

Highlights

- Neural trajectories in the hippocampus exhibited greater variability during a working memory (WM) task compared to those in the entorhinal cortex and amygdala regions.
- The distance of neural trajectories between encoding and retrieval states in the hippocampus was memory-load dependent during a WM task.
- Hippocampal neural trajectories fluctuated between the encoding and retrieval states in a task-dependent manner during both baseline and sharp-wave ripple (SWR) periods.
- Hippocampal neural trajectories shifted from encoding to retrieval states during SWR period.

Hippocampal neural fluctuations during encoding and retrieval in a human working memory task: Encoding-to-retrieval shift during sharp-wave ripples

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Abstract

Working memory (WM) is indispensable for numerous cognitive functions, although the underlying neural mechanisms have not been fully elucidated. While the hippocampus and sharp-wave ripple complexes (SWRs) – brief, synchronous neural events within the hippocampus – are acknowledged for their importance in memory consolidation and retrieval, their association with WM tasks remains unclear. The present study posits that hippocampal multiunit activity patterns, in concert with SWRs, manifest distinct dynamics during WM tasks. We analyzed a dataset comprising intracranial electroencephalogram recordings from the medial temporal lobe (MTL) of nine epilepsy patients engaged in an eight-second Sternberg task. Gaussian-process factor analysis was utilized to extract low-dimensional neural representations, or ‘trajectories’, within MTL regions during the WM task. Our results reveal that the hippocampus demonstrates the most significant fluctuation in neural trajectory relative to the entorhinal cortex and amygdala. Furthermore, trajectories’ dissimilarity measured between encoding and retrieval phases is dependent on memory load. Notably, hippocampal trajectories fluctuate during the retrieval phase and exhibit task-dependent shifts between encoding and retrieval states, both during baseline and SWR events. This fluctuation transitions from encoding to retrieval states in the presence of SWRs. These findings reinforce the pivotal role of the hippocampus in WM tasks and suggest a new hypothesis: the hippocampus alters its functional state from encoding to retrieval during SWRs.

Keywords: working memory, WM, memory load, hippocampus, sharp-wave ripples, SWR, humans

1. Introduction

Working memory (WM) is crucial in everyday life; however, its neural mechanism has yet to be fully elucidated. Specifically, the role of the hippocampus, an essential brain region contributing to memory has been an ongoing topic [1] [2] [3] [4] [5] [6] [7] [8] [9]. Understanding the hippocampus’ role in working memory is instrumental in deepening our knowledge of cognitive processes, ultimately aiding in developing cognitive training strategies and interventions.

It is known that a transient and synchronous oscillation

called sharp-wave ripple (SWR) [10] is associated with various cognitive functions, including memory replay [11] [12] [13] [14] [15], memory consolidation [16] [17] [18] [19], memory recall [20] [21] [22], and neural plasticity [23] [24]. Thus, SWR might be a fundamental representation of processing in the hippocampus and contribute to working memory performances. However, investigations into the effects of SWRs on working memory remain infrequent ([25] and limited to rodent models using navigation tasks, in which precise timings of memory acquisition and recall are not separated.

Moreover, it is getting discovered that hippocampal neurons exhibit low-dimensional representations during

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WM tasks. For instance, the firing patterns of place cells [26] [27] [28] [29] [30] in the hippocampus were embedded within a dynamic, nonlinear three-dimensional hyperbolic geometry in rodent [31]. Furthermore, grid cells in the entorhinal cortex (EC) — the primary gateway to the hippocampus [32] [33] [34] — exhibited toroidal topology during exploration [35]. However, again, these experiments are limited to spatial navigation tasks in rodents so that the temporal resolution of WM tasks is constrained. Moreover, whether these findings are generalized to humans and beyond navigation tasks, are not investigated yet.

Given these backgrounds, in this study, we investigated the hypothesis that hippocampal neurons exhibit distinct representations in low-dimensional spaces as ‘neural trajectory’ during WM tasks, with a specific focus on SWR periods. To test this hypothesis, we utilized a dataset of patients performing an eight-second Sternberg task with high temporal resolution (1 s for fixation, 2 s for encoding, 3 s for maintenance, and 2 s for retrieval) while their intracranial electroencephalography signals (iEEG) in the medial temporal lobe (MTL) were recorded [36]. To explore low-dimensional neural trajectories, we employed Gaussian-process factor analysis (GPFA) based on multiunit activities, a proven tool for the analysis of neural population dynamics [37].

2. Results

2.1. iEEG recording and neural trajectory in MTL regions during a Sternberg task

We employed a publicly available dataset [36] for this analysis. This dataset includes LFP signals (Figure 1A) within MTL regions (Table 1) during a modified Sternberg task. SWR⁺ candidates were detected from LFP signals passed through the ripple band (80–140 Hz) (Figure 1B) within all hippocampal regions (refer to Methods), while SWR[−] candidates were defined at identical timestamps of SWR⁺ candidates but with shuffled across different trials (Figure 1). The multiunit spikes (Figure 1C) are included in the dataset as well, being established using a spike sorting algorithm [38]. Using the 50-ms binned multiunit activity without overlaps, we employed GPFA [37] to determine the neural trajectory (or factors) of the MTL regions by session and region (Figure 1D). Each factor was z-normalized by session and region (for example, session #2 in AHL

of subject #1). The Euclidean distance from the origin (*O*) was calculated (Figure 1E).

2.2. Hippocampal neural trajectory correlated with a Sternberg task

In Figure 2A, the median neural trajectories of 50 trials were depicted as point clouds within the three major factor space. The optimal embedding dimension for the GPFA model was determined to be three using the elbow method (Figure 2B). The trajectory distance from the origin (*O*) ($\|g_F\|$, $\|g_E\|$, $\|g_M\|$, and $\|g_R\|$) of the hippocampus was larger than those of the EC and amygdala (Figure C & D).¹

Similarly, the distance among geometric medians of the four phases were calculated: $\|g_F g_E\|$, $\|g_F g_M\|$, $\|g_F g_R\|$, $\|g_E g_M\|$, $\|g_E g_R\|$, and $\|g_M g_R\|$. Again, the hippocampus showed larger distances among phases compared to both the EC and amygdala.²

2.3. Memory load-dependent neural trajectory distance between the encoding and retrieval states in the hippocampus

Regarding the memory load of the Sternberg task, correct rate of trials and set size (= the number of alphabetical letters to encode) were negatively correlated (Figure 3A).³ Similarly, response time and set size were positively correlated (Figure 3B).⁴

Furthermore, similarly, set size and the trajectory distance between the encoding and retrieval phases

¹Hippocampus: Distance = 1.11 [1.01], median [IQR], $n = 195,681$ timepoints; EC: Distance = 0.94 [1.10], median [IQR], $n = 133,761$ timepoints; Amygdala: Distance = 0.78 [0.88], median [IQR], $n = 165,281$ timepoints.

²Hippocampus: Distance = 0.60 [0.70], median [IQR], $n = 8,772$ combinations; EC: Distance = 0.28 [0.52], median [IQR], $n = 5,017$ combinations ($p < 0.01$; Brunner–Munzel test); Amygdala: Distance = 0.24 [0.42], median [IQR], $n = 7,466$ combinations ($p < 0.01$; Brunner–Munzel test).

³Correct rate: set size four (0.99 ± 0.11 , mean \pm SD; $n = 333$ trials) vs. set size six (0.93 ± 0.26 ; $n = 278$ trials; $p < 0.001$, Brunner–Munzel test with Bonferroni correction) and set size eight (0.87 ± 0.34 ; $n = 275$ trials; $p < 0.05$; Brunner–Munzel test with Bonferroni correction). Overall, $p < 0.001$ for Kruskal–Wallis test; correlation coefficient = -0.20 , $p < 0.001$.

⁴Response time: set size four (1.26 ± 0.45 s; $n = 333$ trials) vs. set size six (1.53 ± 0.91 s; $n = 278$ trials) and set size eight (1.66 ± 0.80 s; $n = 275$ trials). All comparisons $p < 0.001$, Brunner–Munzel test with Bonferroni correction; $p < 0.001$ for Kruskal–Wallis test; correlation coefficient = 0.22 , $p < 0.001$.

($\log_{10}||g_{EGR}||$) were positively correlated (Figure 3C).⁵, while distances between other phase combinations did not yield no significant correlations (Figures 3D & S2).

2.4. Detection of hippocampal SWR from putative CA1 regions

Under the aim to improve the precision of recording sites and the detection of SWRs, we estimated electrodes in CA1 regions of the hippocampus based on observing distinct multiunit spike patterns during SWR events. For each session and hippocampal region, SWR^+/SWR^- candidates were embedded into a two-dimensional space via UMAP (Figure 4A).⁶ We calculated the silhouette score as a measure of clustering quality (Figure 4B & Table 2). Recording sites with an average silhouette score across sessions exceeding 0.6 were defined as putative CA1 regions⁷ (Tables 2 & 3). Five putative CA1 regions were identified, and the four of them had not been labeled as seizure onset zones (Table 1).

Subsequently, SWR^+/SWR^- candidates within these putative CA1 regions were labeled SWR^+ and SWR^- , respectively⁸ (Table 3). Both SWR^+ and SWR^- exhibited an identical duration⁹ (Figure 4C) due to their definitions, following a log-distribution profile. A increase in SWR^+ incidence was detected during the initial 400 ms of the retrieval phase¹⁰ (Figure 4D). Additionally, the peak ripple band amplitude of SWR^+ exceeded that of SWR^- and followed a log-normal distribution (Figure 4E).¹¹.

⁵Correlation between set size and $\log_{10}||g_{EGR}||$: correlation coefficient = 0.05, $p < 0.001$. Specific values: $||g_{EGR}|| = 0.54$ [0.70] for set size four trials, $n = 447$; $||g_{EGR}|| = 0.58$ [0.66] for set size six trials, $n = 381$; $||g_{EGR}|| = 0.61$ [0.63] for set size eight trials, $n = 395$.

⁶For illustrative purposes, consider the AHL in session #1 of subject #1.

⁷The identified regions were: AHL of subject #1, AHR of subject #3, PHL of subject #4, AHL of subject #6, and AHR of subject #9.

⁸Definitions lead to equal counts for both categories: SWR^+ ($n = 1,170$) and SWR^- ($n = 1,170$).

⁹Definitions lead to equal duration for both categories: SWR^+ (93.0 [65.4] ms) and SWR^- (93.0 [65.4] ms).

¹⁰ SWR^+ increased against the bootstrap sample; 95th percentile = 0.42 [Hz]; $p < 0.05$.

¹¹ SWR^+ (3.05 [0.85] SD of baseline, median [IQR]; $n = 1,170$) vs. SWR^- (2.37 [0.33] SD of baseline, median [IQR]; $n = 1,170$; $p < 0.001$; Brunner–Munzel test).

2.5. Transient neural trajectory change in the hippocampus during SWR

The *distances* of trajectory from the origin (O) during SWR events in both the encoding and retrieval phases were calculated (Figure 5A). Given the distance increase during SWR as shown in Figure 5A, we categorized each SWR into three stages: pre-, mid-, and post-SWR. Subsequently, the distances from O during these SWR periods are represented as $||pre-eSWR^+||$, $||mid-eSWR^+||$, and so on.

$||mid-eSWR^+||$ ¹² was larger than $||pre-eSWR^+||$ ¹³, and $||mid-rSWR^+||$ ¹⁴ was larger than $||pre-rSWR^+||$ in both Match IN and Mismatch OUT tasks.¹⁵

2.6. Visualization of hippocampal neural trajectory during SWR in two-dimensional spaces

Based on our observations of neural trajectory 'jump' during SWR (Figure 5), we visualized the three-dimensional trajectories of pre-, mid-, and post-SWR events during the encoding and retrieval phases (Figure 6), the distance between which was memory-load dependent (Figure 3).

To achieve the visualization in two dimension spaces, peri-SWR trajectories were linearly aligned by positioning g_E at the origin (0, 0) and g_R at ($||g_{EGR}||$, 0). These aligned trajectories were rotated around the g_{EGR} axis (= x-axis). Thus, distances from the origin O and angles between g_{EGR} in the original three-dimensional spaces are preserved in these two-dimensional ones.

The scatter plot in these two-dimensional spaces illustrates characteristic distributions of peri-SWR trajectories based on phases and task types. For instance, it is observable that $||mid - eSWR^+||$ is larger than $||pre - eSWR^+||$ (Figure 6B), consistent with our earlier findings (Figure 5).

2.7. Fluctuations of hippocampal neural trajectories between encoding and retrieval states

Subsequently, we checked trajectory *directions* based on $\overrightarrow{g_{EGR}}$. SWR directions were defined by neural tra-

¹²1.25 [1.30], median [IQR], $n = 1,281$, in Match IN task; 1.12 [1.35], median [IQR], $n = 1,163$, in Mismatch OUT task

¹³1.08 [1.07], median [IQR], $n = 1,149$, in Match IN task; 0.90 [1.12], median [IQR], $n = 1,088$, in Mismatch OUT task

¹⁴1.32 [1.24], median [IQR], $n = 935$, in Match IN task; 1.15 [1.26], median [IQR], $n = 891$, in Mismatch OUT task

¹⁵1.19 [0.96], median [IQR], $n = 673$, in Match IN task; 0.94 [0.88], median [IQR], $n = 664$, in Mismatch OUT task

jectory at -250 ms and $+250$ ms from their center (*i.e.*, $\overrightarrow{eSWR^+}$).

The density of $\overrightarrow{eSWR} \cdot \overrightarrow{g_{EGR}}$, $\overrightarrow{rSWR} \cdot \overrightarrow{g_{EGR}}$, and $\overrightarrow{eSWR} \cdot \overrightarrow{rSWR}$ were calculated (Figure 7A–D). $\overrightarrow{rSWR} \cdot \overrightarrow{g_{EGR}}$ showed biphasic distributions.

By taking the differences between the distribution of $\overrightarrow{rSWR^+} \cdot \overrightarrow{g_{EGR}}$ (Figure 7A & B) and those of $\overrightarrow{rSWR^-} \cdot \overrightarrow{g_{EGR}}$ (Figure 7C & D), the contributions of SWR was calculated (Figure 7E & F), revealing a shift in the direction of $\overrightarrow{g_{EGR}}$ (Figure 7E & F; *red rectangles*).

Additionally, only in Mismatch OUT task, $\overrightarrow{eSWR^+} \cdot \overrightarrow{rSWR^+}$ was less than $\overrightarrow{eSWR^+} \cdot \overrightarrow{rSWR^-}$ (baseline periods) (Figure 7F; *pink circles*); in other words, \overrightarrow{eSWR} and \overrightarrow{rSWR} directed in the adverse direction solely in Mismatch OUT task but in Match IN task (Figure 7E; *pink circles*).

3. Discussion

This study hypothesized that hippocampal neurons exhibit distinct representations, or trajectories, in low-dimensional spaces during a WM task in humans, particularly during SWR periods. First, we projected the multiunit spikes in MTL regions during a Sternberg task onto three-dimensional spaces by GPFA (Figures 1D–E and Figure 2A). The distance of trajectory among WM phases ($\|\overrightarrow{g_{FGE}}\|$, $\|\overrightarrow{g_{FGM}}\|$, $\|\overrightarrow{g_{FGR}}\|$, $\|\overrightarrow{g_{EGM}}\|$, $\|\overrightarrow{g_{EGR}}\|$, and $\|\overrightarrow{g_{MGR}}\|$) was larger in the hippocampus than the EC and amygdala (Figure 2E), showing more dynamical neural activity in the hippocampus during the WM task. Additionally, the distance of trajectory between the encoding and retrieval phases in the hippocampus ($\|\overrightarrow{g_{FGE}}\|$) was positively correlated with memory load (Figures 3C & D), indicating it as a reflection of WM processing. Furthermore, the neural trajectory in the hippocampus showed transient increase during SWRs (Figure 5). Finally, the hippocampal neural trajectory fluctuated between encoding and retrieval states, with a shift from encoding to retrieval during SWR events (Figure 7). In sum, these results demonstrated the hippocampal neural behavior in a WM task in humans.

First, we found that the distance of the neural trajectory among the four phases ($\|\overrightarrow{g_{FGE}}\|$, $\|\overrightarrow{g_{FGM}}\|$, $\|\overrightarrow{g_{FGR}}\|$, $\|\overrightarrow{g_{EGM}}\|$, $\|\overrightarrow{g_{EGR}}\|$, and $\|\overrightarrow{g_{MGR}}\|$) was longer in the hippocampus compared to the EC and amygdala, even considering the distance from O ($\|\overrightarrow{g_F}\|$, $\|\overrightarrow{g_E}\|$, $\|\overrightarrow{g_M}\|$, and

$\|\overrightarrow{g_R}\|$) in those regions (Figures 2C–E). These results indicate hippocampal participation in the WM task, which is partially supported by previous findings of hippocampal persistent firing in the maintenance phase [3] [4] [5] [6]. However, by applying GPFA to multiunit activity during the 1-s level resolution of WM task, we revealed that the neural trajectory in low dimensional space displayed memory-load dependency between the encoding and retrieval phase, represented as $\|\overrightarrow{g_{EGR}}\|$ (Figure 3). Overall, these results provide evidence that the hippocampus is linked to WM processing.

The validity of our analysis of confining to putative CA1 regions (Figure 4) is supported by several factors. First, this targeted approach stems from well-established observations that SWRs are time-locked to synchronous spike bursts of interneurons and pyramidal neurons [39] [40] [41] [42], potentially around $50 \mu\text{m}$ radius of the recording site [43]. Additionally, in the present study, we found the increase in SWRs' incidence at 0–400 ms of the retrieval phase (Figure 4D). This result is consistent with previous reports showing increased SWR occurrence before spontaneous verbal recall [21] [22]. Thus, our result is not only consistent but also extends the finding to a triggered retrieval condition. Moreover, the log-normal distributions of SWR duration and ripple band peak amplitude observed in this study (Figure 4C & E) align with the consensus in this field [44]. Therefore, our approach of limiting recording sites for putative CA1 regions would have contributed to precision of SWR detection. One limitation is that the increase in trajectory distance from O during SWR (Figure 5) would have been biased to greater due to the channel selection; however, this is not critical for our major findings.

Interestingly, the trajectory directions in the retrieval phase oscillated between the encoding and retrieval states both in baseline and SWR periods (Figures 7C & D). In addition, the balance of such fluctuation was shifted from the encoding to retrieval state during SWR (Figures 7E & F). These results are again consistent with previous reports suggesting SWR's role in memory recall [21] [22]. Our result adds another layer of understanding, that is, SWR occurs when hippocampal representation proceeds "from encoding" to retrieval states. Therefore, our results provide new aspects of hippocampal representations: (i) neural fluctuations between encoding and retrieval states during a WM task and (ii)

SWR as a switching representation from encoding to retrieval states.

Moreover, our study reveals WM-task type specific directions between encoding- and retrieval-SWRs (Figure 7E–F). Specifically, eSWR and rSWR directed in the adverse direction not in Match IN but in Mismatch OUT task. These result might be explained by the memory engram theory [45]. In fact, Match In task exposed subjects to once-seen letter, while Mismatch OUT task a novel letter which was not included in the encoding phase. These results suggest that SWR is related to working cognitive processess in humans.

In conclusion, our study has demonstrated that hippocampal activity fluctuates between encoding and retrieval states during a WM task and exhibits a significant transition "from encoding" to retrieval during SWR periods.

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Contributors

Y.W. and T.Y. conceptualized the study; Y.W. performed the data analysis; Y.W. and T.Y. wrote the original draft; and all authors reviewed the final manuscript.

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Declaration of Interests

The authors declare that they have no competing interests.

Data and code availability

The data is available on G-Node (<https://doi.gin.g-node.org/10.12751/g-node.d76994/>).

The source code is available on GitHub (<https://github.com/yanagisawa-lab/hippocampal-neural-fluctuation-during-a-WM-task-in-hum>)

Inclusion and Diversity Statement

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

Declaration of Generative AI in Scientific Writing

The authors employed ChatGPT, provided by OpenAI, for enhancing the manuscript's English language quality. After incorporating the suggested improvements, the authors meticulously revised the content. Ultimate responsibility for the final content of this publication rests entirely with the authors.

Tables

		Hipp. head		Hipp. Body		EC		Amy.			
Subject ID	# of										
sessions	AHL	AHR	PHL	PHR	ECL	ECR	AL	AR		SOZ	
#1	4	o	x	o	o	o	x	o	x		AHR, LR
#2	7	o	o	o	o	o	o	o	o		AHR, PHR
#3	3	o	o	o	o	o	o	o	x		AHL, PHL
#4	2	o	o	o	o	o	o	o	o		AHL, AHR, PHL, PHR
#5	3	o	x	x	o	x	x	o	x		DRR
#6	6	o	o	o	o	o	o	o	o		AHL, PHL, ECL, AL
#7	4	o	o	o	o	o	o	o	o		AHR, PHR
#8	5	o	o	o	o	o	o	o	o		ECR
#9	2	o	o	o	o	o	o	o	o		ECR, AR

Table 1 – Electrode positions of the dataset

The electrode positions and the seizure onset zones. Regions marked as "o" were available, but those marked as "x" (*navy*) were not available in the dataset. Abbreviations: AHL, left hippocampal head; AHR, right hippocampal head; PHL, left hippocampal body; PHR, right hippocampal body; ECL, left entorhinal cortex; ECR, right entorhinal cortex; AL, left amygdala; AR, right amygdala, SOZ: seizure onset zone.

	Hipp. head		Hipp. body	
Subject	AHL	AHR	PHL	PHR
#1	0.60 ± 0.14	n.a.	n.a.	0.1 ± 0
#2	0.21 ± 0.16	0.17 ± 0.21	0.18 ± 0.22	0.20 ± 0.15
#3	0.40 ± 0.42	0.83 ± 0.12	n.a.	n.a.
#4	0.10 ± 0.00	0.10 ± 0.00	0.90 ± 0.00	0.10 ± 0.14
#5	n.a.	n.a.	n.a.	n.a.
#6	0.63 ± 0.06	n.a.	n.a.	0.27 ± 0.06
#7	0.10 ± 0.00	0.35 ± 0.35	0.37 ± 0.47	0.10 ± 0.00
#8	0.13 ± 0.10	n.a.	0.28 ± 0.49	n.a.
#9	n.a.	0.85 ± 0.07	0.15 ± 0.07	n.a.

Table 2 – The silhouette score of UMAP clustering between SWR^+ candidates and SWR^- candidates

The silhouette scores (mean ± SD for sessions by subject) of UMAP clustering on SWR^+ candidates and SWR^- candidates (Figure 4A) were based on their underlying multiunit spike patterns (mean values were 0.205 [0.285], median [IQR]; Figure 4B).

	# of sessions	# of trials	ROI	# of SWRs	SWR incidence [Hz]
#1	2	100	AHL	274	0.34
#3	2	97	AHR	325	0.42
#4	2	99	PHL	202	0.26
#6	2	100	AHL	297	0.37
#9	2	97	AHR	72	0.092784
	Total = 10	Total = 493		Total = 1,170	0.30 ± 0.13
(mean ± SD)					

Table 3 – The number of defined SWR events

The table summarizes the statistics of putative CA1 regions and SWRs. Only the first two sessions (sessions 1 and 2) from each subject were utilized to reduce the sampling bias.

Figures

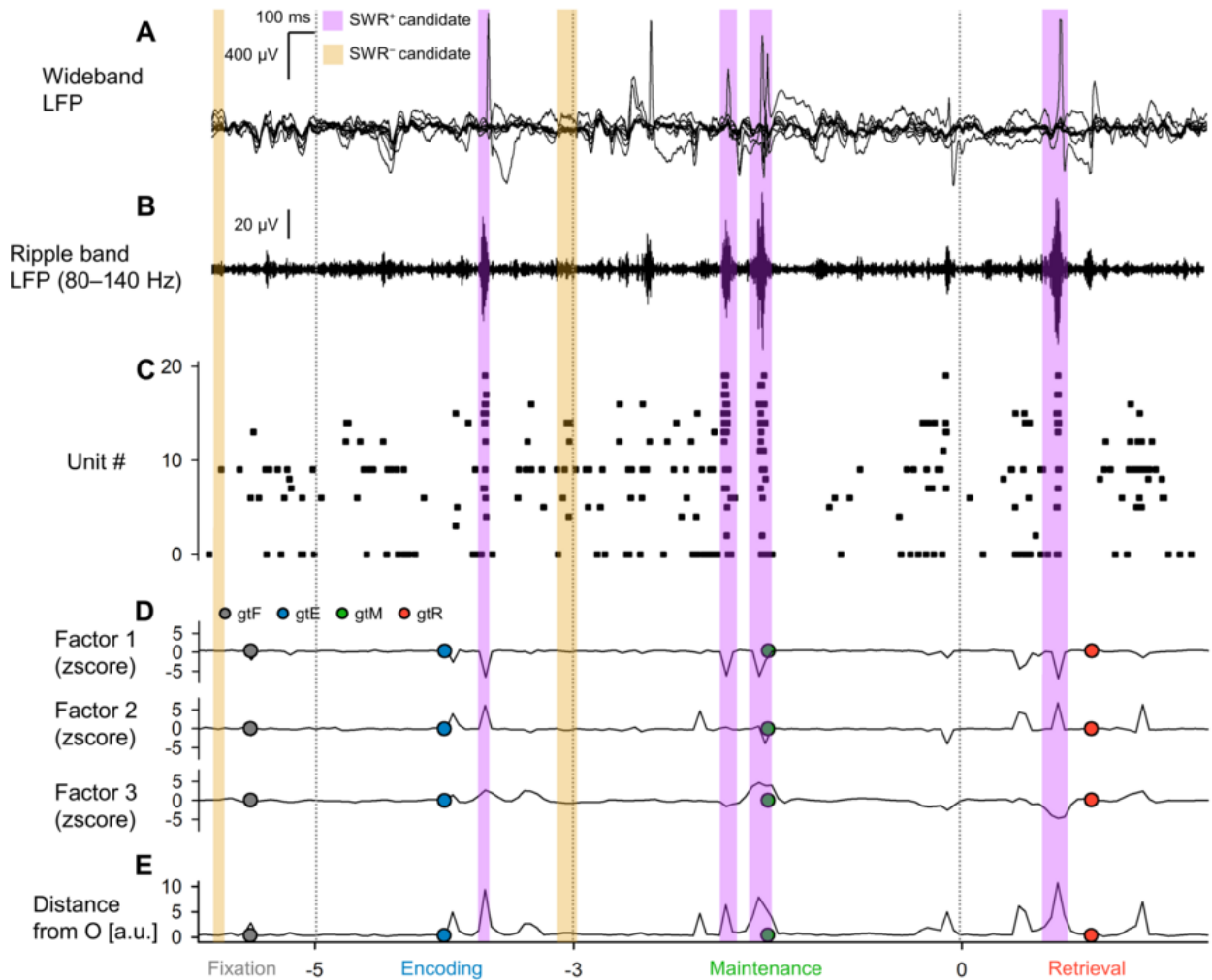


Figure 1 – Local field potential (LFP), multiunit activity, and neural trajectory of the hippocampus during a modified Sternberg task

A. Wideband LFP Trace: Displaying the intracranial encephalogram (iEEG) data for subject #6, session #2, trial #5, located in the left hippocampal head. This trace showcases the varying phases of the modified Sternberg working memory task, including fixation (gray, 1 s), encoding (blue, 2 s), maintenance (green, 3 s), and retrieval (red, 2 s). **B.** Ripple Band LFP: This presents the LFP trace in the ripple frequency band from the same trial and session. **C.** Multiunit Spike Raster: A plot illustrating multiunit spikes recorded during the mentioned trial. **D.** Neural Trajectory Factors: Depicting the first three factors of the hippocampal trajectory for the trial, determined by Gaussian-process factor analysis on spike counts with 50-ms bins. The dot circles signify the geometric median coordinates for each of the four task phases. **E.** Trajectory Distance Analysis: Shows the distance of the initial three trajectory factors from the origin (*O*). Note that highlighted rectangles overlaying the figure indicate the timings for SWR⁺ candidates (in purple) and SWR[−] candidates (in yellow).

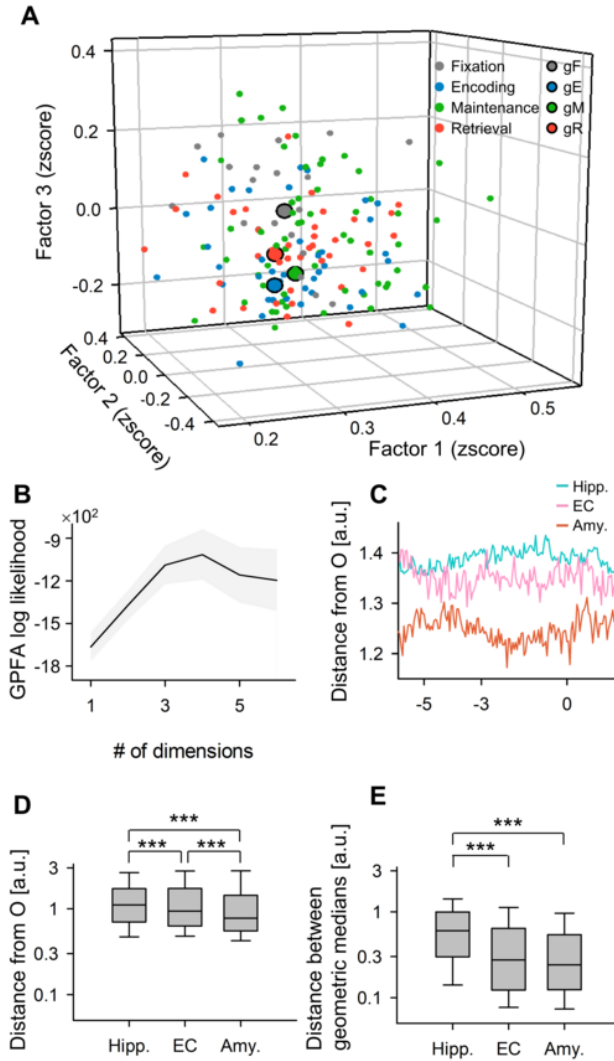


Figure 2 – State-dependent hippocampal neural trajectory

A. Neural Trajectory Visualization: This three-dimensional plot represents the neural trajectory of the left hippocampus derived from the Gaussian-process factor analysis (GPFA). Smaller points indicate coordinates of 50-ms neural trajectory bins within a session (median of 50 trials). Larger points with *black* edges represent geometric medians for each phase of the Sternberg working memory task, distinguished by colors for fixation (*gray*), encoding (*blue*), maintenance (*green*), and retrieval (*red*). **B. GPFA Model Log-likelihoods:** The graph showcases the log-likelihood predictions of GPFA models concerning the number of dimensions. Notably, the optimal dimension was identified as three using the elbow method. **C. Distance Analysis from Origin:** The plot represents the trajectory distance from the origin (*O*) for the hippocampus (Hipp.), entorhinal cortex (EC), and amygdala (Amy.), plotted against the time since probe onset. **D. Trajectory Distance in Medial Temporal Regions:** The box plots display the trajectory distances from *O* for each region, with the hippocampus showing the greatest distance, followed by the EC and the Amygdala. **E. Inter-phase Trajectory Distances:** This visualizes the distance variations between trial trajectory geometric medians of task phases. The distances in the hippocampus were observed to be more pronounced compared to the EC or Amygdala.

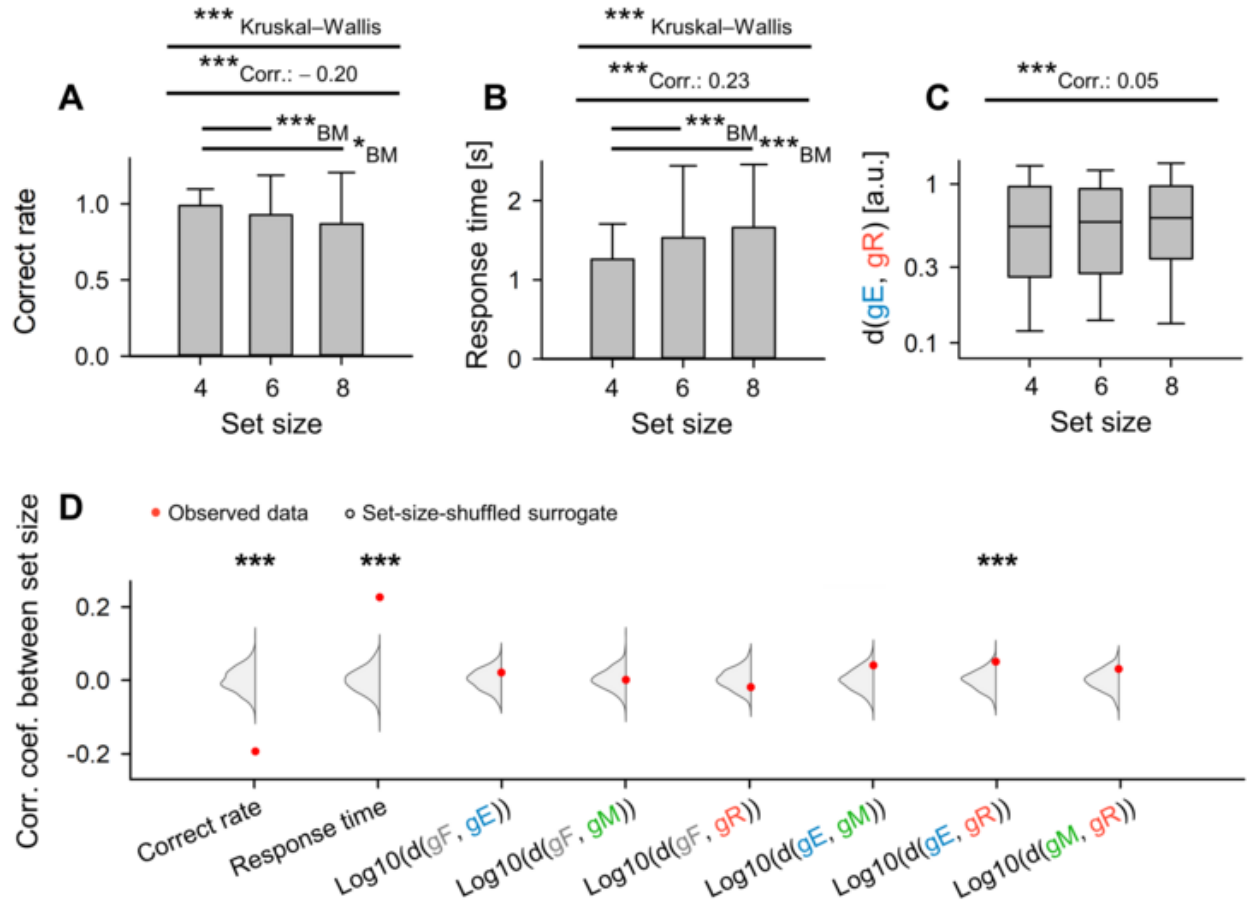


Figure 3 – Influence of memory load on encoding and retrieval states in the hippocampus

A. Relationship between set size (number of letters encoded) and correct rate in the WM task. Notably, a negative correlation was observed (coefficient = -0.20 , $***p < 0.001$), analyzed using set-size-shuffled surrogate. Significance was determined using Kruskal–Wallis test, followed by the Brunner–Munzel test with Bonferroni correction. **B.** Association between set size and response time post-probe initiation. A positive correlation was detected (coefficient = 0.23 , $***p < 0.001$), analyzed using set-size-shuffled surrogate. Statistical tests included the Kruskal–Wallis and post-hoc Brunner–Munzel tests with Bonferroni correction. **C.** Analysis of set size against the distance between geometric medians during encoding and retrieval phases (represented as $\|g_E g_R\|$). A correlation of 0.05 was noted for set size and $\log_{10}\|g_E g_R\|$, analyzed using set-size-shuffled surrogate. **D.** Experimentally observed correlations between set size and various parameters are illustrated by *red* dots. The *gray* violin plots contrast these findings with set-size-shuffled surrogate data ($n = 1,000$), emphasizing the significant observed correlation coefficients ($***p < 0.001$). Abbreviations: g_F , g_E , g_M , g_R represent the geometric median of trajectories during fixation, encoding, maintenance, and retrieval phases, respectively.

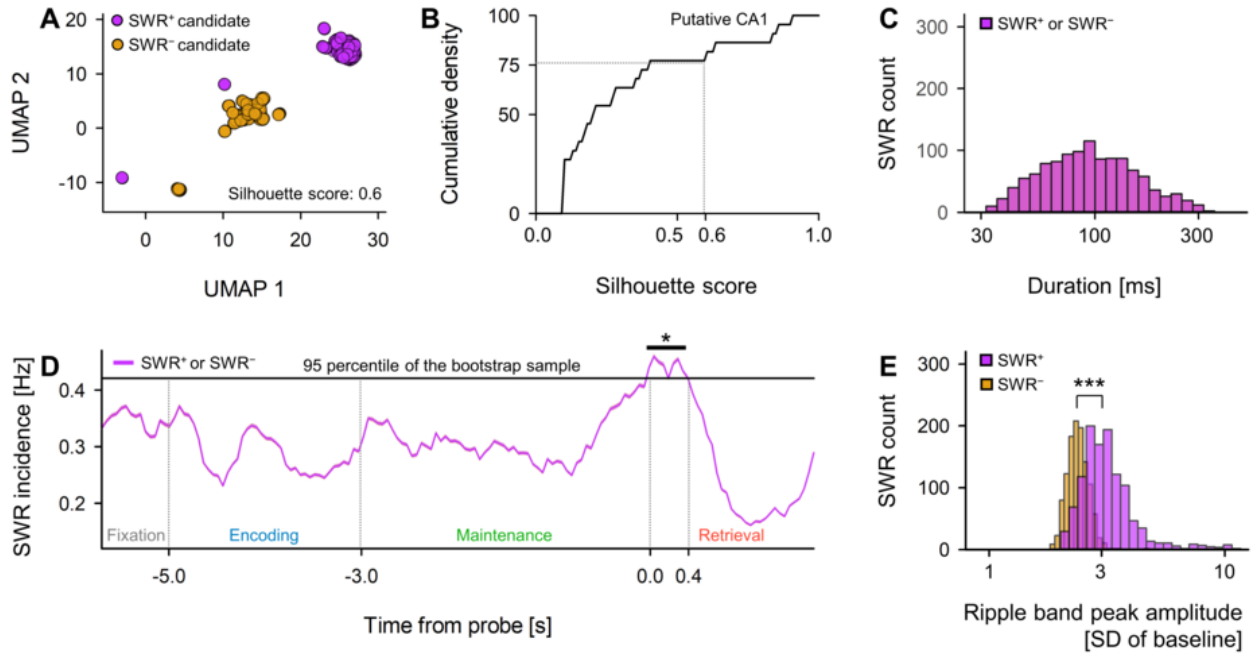


Figure 4 – SWR detection in putative CA1 regions

A. Two-dimensional UMAP (uniform manifold approximation and projection)[46] representation of unit activities during SWR⁺ candidates (*purple*; potential SWR events) and SWR⁻ candidates (*yellow*; controls for SWR⁺ candidates). **B.** Cumulative density plot illustrating silhouette scores for different hippocampal regions (refer to Table 2). Regions with silhouette scores exceeding 0.60 (75th percentile) are considered as putative CA1 regions. Within these regions, SWR⁺ and SWR⁻ candidates are identified and categorized as SWR⁺ ($n = 1,170$) and SWR⁻ ($n = 1,170$), respectively. **C.** Overlapping duration distributions [ms] for SWR⁺ (*purple*; 93.0 [65.4], median [IQR]) and SWR⁻ (*yellow*; 93.0 [65.4], median [IQR]). The identical overlap results from their definition criteria. **D.** Depiction of ripple incidence [Hz] for both SWR⁺ (*purple*) and SWR⁻ (*yellow*) relative to time from the probe, represented as mean \pm 95% confidence interval. However, the 95% confidence interval might not be readily discernible due to its narrow range. Note the significant elevation in SWR incidence between 0–400 ms post-probe, surpassing the 95th percentile of bootstrap samples (0.421 [Hz], $*p < 0.05$). **E.** Histogram showcasing ripple band peak amplitude distributions for SWR⁻ (*yellow*; 2.37 [0.33], median [IQR]) versus SWR⁺ (*purple*; 3.05 [0.85], median [IQR]). A notable difference is present between the two, confirmed with $***p < 0.001$ using the Brunner–Munzel test.

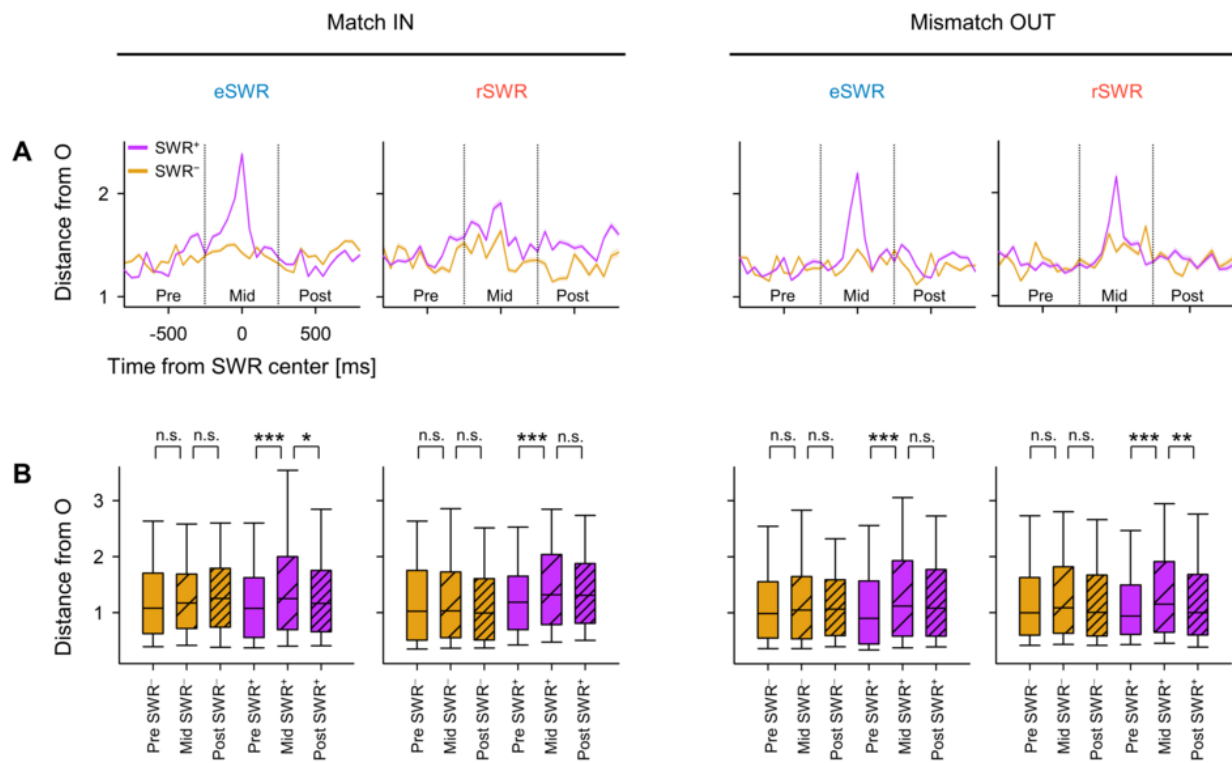


Figure 5 – Transient neural trajectory change during SWR

A. Distance from the origin (*O*) of the peri-sharp-wave-ripple trajectory (mean \pm 95% confidence interval; however, the interval may not be easily visible due to the narrow range.). **B.** The distance from the origin during the mid-SWR⁺ period was found to be longer compared to the corresponding pre-SWR⁺ period (* p < 0.05, ** p < 0.01, *** p < 0.001; Brunner–Munzel test). Abbreviations: SWR, sharp-wave ripple events; eSWR, SWR during the encoding phase; rSWR, SWR during the retrieval phase, SWR⁺, SWR event; SWR⁻ control events for SWR⁺; pre-SWR, mid-SWR, or post-SWR, the time interval from -800 to -250 ms, from -250 to +250 ms, or from +250 to +800 ms from the center of SWR.

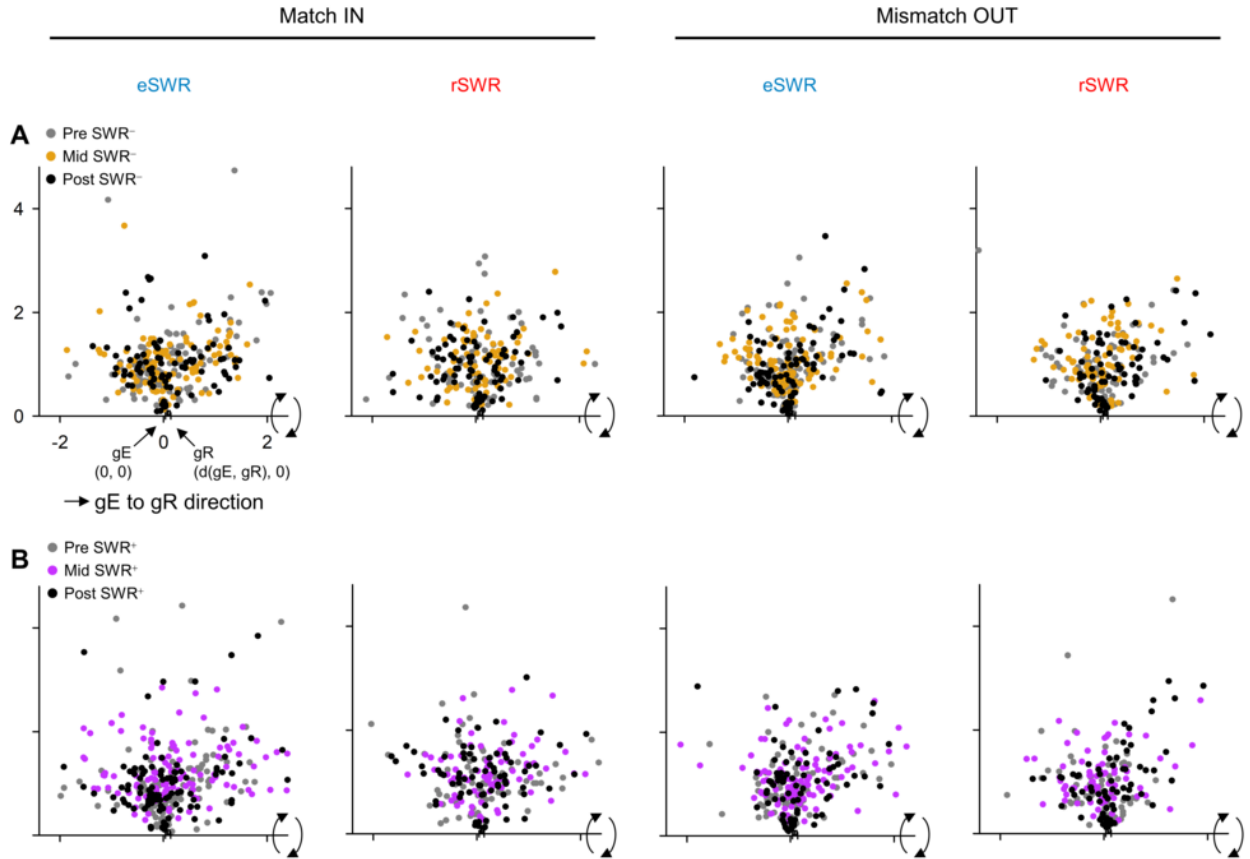


Figure 6 – Coordinates of neural trajectory during sharp-wave ripple aligned by encoding and retrieval states.

A. Hippocampal neural trajectories during pre- (gray), mid- (yellow), and post-SWR⁻ (black) in Match IN (left) and Mismatch OUT task (right). **B.** The equivalents for SWR⁺ instead of SWR⁻, though mid-SWR⁺ is depicted with purple. All data points underwent adjustments and rotations to fit a two-dimensional representation, positioning g_E at (0, 0) and g_R at ($\|g_E g_R\|$, 0). The $\|g_E g_R\|$ metric varies across sessions, and its median \pm IQR (with medians approximately 0.2) is presented on the x-axes. Note: some intervals might be challenging to discern because of their narrow range. In this two-dimensional depiction, both the distances and angles preserve their relationships as in the original three-dimensional space. Abbreviations: SWR, sharp-wave ripple events; eSWR, SWR during the encoding phase; rSWR, SWR during the retrieval phase; SWR⁺, SWR event; SWR⁻ control events for SWR⁺; pre-SWR, mid-SWR, or post-SWR, the time interval from -800 to -250 ms, from -250 to +250 ms, or from +250 to +800 ms from the center of SWR.

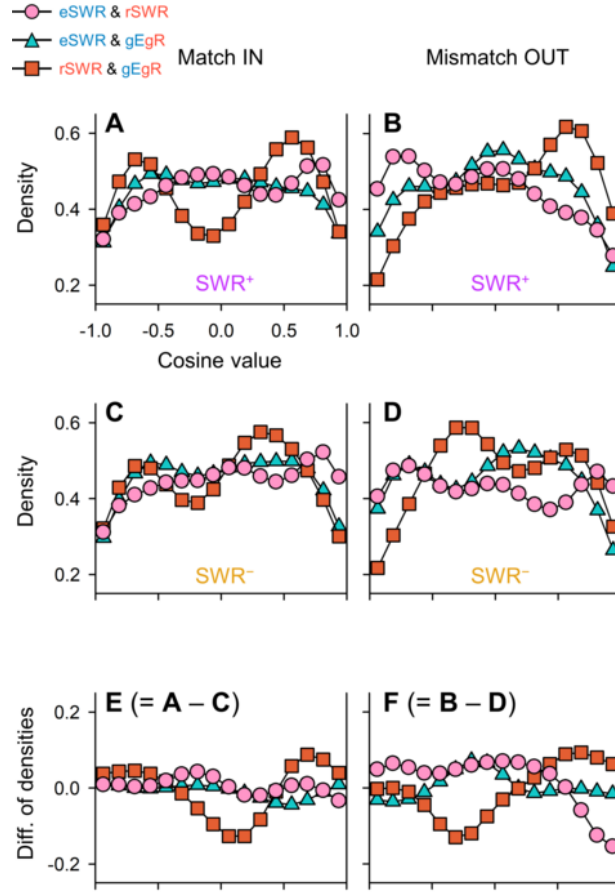


Figure 7 – Analysis of Neural Trajectory Directions during SWR between Encoding, and Retrieval States.

A–B Kernel density estimation (KDE) distribution of $\overrightarrow{eSWR^+} \cdot \overrightarrow{rSWR^+}$ (pink circles), $\overrightarrow{eSWR^+} \cdot \overrightarrow{gEgR}$ (blue triangles), and $\overrightarrow{rSWR^+} \cdot \overrightarrow{gEgR}$ (red rectangles) in Match In (**A**) and Mismatch OUT task (**B**). **C–D**. The corresponding distributions of SWR^- in response to those of SWR^+ in **A–B**. **E–F**. The differences in distributions, highlighting the SWR components (**E** = **C** – **A**; **F** = **B** – **D**). Note the inverse directionality between $\overrightarrow{eSWR^+}$ and $\overrightarrow{rSWR^+}$ only in Mismatch OUT task (pink circles in **E–F**). Additionally, the shifts from the retrieval to encoding states were observed for SWR components both in Match IN and Mismatch OUT tasks (red rectangles in **E–F**).