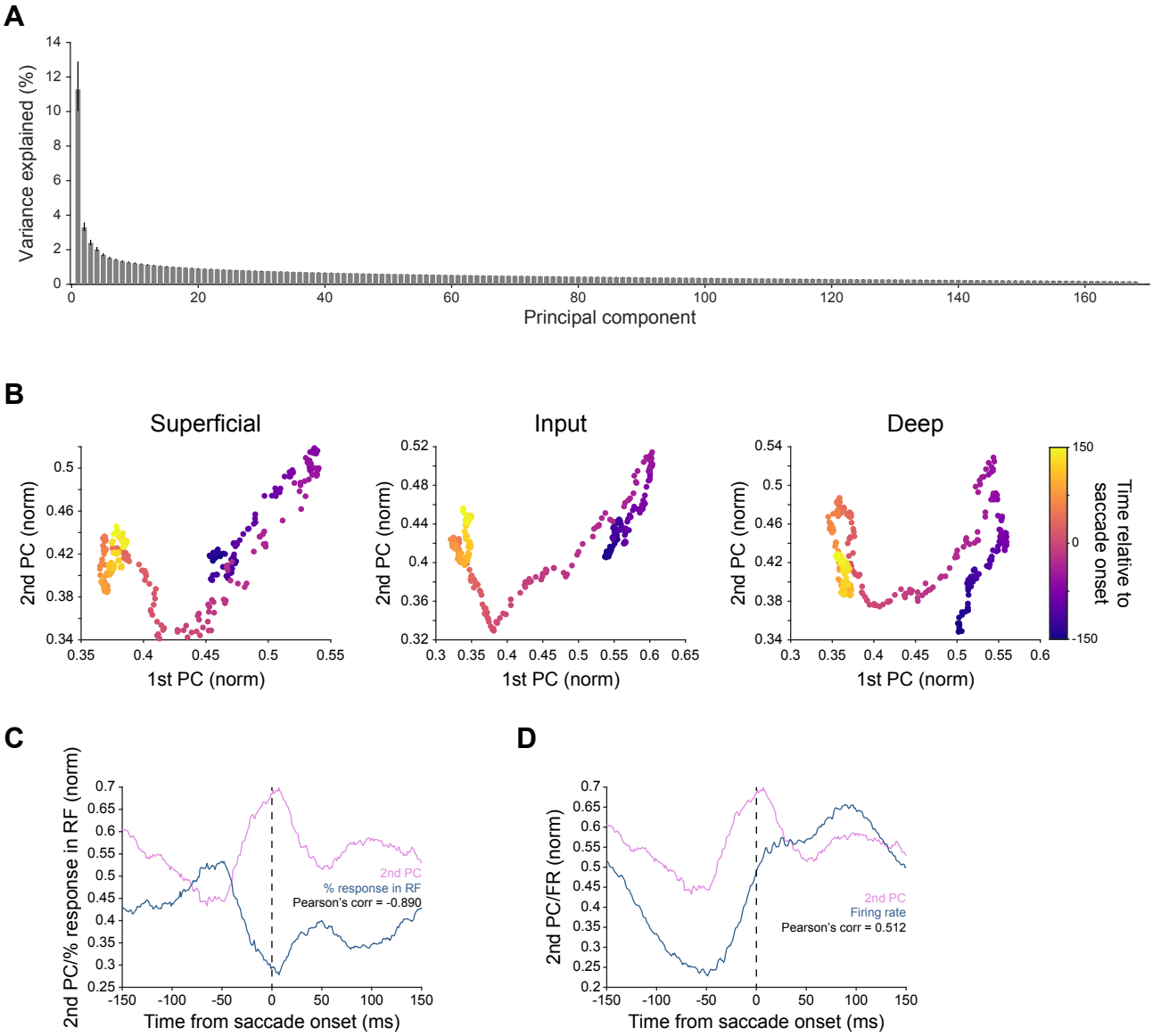

Figure S4. Remapping Trajectories are Tracked in Principal Component Space
(A) Variance explained by each principal component, averaged across sessions and targets. Error bars indicate bootstrapped 95% confidence intervals.
(B) Averaged remapping trajectory in principal component space across all single units for both targets when PC analysis is performed on units from a given layer.
(C) Correlation between values along the 2nd principal component axis and the percent of total response contained within the current and future fields (as opposed to firing evoked by stimuli landing outside the receptive field). Pearson's correlation = -0.890.
(D) Correlation between values along the 2nd principal component axis and firing rate. Pearson's correlation = 0.512.

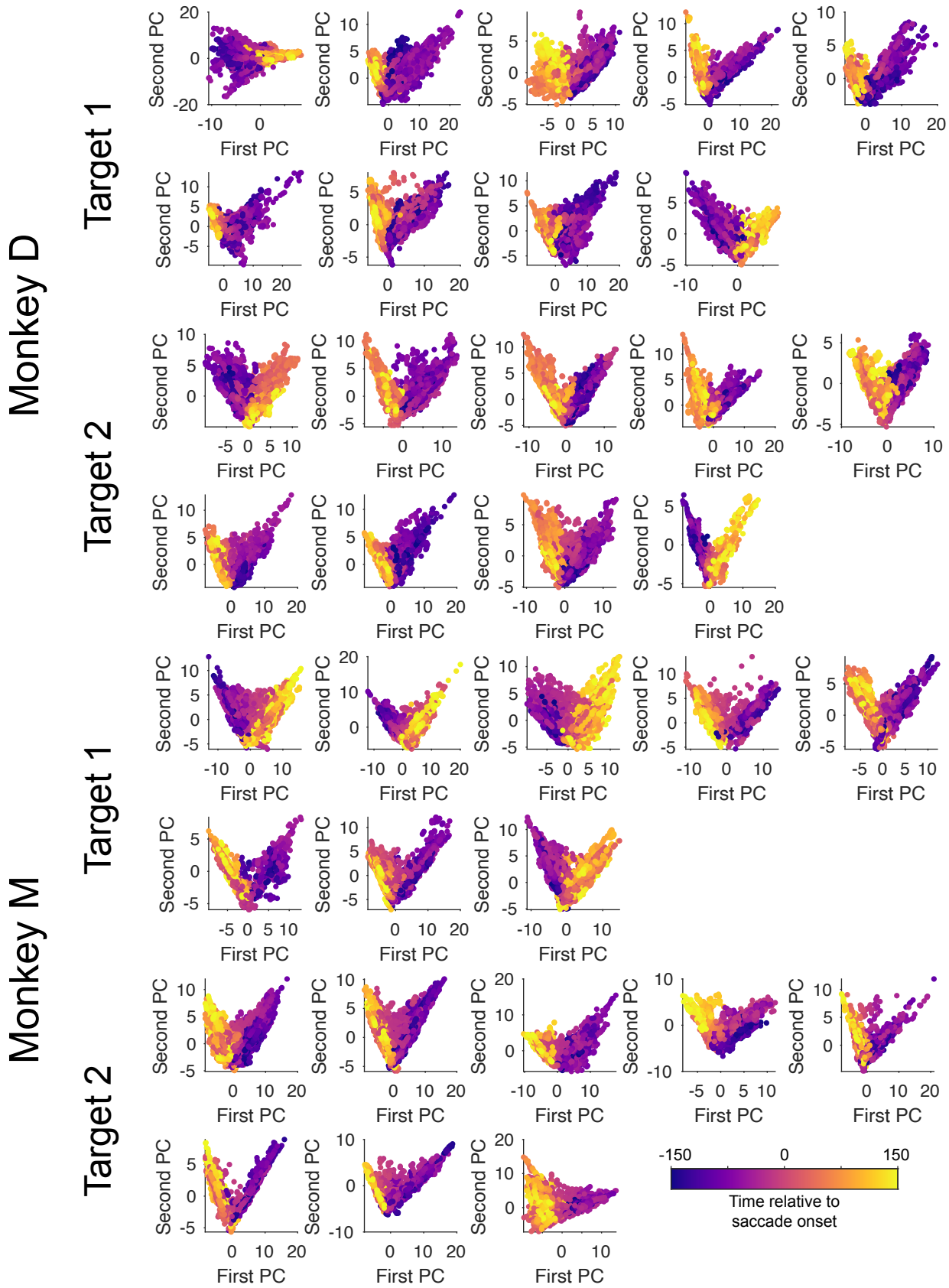

Figure S5. Principle Component Trajectories of All Sessions
Receptive field remapping trajectories in principal component space of all sessions. Across sessions, the data consistently shows a V-shaped trajectory in this space as remapping occurs.

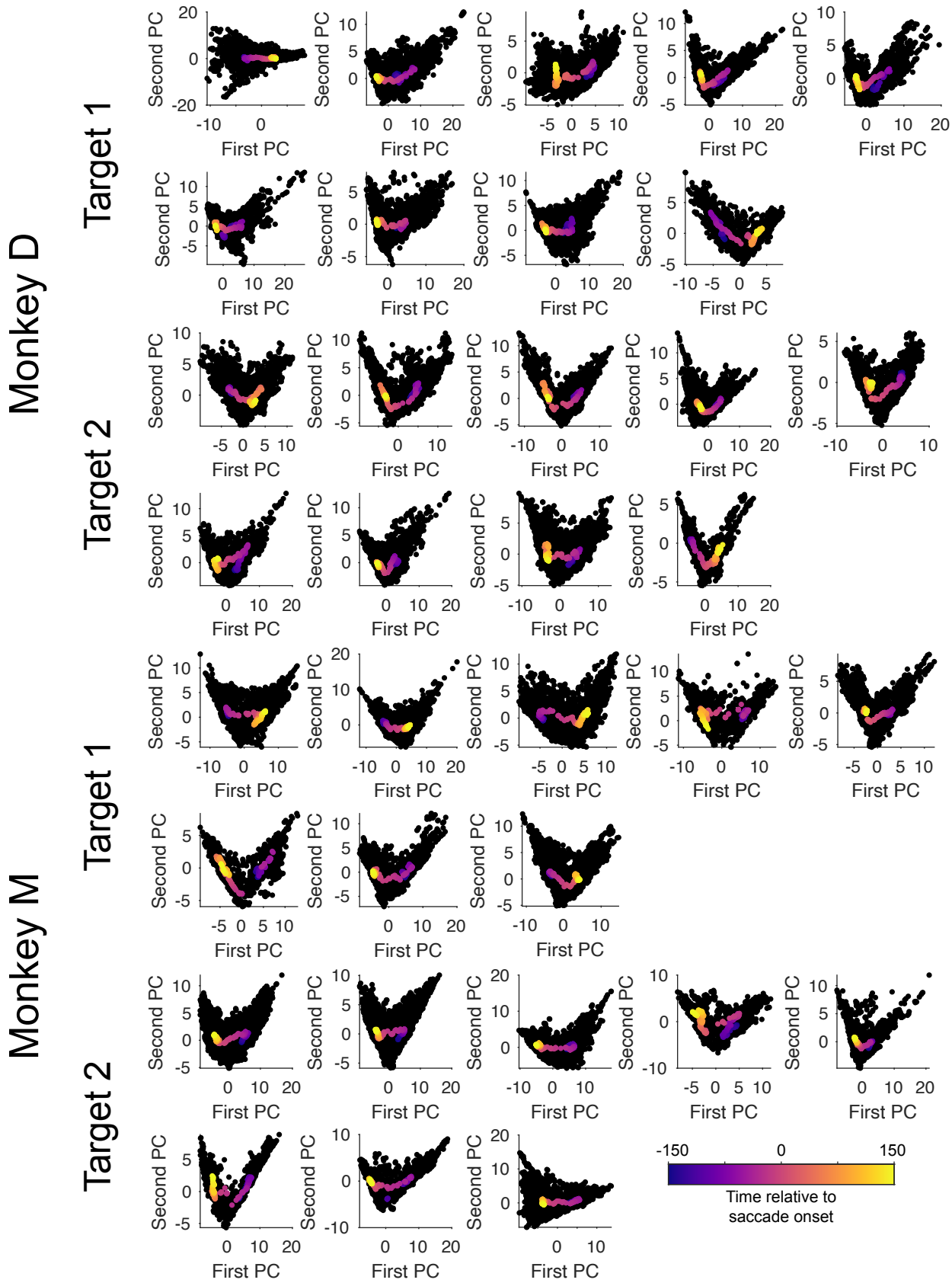

Figure S6. Average Principle Component Trajectories of All Sessions
Average receptive field remapping trajectories in principal component space of all sessions.