

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 Between Memory Encoding and Retrieval States During a Working Memory Task in Humans

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Abstract

Working memory (WM) is crucial for a multitude of cognitive functions, but its underlying neural mechanisms at play remain to be fully understood. Interestingly, the hippocampus and sharp-wave ripples (SWRs)— transient and synchronous oscillation observed in the hippocampus — garner recognition for their significance in memory consolidation and retrieval, albeit their correlation with WM tasks remains to be elucidated. Here we show that during a WM task, multi-unit activity patterns in the hippocampus display distinctive dynamics, especially during SWR periods. This study analyzed intracranial electroencephalography data from the medial temporal lobe (MTL) of nine patients with epilepsy, recorded during an eight-second Sternberg task. Utilizing Gaussian-process factor analysis, we extracted low-dimensional neural representations or trajectories (NT) within MTL regions

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during the Sternberg WM task. The results revealed significant variations in NT of the hippocampus compared to those of the entorhinal cortex and the amygdala. Additionally, the distance of the NT between the encoding and retrieval phases was dependent on memory load. Crucially, hippocampal NT oscillated during the retrieval phase between encoding and retrieval states in a task-dependent manner. Notably, these fluctuations shifted from encoding to retrieval states during SWR episodes. These findings reaffirm the role of the hippocampus in WM tasks and propose a hypothesis: the hippocampus transitions its functional state from encoding to retrieval during SWRs in WM tasks.

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

8 figures, 3 tables, 221 words for abstract, and 3178 words for main text

1. Introduction

Working memory (WM) is crucial in everyday life; however, its neural mechanism has yet to be fully elucidated. Specifically, the hippocampus’s involvement in WM processing, a pivotal region for memory, is the subject of ongoing research [1, 2, 3, 4, 5, 6, 7, 8, 9]. Understanding the hippocampus’ role in working memory is instrumental in deepening our comprehension of cognitive processes and could potentially enhance cognitive abilities.

Current evidence suggests that transient, synchronized oscillations, termed sharp-wave ripples (SWRs) [10], are associated with various cognitive functions. SWRs have traditionally been linked with long-term memory functions such as memory replay [11, 12, 13, 14, 15], memory consolidation [16, 17, 18, 19], memory recall [20, 21, 22], and neural plasticity [23, 24]. However, only a subset of studies has investigated the role of SWRs in WM tasks [25, 26]. This gap in our understanding motivates the current study to further investigate the potential involvement of SWRs in WM, particularly given their fundamental computational manifestation in hippocampal

processing.

Recent studies have found that low-dimensional representations in hippocampal neurons can explain WM task performances. Specifically, the firing patterns of place cells [27, 28, 29, 30, 31], found in the hippocampus, have been identified within dynamic, nonlinear three-dimensional hyperbolic spaces in rats [32]. Additionally, grid cells in the entorhinal cortex (EC), which is the main pathway to the hippocampus [33, 34, 35], exhibited a toroidal geometry during exploration in rats [36].

However, these existing studies predominantly focus on spatial navigation in rodents, presenting several limitations. First, the temporal resolution of navigation tasks is insufficient, obscuring the precise timing of memory acquisition and recall. Second, the presence of noise in signals recorded during rodent movement complicates the detection of SWRs [37]. Third, the generalization to humans and tasks other than spatial navigation remains unclear. Given these limitations, it is crucial to explore SWRs in a controlled, less noisy environment to better understand their potential role in WM tasks in humans.

Considering these factors, this study investigates the hypothesis that hippocampal neurons in humans exhibit low-dimensional neural trajectories (NTs) that depend on WM load, particularly during SWR periods. To test this hypothesis, we employed a dataset of patients performing an eight-second Sternberg task (1 s for fixation, 2 s for encoding, 3 s for maintenance, and 2 s for retrieval) with high temporal resolution. Intracranial electroencephalography (iEEG) signals within the medial temporal lobe (MTL) were recorded for these patients [38]. To investigate low-dimensional NTs, we utilized Gaussian-process factor analysis (GPFA), an established method for analyzing neural population dynamics [39, 40, 41, 42, 43, 44, 45, 46, 47].

2. Methods

2.1. Dataset

The dataset used in this study, which is publicly available, comprises nine epilepsy patients performing a modified the Sternberg task [38]. This task includes four phases: fixation (1s), encoding (2s), maintenance (3s), and retrieval (2s). During the encoding phase, participants were presented with a set of four, six, or eight alphabet letters. They were then tasked with determining whether a probe letter displayed during the retrieval phase had previously appeared (the correct response for Match IN task) or not (the correct response for Mismatch OUT task). Intracranial electroencephalography (iEEG) signals were captured with a 32 kHz sampling rate within a 0.5–5,000 Hz frequency range, using depth electrodes in the MTL regions: the anterior head of the left and right hippocampus (AHL and AHR), the posterior body of the hippocampus (PHL and PHR), the entorhinal cortex (ECL and ECR), and the amygdala (AL and AR), as depicted in Figure 1A and Table 1. The iEEG signals were subsequently downsampled to 2 kHz. Correlations between variables such as set size and correct rate were examined (Figure ??S1). Multiunit spike timing was determined via a spike sorting algorithm [48] using the Combinato package (<https://github.com/jniediek/combinato>)(Figure 1C).

2.2. Calculation of NT using GPFA

NTs, also referred to as 'factors', in the hippocampus, EC, and amygdala were determined using GPFA [39] applied to the multi-unit activity data for each session, performed with the elephant package (<https://elephant.readthedocs.io/en/latest/reference/gpfa.html>). The bin size was set to 50 ms, without overlaps. Each factor was z-normalized across all sessions, and the Euclidean distance from the origin (O) was then computed.

For each NT within a region such as AHL, geometric medians (g_F for fixation, g_E for encoding, g_M for maintenance, and g_R for retrieval phase) were calculated by determining the median coordinates of the NT during the

four phases. An optimal GPFA dimensionality was found to be three using the elbow method obtained by examining the log-likelihood values through a three-fold cross-validation approach (Figure 2B).

2.3. Identifying SWR candidates from hippocampal regions

Potential SWR events within the hippocampus were detected using a widely used method [49]. LFP signals from a region of interest (ROI) like AHL, were re-referenced by deducting the averaged signal from locations outside the ROI (for instance, AHR, PHL, PHR, ECL, ECR, AL, and AR). The re-referenced LFP signals were then filtered with a ripple-band filter (80–140 Hz) to determine SWR candidates, marked as SWR^+ candidates. SWR detection was carried out using a published tool (https://github.com/Eden-Kramer-Lab/ripple_detection) [50], with the bandpass range adjusted to 80–140 Hz for humans [21, 22, 49], unlike the initial 150–250 Hz range typically applied to rodents [10].

Control events for SWR^+ candidates, labeled as SWR^- candidates, were detected by randomly shuffling the timestamps of SWR^+ candidates across all trials and subjects. The resulting $\text{SWR}^+/\text{SWR}^-$ candidates were then visually inspected.

2.4. Defining SWRs from putative hippocampal CA1 regions using UMAP clustering

Potential SWRs were differentiated from SWR candidates in putative CA1 (cornu Ammonis 1) regions. These regions were initially defined as follows: $\text{SWR}^+/\text{SWR}^-$ candidates in the hippocampus were projected into a two-dimensional space based on overlapping spike counts per unit using a supervised method, UMAP (Uniform Manifold Approximation and Projection) [51]. Clustering validation was performed by calculating the silhouette score [52] from clustered samples. Regions in the hippocampus, which scored above 0.6 on average across sessions (75th percentile), were identified as putative CA1 regions, resulting in the identification of five electrode positions from five patients.

SWR⁺/SWR⁻ candidates in these predetermined CA1 regions were categorized as SWR⁺/SWR⁻, and thus they no longer retained their candidate status. The duration and ripple band peak amplitude of SWRs were found to follow log-normal distributions. Each time period of SWR was partitioned relative to the time from the SWR center into pre- (at -800 ms to -300 ms from the SWR center), mid- (at -250 to +250 ms), and post-SWR (at +300 to +800 ms) times.

2.5. Statistical Evaluation

Both the Brunner–Munzel test and the Kruskal–Wallis test were executed using the SciPy package in Python [53]. Correlational analysis was conducted by determining the rank of the observed correlation coefficient within its associated set-size-shuffled surrogate using a customized Python script. The bootstrap test was implemented with an in-house Python script.

3. Results

3.1. *i*EEG Recording and NT in MTL Regions during the Sternberg Task

Our analysis employed a publicly accessible dataset [38], which comprises LFP signals (Figure 1A) from MTL regions (Table 1) recorded during the execution of the modified Sternberg task. We extracted SWR⁺ candidates from LFP signals that were filtered in the 80–140 Hz ripple band (Figure 1B), originating in all hippocampal regions (refer to Methods section). Meanwhile, SWR⁻ candidates, control events for SWR⁺ candidates, were defined at the same timestamps but distributed across different trials (Figure 1). The dataset encompassed multi-unit spikes (Figure 1C), recognized via a spike sorting algorithm [48]. Employing GPFA [39], we applied this to 50-ms windows of binned multi-unit activity without overlaps to determine the NTs, or factors, of MTL regions by session and region (Figure 1D). We normalized each factor per session and region, for instance, session #2 in AHL of subject #1. We then calculated the Euclidean distance from the origin (*O*) (Figure 1E).

3.2. Correlation of Hippocampal NT with the Sternberg Task

Figure 2A exhibits a distribution of median NTs, comprising 50 trials, within the three main factor spaces. Utilizing the elbow method, we established the optimal embedding dimension for the GPFA model as three (Figure 2B). The NT distance from the origin (O) — represented as $\|g_F\|$, $\|g_E\|$, $\|g_M\|$, and $\|g_R\|$ — in the hippocampus surpassed the corresponding distances in the EC and amygdala (Figure 2C & D).¹

Similarly, we computed the distances between the geometric medians of four phases, namely $\|g_F g_E\|$, $\|g_F g_M\|$, $\|g_F g_R\|$, $\|g_E g_M\|$, $\|g_E g_R\|$, and $\|g_M g_R\|$. The hippocampus showed larger distances between phases than those in the EC and amygdala.²

3.3. Memory-Load-Dependent NT Distance between Encoding and Retrieval States in the Hippocampus

Regarding memory load in the Sternberg task, we observed a negative correlation between the correct rate of trials and the set size, which denotes the number of letters to be encoded (Figure 3A).³ Concomitantly, a positive correlation was noted between the response time and set size (Figure 3B).⁴

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). Generally, $p < 0.001$ 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;

Next, we discovered a positive correlation between set size and the NT distance separating the encoding and retrieval phases ($\log_{10}\|\mathbf{g}_{\text{EGR}}\|$) (Figure 3C).⁵ However, distances between other phase combinations did not highlight statistically significant correlations (Figures 3D and ??).

3.4. Detection of Hippocampal SWRs from Putative CA1 Regions

To enhance the precision of recording sites and SWR detection, we approximated the electrode placements in the CA1 regions of the hippocampus using distinguished multi-unit spike patterns during SWR events. $\text{SWR}^+/\text{SWR}^-$ candidates from each session and hippocampal region were embedded in two-dimensional space using UMAP (Figure 4A).⁶ With the silhouette score as a quality metric for clustering (Figure 4B and Table 2), recording sites demonstrating an average silhouette score exceeding 0.6 across all sessions were identified as putative CA1 regions.⁷ (Tables 2 and 3). We identified five putative CA1 regions, four of which were not indicated as seizure onset zones (Table 1).

Subsequently, $\text{SWR}^+/\text{SWR}^-$ candidates within these putative CA1 regions were labeled as SWR^+ and SWR^- , respectively⁸ (Table 3). Both SWR^+ and SWR^- manifested identical durations⁹ due to their definitions and followed a log-normal distribution (Figure 4C). During the initial 400 ms of the retrieval phase, an increase in SWR^+ incidence was found¹⁰ (Figure 4D). The

correlation coefficient = 0.22, $p < 0.001$

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

⁶Consider the AHL in session #1 of subject #1 as a case in point.

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

⁸These definitions produced equal counts for both categories: SWR^+ ($n = 1,170$) and SWR^- ($n = 1,170$).

⁹These definitions result in equal durations 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]; $p < 0.05$.

peak ripple band amplitude of SWR^+ surpassed that of SWR^- and followed a log-normal distribution (Figure 4E).¹¹.

3.5. Transient Change in Hippocampal NT during SWR

We assessed the 'distance' of the NT from the origin (O) during SWR events in both encoding and retrieval phases (Figure 5A). Observing the increase in distance during SWR, as illustrated in Figure 5A, we categorized each SWR into three stages: pre-, mid-, and post-SWR. Hence, the distances from the origin O during those SWR intervals are identified as $\|\text{pre-eSWR}^+\|$, $\|\text{mid-eSWR}^+\|$ and others.

As a result, $\|\text{mid-eSWR}^+\|$ ¹² exceeded $\|\text{pre-eSWR}^+\|$ ¹³, and $\|\text{mid-rSWR}^+\|$ ¹⁴ was larger than $\|\text{pre-rSWR}^+\|$ in both the Match IN and Mismatch OUT tasks.¹⁵.

3.6. Visualization of Hippocampal NT during SWR in Two-Dimensional Spaces

Having observed NT 'jumping' during SWR (Figure 5), we visualized the three-dimensional NTs of pre-, mid-, and post-SWR events during the encoding and retrieval phases (Figure 6). The distance between these was found to be memory-load dependent (Figure 3).

To provide two-dimensional visualization, we linearly aligned peri-SWR NTs by setting \mathbf{g}_E at the origin (0, 0) and \mathbf{g}_R at $(\|\mathbf{g}_{EGR}\|, 0)$. Subsequently, we rotated these aligned NTs around the \mathbf{g}_{EGR} axis (the x axis), ensuring that the distances from the origin O in the original three-dimensional spaces

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).

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

and angles from $\overrightarrow{g_{EGR}}$ are retained in the two-dimensional equivalent.

Scatter plot visualization of NTs within these two-dimensional spaces revealed distinct distributions of peri-SWR NTs based on phases and task types. A notable example of this is the observation that the magnitude of $\|\text{mid-eSWR}^+\|$ exceeds that of $\|\text{pre-eSWR}^+\|$ (Figure 6B), which is consistent with our previous observations (Figure 5).

3.7. Fluctuations of Hippocampal NTs between Encoding and Retrieval States

Subsequently, we investigated the 'direction' of the NT in relation to $\overrightarrow{g_{EGR}}$, which was found to be dependent on memory load (Figure 3). The directions of the SWRs were determined by the NT at -250 ms and $+250$ ms from their center, denoted as, for example, \overrightarrow{eSWR}^+ . We calculated the cosine similarities between $\overrightarrow{g_{EGR}}$, \overrightarrow{eSWR} , and \overrightarrow{rSWR} in both SWR (SWR^+) and baseline periods (SWR^-) (Figure 7A–D).

$\overrightarrow{rSWR}^+ \cdot \overrightarrow{g_{EGR}}$ exhibited a biphasic distribution. By computing the difference between the distribution of $\overrightarrow{rSWR}^+ \cdot \overrightarrow{g_{EGR}}$ (Figure 7A & B) and that of $\overrightarrow{rSWR}^- \cdot \overrightarrow{g_{EGR}}$ (Figure 7C & D), we were able to determine the contributions of SWR (Figure 7E & F), which indicated a shift in the direction of $\overrightarrow{g_{EGR}}$ (Figure 7E & F: *red rectangles*).

Furthermore, $\overrightarrow{eSWR}^+ \cdot \overrightarrow{rSWR}^+$ was less than $\overrightarrow{eSWR}^- \cdot \overrightarrow{rSWR}^-$ strictly in Mismatch OUT task (Figure 7F: *pink circles*). In other words, eSWR and rSWR pointed in the opposite direction exclusively in Mismatch OUT task but not Match IN task (Figure 7E: *pink circles*).

4. Discussion

This study hypothesizes that in low-dimensional spaces during a WM task in humans, hippocampal neurons form unique NTs, primarily during SWR periods. Initially, multi-unit spikes in the MTL regions were projected onto three-dimensional spaces during a Sternberg task using GPFA (Figure 1D–E & Figure 2A). The NT distances across WM phases ($\|\overrightarrow{g_{FGE}}\|$, $\|\overrightarrow{g_{FGM}}\|$, $\|\overrightarrow{g_{FGR}}\|$, $\|\overrightarrow{g_{EGM}}\|$, $\|\overrightarrow{g_{EGR}}\|$, and $\|\overrightarrow{g_{MGR}}\|$) were significantly larger in

the hippocampus compared to the EC and amygdala (Figure 2C–E), indicating dynamic and responsive neural activity in the hippocampus during the WM task. Also, in the hippocampus, the NT distance between the encoding and retrieval phases ($\|\mathbf{g}_F\mathbf{g}_E\|$) correlated positively with memory load (Figure 3C–D), reflecting WM processing. The hippocampal neural NT transiently expanded during SWRs (Figure 5). Lastly, the hippocampal neural NT alternated between encoding and retrieval states, transitioning from encoding to retrieval during SWR events (Figure 7). These findings explain aspects of hippocampal neural activity during a WM task in humans and offer new insights into SWRs as a state-switching element in hippocampal neural states.

The distance of the neural NT across the phases was significantly longer in the hippocampus compared to the EC and amygdala, even when considering the distance from O in these regions (Figure 2C–E). This establishes the involvement of the hippocampus in the WM task, corroborating previous studies indicating hippocampal persistent firing during the maintenance phase [4, 5, 6, 3]. However, in the present study, applying GPFA to multi-unit activity during a one-second level resolution of the WM task revealed that the neural NT in low-dimensional space presented a memory-load dependency between the encoding and retrieval phases, denoted as $\|\mathbf{g}_E\mathbf{g}_R\|$ (Figure 3). These findings support the association of the hippocampus with WM processing.

Our analysis focused on putative CA1 regions (Figure 4) in order to enhance the validity of recording site and the true positive rate of SWR detection. This criterion is supported by accumulated evidence. For instance, SWRs synchronize with spike bursts of interneuron and pyramidal neuron [54, 55, 56, 57], potentially within a 50 μm radius of the recording site [58]. Additionally, we identified increased incidence of SWRs during the first 0–400 ms of the retrieval phase (Figure 4D). This finding aligns with previous reports of heightened SWR occurrence preceding spontaneous verbal recall [21, 22], extending our understanding to a triggered retrieval condition.

Moreover, the log-normal distributions of both SWR duration and ripple band peak amplitude observed in this study (Figure 4C & E) coincide with the consensus in this field [49]. Therefore, these results support the electrode placement and detected SWRs in this study. One could argue that the neural trajectory (NT) distance increase from O during sharp wave-ripples (SWRs) (Figure 5) may be artificially inflated towards higher values due to channel selection using UMAP clustering on spike counts. However, this potential bias does not affect the direction of NT, the memory-load dependency, nor the WM task dependency identified in this study.

Interestingly, during the retrieval phase, NT directions alternated between encoding and retrieval states during both baseline and SWR periods in a task-dependent manner (Figure 7C & D). Additionally, the balance of this fluctuation transitioned from encoding to retrieval state during SWR events (Figure 7 E & F). These results align with previous studies on the role of SWR in memory retrieval [21, 22]. Our findings suggest that (i) neuronal oscillation between encoding and retrieval states occurs during a WM task and (ii) SWR events serve as indicators of the transition in hippocampal neural states from encoding to retrieval during a WM task.

Moreover, our study noted differences specific to the WM-task type between encoding- and retrieval-SWRs (Figure 7E–F). Notably, opposing movements of encoding-SWR (eSWR) and retrieval-SWR (rSWR) were not observed in Match IN task but were apparent in Mismatch OUT task. Memory engram theory [59] might explain these observations: Match In task presented participants with previously shown letters, while Mismatch OUT task introduced a new letter absent in the encoding phase. This explanation underscores the significant role of SWR in human cognitive processes.

In conclusion, this study illustrates that during a WM task in humans, hippocampal activity fluctuate between encoding and retrieval states, uniquely transitioning from encoding to retrieval during SWR events. These findings offer novel insights into the neural correlates and functionality of working memory within the hippocampus.

Data Availability Statement

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-fluctuations-during-a-WM-task-in-humans>).

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Ethics Declarations

All study participants provided their written informed consent, subsequent to the approval from the pertinent institutional ethics review board (Kantonale Ethikkommission Zürich, PB 2016–02055).

Author Contributions

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.

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

Subject ID	# of sessions	AHL	AHR	PHL	PHR	ECL	ECR	AL	AR	
1	4	✓	n.a.	✓	✓	✓	n.a.	✓	n.a.	A
2	7	✓	✓	✓	✓	✓	✓	✓	✓	AF
3	3	✓	✓	✓	✓	✓	✓	✓	n.a.	AF
4	2	✓	✓	✓	✓	✓	✓	✓	✓	AHL & AF
5	3	✓	n.a.	n.a.	✓	n.a.	n.a.	✓	n.a.	
6	6	✓	✓	✓	✓	✓	✓	✓	✓	AHL & P
7	4	✓	✓	✓	✓	✓	✓	✓	✓	AF
8	5	✓	✓	✓	✓	✓	✓	✓	✓	
9	2	✓	✓	✓	✓	✓	✓	✓	✓	E

Table 1 – Electrode Distribution within the Dataset

This figure denotes the placements of electrodes and seizure onset zones. Regions marked with ✓ were included in the dataset, while those imprinted with n.a. were absent. The abbreviations used are as follows: 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 represents the seizure onset zone.

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 – Silhouette Score of UMAP Clustering for SWR^+ Candidates and SWR^- Candidates

The silhouette scores (mean \pm SD across sessions per subject) for UMAP clustering of SWR^+ candidates and SWR^- candidates are based on their respective multi-unit spike patterns (Figure 4A). The mean scores were 0.205 and the standard deviation was 0.285, calculated for the interquartile range (IQR; Figure 4B).

Subject ID	# 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	0.09
Total = 10		Total = 493		Total = 1170	0.30 \pm 0.13 (mean \pm SD)

Table 3 – Summary of Detected SWRs

The table provides statistics of presumptive CA1 regions and SWR events. Only the initial two sessions (sessions 1 and 2) from each subject were included in our analysis to reduce sampling bias.

Figures

Figure 1 – Local Field Potentials, Multiunit Activity, and Neural Trajectories in the Hippocampus During a Modified Sternberg Task

A. Representative wideband LFP signals for intracranial EEG recording from the left hippocampal head are presented. This recording took place while the subject performed a modified Sternberg working memory task. Task stages included fixation (1 s, *gray*), encoding (2 s, *blue*), maintenance (3 s, *green*), and retrieval (2 s, *red*). **B.** Displays the associated ripple band LFP traces. Note *purple* and *yellow* rectangles, which denote the timings for SWR^+ candidates and SWR^- candidates, respectively (the latter serving as control events for SWR^+). **C.** A raster plot illustrates multi-unit spikes from the LFP traces. These spikes have been sorted using a spike algorithm [48]. **D.** Shows neural trajectories (NTs) computed by GPFA[39] based on spike counts per unit with 50-ms bins. The geometric median of each phase is marked by dot circles. **E.** Indicates the distance of the NT from the origin point O .

Figure 2 – State-Dependent Neural Trajectory of Hippocampal Neurons

A. Neural trajectories (NTs) depicted as a point cloud within the first three-dimensional factors derived from GPFA [39]. The smaller dots represent 50-ms NT bins, and the larger dots with *black* edges denote the geometric medians for each phase in the Sternberg working memory task: fixation ($\|\mathbf{g}_F\|$, *gray*), encoding ($\|\mathbf{g}_E\|$, *blue*), maintenance ($\|\mathbf{g}_M\|$, *green*), and retrieval ($\|\mathbf{g}_R\|$, *red*). **B.** The figure presents the log-likelihood of the GPFA models versus the number of dimensions used to embed multi-unit spikes found in the medial temporal lobe (MTL) regions. Specifically, the elbow method identified three as the optimal dimension. **C.** This panel displays the distance of the NTs from the origin (O) for the hippocampus (Hipp.), entorhinal cortex (EC), and amygdala (Amy.), plotted against the time elapsed from the probe onset. **D.** The NT distance from O within the MTL regions is shown. The hippocampus has the greatest distance, followed by the EC and the Amygdala. **E.** The box plot illustrates inter-phase NT distances within the MTL regions.

Figure 3 – Positive Correlation between Memory Load and Neural Trajectory Distance in the Hippocampus between Encoding and Retrieval Phases

A. The relationship between set size (number of letters to be encoded) and accuracy in the working memory task (coefficient = -0.20 , $***p < 0.001$). **B.** The correlation between set size and response time (coefficient = 0.23 , $***p < 0.001$). **C.** The correlation between set size on the inter-phase distances between the encoding and retrieval phases ($\|g_{EGR}\|$) (correlation coefficient = 0.05 , $***p < 0.001$). **D.** Experimental observations of correlations between set size and the following parameters: accuracy, response time, $\log_{10} \|g_{FGE}\|$, $\log_{10} \|g_{FGM}\|$, $\log_{10} \|g_{FGR}\|$, $\log_{10} \|g_{EGM}\|$, $\log_{10} \|g_{EGR}\|$, and $\log_{10} \|g_{MGR}\|$ represented by *red* dots. The kernel density plots (*gray*) illustrate the corresponding shuffled surrogate with set size ($n = 1,000$) ($***ps < 0.001$).

Figure 4 – Detection of SWRs in Putative CA1 Regions

A. Two-dimensional UMAP [51] projection displays multi-unit spikes during SWR^+ candidates (*purple*) and SWR^- candidates (*yellow*). **B.** A cumulative density plot indicates silhouette scores, reflecting UMAP clustering quality (see Table 2). Hippocampal regions with silhouette scores exceeding 0.60 (equivalent to the 75th percentile) are identified as putative CA1 regions. SWR^+ and SWR^- candidates, which were recorded from these regions, are classified as SWR^+ and SWR^- respectively ($n_s = 1,170$). **C.** Identical distributions of SWR^+ (*purple*) and SWR^- (*yellow*) distributions, based on their definitions (93.0 [65.4] ms, median [IQR]). Note that these distributions exhibit log-normality. **D.** Identical SWR incidence for both SWR^+ (*purple*) and SWR^- (*yellow*), relative to the probe’s timing (mean $\pm 95\%$ confidence interval). However, 95% confidence interval may not be visibly apparent due to their narrow ranges. Note that a significant SWR incidence increase was detected during the initial 400 ms of the retrieval phase (0.421 [Hz], $*p < 0.05$, bootstrap test). **E.** Distributions of ripple band peak amplitudes for SWR^- (*yellow*; 2.37 [0.33] SD of baseline, median [IQR]) and SWR^+ (*purple*; 3.05 [0.85] SD of baseline, median [IQR]) are manifested ($***p < 0.001$, the Brunner–Munzel test). Note the log-normality for SWR^+ events.

Figure 5 – Transient Change in Neural Trajectory during SWR

A. The distance from origin (O) of the peri-sharp-wave-ripple neural trajectory (mean $\pm 95\%$ confidence interval). The intervals may be obscured due to their minimal ranges. **B.** The distance from the origin (O) during the pre-, mid-, and post-SWR periods is demonstrated ($*p < 0.05$, $**p < 0.01$, $***p < 0.001$; Brunner–Munzel test applied). Abbreviations: SWR, sharp-wave ripple events; eSWR, SWR during the encoding phase; rSWR, SWR within the retrieval phase; SWR^+ , positive SWR event; SWR^- , control events for SWR^+ ; pre-, mid-, or post-SWR refer to the time intervals from -800 to -250 ms, from -250 to $+250$ ms, or from $+250$ to $+800$ ms, respectively, all relative to the SWR center.

Figure 6 – Visualization of Neural Trajectory During SWR in Two-Dimensional Space

The panels depict hippocampal neural trajectories (NTs) during SWR projected onto two-dimensional spaces. **A.** Shows the hippocampal NTs as point clouds during pre-SWR⁻ (*gray*), mid-SWR⁻ (*yellow*), and post-SWR⁻ (*black*). **B.** Conveys the equivalent for SWR⁺ rather than SWR⁻. The projection was executed as follows: First, a linear transformation placed \mathbf{g}_E at the origin O (0,0), and \mathbf{g}_R at $(\|\mathbf{g}_{ER}\|, 0)$. The point cloud was subsequently rotated around the \mathbf{g}_{ER} axis (similar to the x axis) for adaptation to two-dimensional spaces. Thus, within these two-dimensional spaces, the distances from point O and the angles for the \mathbf{g}_{ER} axis are retained as in the original three-dimensional spaces created by GPFA. Abbreviations: SWR denotes sharp-wave ripple events; eSWR refers to SWR during the encoding phase; rSWR signals SWR during the retrieval phase; SWR⁺, characterizes an SWR event; SWR⁻ signifies control events for SWR⁺; pre-SWR, mid-SWR, or post-SWR, represent the time intervals from -800 to -250 ms, from -250 to $+250$ ms, or from $+250$ to $+800$ ms from the center of the SWR.

Figure 7 – Direction of Neural Trajectory During SWR Based on Encoding and Retrieval States

A–B The kernel density estimation distributions of $\overrightarrow{\text{eSWR}^+} \cdot \overrightarrow{\text{rSWR}^+}$ (*pink circles*), $\overrightarrow{\text{eSWR}^+} \cdot \overrightarrow{\text{gEGR}}$ (*blue triangles*), and $\overrightarrow{\text{rSWR}^+} \cdot \overrightarrow{\text{gEGR}}$ (*red rectangles*) in Match In (*A*) and Mismatch OUT tasks (*B*). **C–D** The corresponding distributions of SWR^- instead of those of SWR^+ in *A* and *B*. **E–F** The differences in the distributions of SWR^+ and those of SWR^- , showing SWR components ($E = C - A$ & $F = D - B$). Biphasic distributions of $\overrightarrow{\text{rSWR}^-} \cdot \overrightarrow{\text{gEGR}}$ indicate fluctuations between the encoding and retrieval states during the Sternberg task. Also, contradicting directionality between $\overrightarrow{\text{eSWR}^+}$ and $\overrightarrow{\text{rSWR}^+}$ was observed (pink circles) not in Match IN task (**E**), but in Mismatch OUT task (**F**). Lastly, transitions from the retrieval to encoding states are evident in the SWR components in both Match IN and Mismatch OUT tasks (*red rectangles* in *E–F*).