

## Review

## Mesoscopic Neural Representations in Spatial Navigation

Lukas Kunz,<sup>1,7,\*</sup> Shachar Maidenbaum,<sup>2,7,\*</sup> Dong Chen,<sup>3,7</sup> Liang Wang,<sup>3,4,5,8,\*</sup> Joshua Jacobs,<sup>2,8</sup> and Nikolai Axmacher<sup>6,8,\*</sup>

Recent evidence suggests that mesoscopic neural oscillations measured via intracranial electroencephalography exhibit spatial representations, which were previously only observed at the micro- and macroscopic level of brain organization. Specifically, theta (and gamma) oscillations correlate with movement, speed, distance, specific locations, and goal proximity to boundaries. In entorhinal cortex (EC), they exhibit hexadirectional modulation, which is putatively linked to grid cell activity. Understanding this mesoscopic neural code is crucial because information represented by oscillatory power and phase may complement the information content at other levels of brain organization. Mesoscopic neural oscillations help bridge the gap between single-neuron and macroscopic brain signals of spatial navigation and may provide a mechanistic basis for novel biomarkers and therapeutic targets to treat diseases causing spatial disorientation.

### Towards a Multilevel Neural Code of Spatial Navigation

Spatial navigation is a core ability of most animals and humans [1]. Successful navigation requires highly specialized **neural representations** (see [Glossary](#)) that encode information about the shape and content of the environment, neural codes that reflect the location, direction, and speed of the navigating organism, and neural mechanisms that underlie goal-directed behavior within the environment. These spatial representations have been identified at various levels of brain organization ranging from place, head-direction, border, and **grid cells** [2–7] up to the macroscopic level of fMRI, which revealed large-scale representations of locations [8], environments [9], directions [10–12], borders [13], and grid-like hexadirectional orientation [14–16].

Traditionally, there has been a gap between studies of spatial representations at the levels of single neurons and fMRI brain networks. In this review, we focus on the potential of **intracranial electroencephalography** (iEEG) recordings of **local field potentials** (LFPs) to bridge this gap, by providing spatial information at the mesoscopic level of relatively small neural assemblies ([Box 1](#)). These recordings have provided evidence for oscillatory signatures of multiple key components of spatial navigation, ranging from more general signals of movement, speed, and distance, to representations of specific spatial locations, goal proximity to boundaries, and, very recently, to mesoscopic grid-like representations. First, we review these results and discuss their relationship to the neural representations of space obtained at the microscopic, single-neuron level, and at the macroscopic level as observed with fMRI. We then elaborate on the basic relationships between single-neuron spiking activity, LFPs, and the fMRI **blood oxygen level-dependent (BOLD) signal** to address the question of how mesoscopic spatial representations relate to their microscopic and macroscopic counterparts. Finally, we discuss the functional and clinical relevance of mesoscopic representations during spatial navigation.

### Highlights

Neural representations of spatial navigation have mainly been studied at the microscopic level of single neurons or at the macroscopic level of fMRI.

Recent intracranial electroencephalography (iEEG) recordings in patients with epilepsy revealed neural representations of spatial features, including travelled distance, goal proximity to boundaries, and grid-like hexadirectional orientation. These representations occur at the mesoscopic level of brain oscillations, particularly in the theta frequency band.

Mesoscopic representations of space bridge the gap between their micro- and macroscopic counterparts. Experimentally testable scenarios may explain how the mesoscopic spatial representations relate to single-neuron firing, other neural oscillations, and fMRI signals.

Neural spatial representations may offer novel tools for biomarkers of neurological and psychiatric diseases.

<sup>1</sup>Epilepsy Center, Medical Center – University of Freiburg, Faculty of Medicine, University of Freiburg, Freiburg im Breisgau, Germany

<sup>2</sup>Department of Biomedical Engineering, Columbia University, New York City, NY, USA

<sup>3</sup>CAS Key Laboratory of Mental Health, Institute of Psychology, Beijing, China

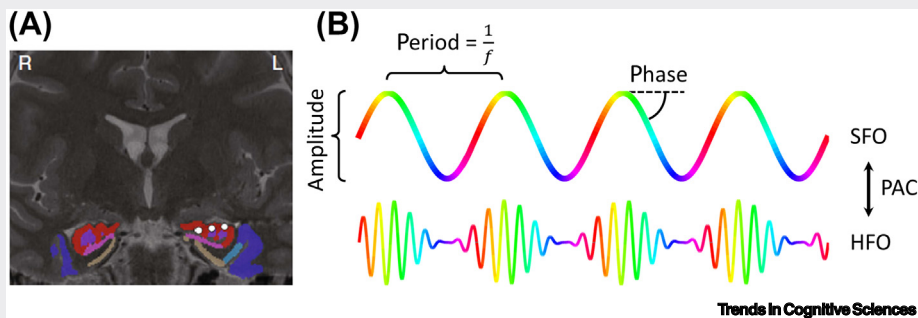
<sup>4</sup>CAS Center for Excellence in Brain Science and Intelligence Technology, Shanghai, China

<sup>5</sup>Department of Psychology, University of Chinese Academy of Sciences, Beijing, China

<sup>6</sup>Department of Neuropsychology, Institute of Cognitive Neuroscience, Faculty of Psychology, Ruhr University Bochum, Bochum, Germany

**Box 1. Detecting and Analyzing Mesoscopic Brain Oscillations**

Since the discovery of the scalp EEG by Hans Berger during the 1920s, it has been known that cognition is accompanied by brain oscillations across a range of frequencies. EEG oscillations are classically divided into frequency bands: delta (1–4 Hz), theta (4–8 Hz), alpha (8–13 Hz), beta (13–30 Hz), low gamma (30–50 Hz), and high gamma (>50 Hz). Although it is more common to record these signals from the scalp, in rare circumstances researchers are able to measure these oscillations directly with high spatiotemporal resolution from the human brain, when neurosurgical patients are implanted with intracranial depth, strip, or grid macroelectrodes for clinical purposes (Figure 1). Contacts from these iEEG macroelectrodes record neuronal activity within a radius of several millimeters when electrode contacts are relatively large [138]. Some patients are implanted with microelectrodes which have smaller conductive surfaces and record neural signals from a region within a few hundred micrometers [98]. These microelectrode recordings also allow for the detection of multi- and single-unit spiking activity. During the implantation period, the patient can perform experimental tasks to identify neural activity related to cognitive functions [139]. Given that patients undergoing iEEG are usually bedridden, these studies often probe spatial navigation via virtual navigation tasks performed on a bedside computer, although differences to real-world navigation have to be taken into account. During the analysis of iEEG data, both univariate and multivariate approaches (e.g., multivariate pattern analysis [140] and representational similarity analysis [141]) have been used effectively to identify general and feature-specific neural representations during spatial navigation.



**Figure 1. Recording and Quantifying Mesoscopic Oscillations.** (A) Coronal MR image showing an intracranial electroencephalography (iEEG) depth electrode in a patient with epilepsy. Three electrodes in hippocampal CA1 are highlighted in white. Regions are colored based on an automatic segmentation procedure. (B) Main characteristics of brain oscillations include amplitude, period, and phase (as indicated by color). Phase-amplitude coupling (PAC) describes a systematic relationship between the phase of a slow-frequency oscillation (SFO) and the amplitude of a high-frequency oscillation (HFO). Abbreviations:  $f$ , frequency; L, left; R, right. Reproduced from [142] (A).

**Navigating the Landscape of Mesoscopic Spatial Representations****Movement, Speed, Time, and Distance**

Theta **oscillations** (Box 2) reliably appear during movement in the hippocampus of both rodents [17] and humans [18], although some characteristics of this signal differ between species. First, theta oscillations are more stable in rodents than in humans, in which theta oscillations occur in bursts of several cycles. Second, theta frequency may usually appear higher in rodents (4–10 Hz) than in humans (1–4 Hz) [19,20], possibly because larger anatomical assemblies tend to synchronize at lower frequencies [21]. However, this latter difference may also be attributed to real-world navigation versus virtual navigation [22,23] or to more frequent electrode placement in anterior parts of the human hippocampus (where theta oscillations may have a lower frequency) [24].

Another oscillatory signal that is involved in spatial navigation is the hippocampal gamma oscillation (30–100 Hz), which increases in power during movement [25,26] and interacts with theta oscillations, such that gamma power varies across the theta cycle [25,27]. This phenomenon of cross-frequency coupling between gamma power and theta phase generalizes to other cognitive domains [28] and is thought to reflect **phase coding** [29] to, for example, bind multiple stimuli into sequences.

<sup>7</sup>These authors contributed equally to this work.

<sup>8</sup>These authors contributed equally to this work.

\*Correspondence: [lukas.kunz@uniklinik-freiburg.de](mailto:lukas.kunz@uniklinik-freiburg.de) (L. Kunz), [shachar.maidenbaum@columbia.edu](mailto:shachar.maidenbaum@columbia.edu) (S. Maidenbaum), [lwang@psych.ac.cn](mailto:lwang@psych.ac.cn) (L. Wang), [joshua.jacobs@columbia.edu](mailto:joshua.jacobs@columbia.edu) (J. Jacobs), and [nikolai.axmacher@rub.de](mailto:nikolai.axmacher@rub.de) (N. Axmacher).

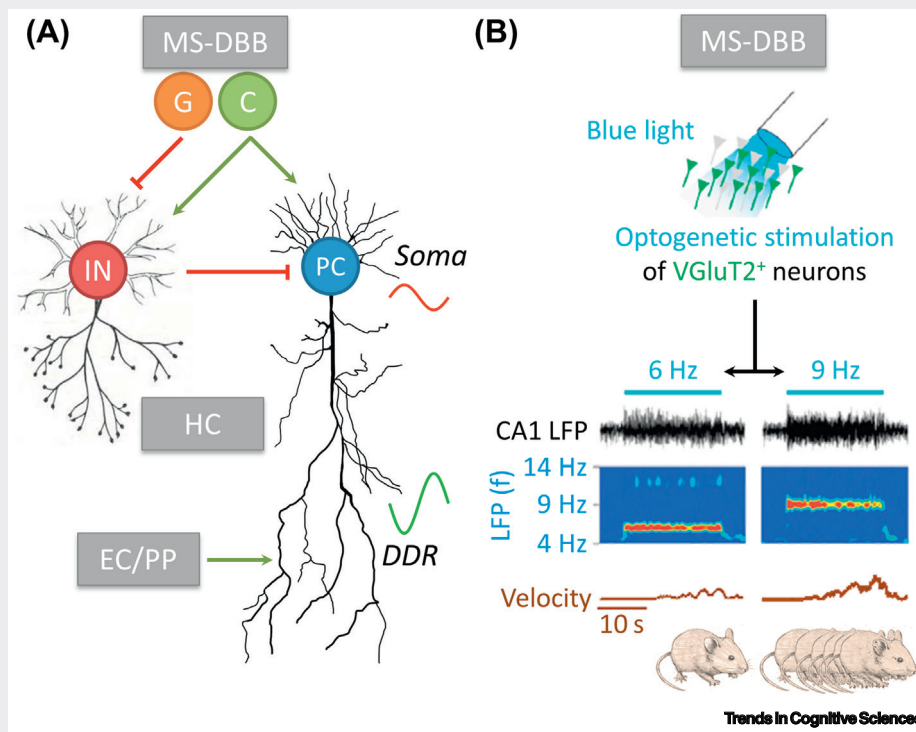
### Box 2. Basics of Theta Oscillations

Despite decades of research, the process of how hippocampal theta oscillations are generated is still under debate. Different models draw a complex picture of contributing brain regions, cell types, transmitters, and receptors, in which the medial septum and the diagonal band of Broca (MS-DBB) occupy major roles [17].

The classic model proposes that theta oscillations are generated by hippocampal pyramidal cells that receive inhibitory and excitatory input from the MS-DBB and the EC/perforant path, respectively (Figure 1). The inhibitory input is induced by glutamatergic [143] and cholinergic [144] MS-DBB neurons and leads, via hippocampal interneurons, to inhibitory postsynaptic potentials (IPSPs) at the soma of the hippocampal pyramidal cells. The excitatory input results in excitatory postsynaptic potentials (EPSPs) at their distal dendritic region. The summed activity of somatic IPSPs and distal dendritic EPSPs then generates theta oscillations with specific amplitudes and phases [17].

Later work suggested multiple modifications of this model [17], of which we describe three. First, different types of theta oscillations may emerge from different mechanisms. For example, atropine-sensitive theta oscillations, which appear during consummatory behaviors, may reflect the CA3 recurrent collateral system, whereas atropine-resistant theta oscillations, which correlate with movement, may be conveyed by entorhinal input. Second, interneuron activity driving the somatic IPSPs may not only depend on MS-DBB inputs, but also on CA1 recurrent collaterals, CA3 afferents, entorhinal afferents, and other interneurons with intrinsic oscillatory properties. These interneurons may comprise multiple classes [145] that target different parts of hippocampal pyramidal neurons. Third, the recurrent circuit of CA3 may form an intrinsic intrahippocampal theta generator and/or pyramidal neurons endowed with intrinsic properties to oscillate at theta frequency may produce single-cell theta oscillations.

Various potential functions have been proposed for hippocampal theta oscillations. Among others, these include information transfer between regions [17] and gating of information flow [146]. Furthermore, theta oscillations may enable the grouping and segregation of neuronal assemblies that were activated at different phases of the theta cycle, for example, during working memory [147]. Theta oscillations may also support mnemonic processes by inducing synaptic modifications of intrahippocampal associational pathways [148] and may allow the creation of temporally ordered memories [149]. Finally, theta oscillations may underlie phase coding by allowing single neurons to represent information by varying the timing of their spiking relative to theta oscillations during perception, navigation, and memory [29].



**Figure 1. Mechanistic Basis of Theta Oscillations.** (A) Classic theta model. See Box 2 text for explanations. Note that many extensions have been suggested to this model. (B) Medial septum and diagonal band of Broca (MS-DBB) VGLUT2<sup>+</sup> neurons as a key player in regulating locomotion. Increasing optogenetic stimulation (blue light) frequency of these neurons (from 6 to 9 Hz) leads to higher theta oscillation frequency in CA1 and is followed by increased velocity of the rodent. Abbreviations: C, cholinergic; DDR, distal dendritic region; EC/PP, entorhinal cortex and perforant path; G, glutamatergic; HC, hippocampus; IN, interneuron; PC, pyramidal cell. Adapted from [17] (A) and [34] (B).

In rodents, faster movement through an environment is associated with increases in both the power [30] and frequency [31] of theta oscillations. Similar results have been obtained in human iEEG recordings, during virtual navigation [19], and during active walking [22,23]. This mesoscopic representation of speed is paralleled by single neurons in the rodent hippocampus and medial EC that linearly increase their firing rate with speed [32,33]. A glutamatergic septal-hippocampal circuit that controls the initiation and velocity of locomotion [34] potentially occupies a key role in coordinating these single-neuron and mesoscopic representations of speed, which may nevertheless be dissociable from each other [35]. Functionally, increased theta frequencies during faster movements may serve to normalize the phase resolution within the theta cycle with respect to higher sampling rates of sensory information during faster movements [34].

To calculate travelled distance, speed must be integrated over time. Therefore, neural representations of time are likely important partners of spatial representations, particularly for accomplishing path integration [36]. In rodents, so-called 'time cells', which activate at particular time points in task intervals, are present in both the hippocampus [37] and lateral EC [38]. At the macroscopic level, hippocampal BOLD patterns reflect the processing of temporal intervals within sequences [39] and precise temporal memories are associated with increased fMRI activity in lateral EC [40]. Interestingly, a neural signature of time may also be visible at the mesoscopic level, since rodent hippocampal theta oscillations exhibit increased power in a temporal discrimination task [41].

Since theta oscillations are related to the processing of movement, speed, and time, it is not surprising that they also contain information about distance travelled: theta power increases during long- versus short-distance 'teleportation' during virtual navigation [42]. Similarly, theta oscillations are stronger before and during long versus short paths [43]. Increased theta power may serve to avoid cognitive overload due to increased sensory input during longer paths, for example by shortening the fraction of the theta cycle available for information processing and, thus, excluding less relevant information from being processed [44]. More broadly, these two studies indicate that theta oscillations are involved in the planning and initiation of movement, a finding that is paralleled by results in rodents [34]. Furthermore, an internal sense of movement appears to be sufficient for evoking hippocampal theta oscillations since they occur during virtual teleportation [42]. At the macroscopic level, hippocampal fMRI activity reflects distances between places [45] and path distance to goal locations [46], but (to our knowledge) no fMRI study so far has directly examined the BOLD correlates of previously travelled distance.

#### Locations, Borders, and Grids

Place cells fire whenever animals or humans are at a specific location within a spatial environment [2,3]. Place cells are mainly observed in the hippocampus [2,3], but they may also exist in extrahippocampal regions [47]. In addition, particularly during sharp-wave ripples, place cell activity often reflects future positions [48], and place cell sequences 'preplay' future paths to goal locations [49]. Furthermore, place cells activate sequentially within theta cycles, allowing theta cycles to simulate time-compressed trajectories through space [50]. Interestingly, mesoscopic oscillations provide additional information about self-location when place cell firing phases are taken into account: adding information about the theta phase at the moment of place cell firing increases one's ability to reconstruct the location of the animal [51,52]. As an extension of these findings, the position of a rat can be accurately decoded from the spatiotemporal structure of the theta rhythm obtained from multisite, high-density hippocampal LFP recordings, without utilizing information from place cell spiking activity [53]. These mesoscopic studies in rodents demonstrate that (multisite) oscillatory theta power and phase contain information regarding self-location. They are complemented by fMRI studies, which suggest that current and future

#### Glossary

##### **Allocentric reference frame:**

representation of the spatial environment with regard to features of the external world independent of the subject; typically contrasted with an egocentric reference frame that is centered on the subject.

##### **Blood oxygen level-dependent (BOLD) signal:**

signal recorded using fMRI that, by measuring blood oxygenation in brain voxels, allows indirect conclusions about the neural activity underlying changes in blood oxygenation.

**Frequency bands:** canonical classes that summarize neural oscillations based on their frequency: delta (<4 Hz), theta (4–8 Hz), alpha (8–13 Hz), beta (13–30 Hz), and gamma (>30 Hz).

**Grid cell:** A type of neuron with a firing field pattern which forms a grid that tessellates the environment into equilateral triangles; mainly observed in medial entorhinal cortex and presumably a key component of path integration. The firing pattern exhibits sixfold rotational symmetry and, thus, may be the source of hexadirectional modulation of meso- and macroscopic brain signals.

##### **Hexadirectional modulation:**

characteristic of a neural signal (e.g., theta power or BOLD amplitude) that varies as a function of a circular variable (e.g., movement direction) after having transformed the circular variable from the classical 360° space into a 60° space.

##### **Intracranial electroencephalography (iEEG):**

technique to measure electrical activity via surgically implanted electrodes from inside the skull. Electrodes comprise grids and strips attached to the surface of the brain (electrocorticography, ECoG) as well as depth electrodes inserted into the brain (stereotactic electroencephalography, SEEG).

##### **Local field potentials (LFPs):**

electrophysiological signals that integrate neural activity within localized brain areas; can be measured via iEEG that records the LFP from a few hundred micrometers up to a couple of millimeters depending on electrode size.

**Neural representations:** patterns of neural activity related to one specific stimulus of the external world or one specific mental content. Here, we define 'macroscopic neural representations' as those measured via fMRI, scalp EEG, or

locations can be decoded from human hippocampus and medial prefrontal cortex [8,54]. However, the possibility of finding reliable macroscopic representations of location in the hippocampus has been challenged [55] and their relationship to single place cells is unclear, because the hippocampus is thought to use sparse coding and hippocampal place cells lack a clear topographical organization [56–58].

Border cells in the medial EC [5] and boundary-vector cells in the subiculum [59] activate adjacent to or at specific distances from a border, respectively. These cells may be instrumental in anchoring place [60] and grid fields [61] to an **allocentric reference frame**. At the macroscopic level, border representations were identified in the parahippocampal place area and retrosplenial cortex [13], and transcranial magnetic stimulation demonstrated a causal role of the occipital place area in representing environmental boundaries [62]. While a direct mesoscopic signal for borders has yet to be found, a closely related representation was recently identified in human subiculum using iEEG: when subjects encoded goal locations near boundaries, power in the theta frequency range (4–10 Hz) was stronger compared with the encoding of goal -locations in the center of the virtual environment [63]. Of note, this study examined the distance between borders and the subject's goal locations (i.e., not the person's own locations) and, thus, provides further evidence that the brain's representations of space generalize from the self to others and also to inanimate objects for which an exact localization is important [64].

Grid cells fire at the vertices of an internally generated grid tiling the entire environment into equilateral triangles [6,7]. Thus far, they have mainly been observed in medial EC, where ~10–20% of all neurons are grid cells [33,65]. Grid cells may help to compute vector relationships and distances between spatial locations [66], potentially constituting the neural basis for path integration [16,67]. In fMRI, there is evidence for macroscopic grid-like representations (i.e., **hexadirectional modulation** of the BOLD signal as a function of movement direction) in healthy participants who were scanned as they moved through a virtual environment [14–16]. fMRI grid-like representations were observed not only in EC, but also in extrahippocampal regions, such as medial prefrontal cortex and parietal cortex ([14]; see Text S1 and Table S1 in the supplemental information online), warranting a brain-wide examination of grid cell density.

Crucially, two recent iEEG studies revealed grid-like representations at the mesoscopic level of brain organization (i.e., a hexadirectional modulation of entorhinal theta power by movement direction in patients with epilepsy who performed a **virtual navigation task**) [68,69]. Both studies found that this hexadirectional modulation was restricted to the EC and not present in neighboring similarly sized regions (hippocampus and amygdala) and other control regions, consistent with single-neuron [7] and fMRI recordings [14,15]. Furthermore, both studies showed that the effect was specific to sixfold rotational symmetry and to the theta frequency range. Importantly, both studies found a link between grid-like representations and spatial memory performance: stronger hexadirectional modulation correlated positively with better spatial memory performance [69], conceptually in agreement with previous fMRI studies [14–16,70], and stronger grid-like representations were demonstrated in later as compared with earlier parts of the experiment after significant learning had occurred [68]. Furthermore, stronger hexadirectional modulation was revealed in parts of the environment closer to the environmental boundary of the virtual arena [68], potentially related to stabilized grid cell firing via error-correcting border cell input after encounters with environmental boundaries [61]. Moreover, mesoscopic grid-like representations were particularly observed during fast movements [68], in line with prior fMRI studies [14,15] and potentially related to speed modulation of grid cells and conjunctive grid by head-direction cells [14,71].

MEG, 'mesoscopic neural representations' as those obtained via iEEG, and 'microscopic neural representation' as those occurring at the level of single neurons.

**Oscillations:** rhythmic fluctuations of neural activity, characterized by frequency (cycles per second), power (squared amplitude), and phase (momentary deflection angle). In general, neural oscillations bias input selection, temporally link neurons into assemblies, and facilitate synaptic plasticity, mechanisms that cooperatively support temporal representation and long-term consolidation of information.

**Phase coding:** neural mechanism by which information is represented via the phase of a (slow) oscillation at which activity of single neurons or cellular assemblies occurs.

**Phase locking** (here in the sense of spike-field coherence): phenomenon in which a neuron fires at consistent phases of a concurrent oscillation across time.

**Phase precession:** phenomenon that a neuron fires at successively earlier LFP phases across time.

**Virtual navigation task:** computerized tasks that let the subject explore virtual environments. This can be done with or without actual body movement, and via desktop or head mounted displays.



Mesoscopic grid-like representations were also observed in broad-band gamma activity: building upon evidence for grid cell activity [72] and fMRI grid-like representations [73,74] during visual exploration in monkeys and humans, respectively, a recent study searched for modulations of oscillatory power as a function of gaze direction [75]. Indeed, hexadirectionally modulated signals were evident in broadband gamma activity (60–120 Hz), both in magnetoencephalography (MEG) and in one patient with epilepsy, who was implanted with iEEG electrodes in the EC.

Taken together, unique patterns of brain oscillations emerge when animals and humans navigate spatial environments, with theta [17–20,22–24,30,31,34,35,42,43,50,76] and gamma oscillations [25–27] being associated most often with characteristics of spatial navigation, such as movement, speed, and distance. Crucially, recent evidence demonstrates that meso-scale neural networks oscillating in these **frequency bands** provide representations of locations, proximity of goals to boundaries, and grid-like hexadirectional orientation that were previously only observed at the microscopic or macroscopic level (Figure 1, Key Figure). In particular, theta [51–53,63,68,69] and gamma oscillations [75] stand out as carriers of this feature-specific information. Although several of these findings are preliminary, and have not yet been demonstrated across species, their possible functional significance merits further investigation. Particularly, a thorough brain-wide examination of both mesoscopic grid-like representations and the prevalence of grid cells would be of interest. Grid-like representations outside the EC may perform complementary computations for spatial navigation or they may support other cognitive functions that exploit similar computational principles as spatial navigation [66,77]. Furthermore, it remains elusive how the different scales of spatial representations are interconnected. This question is closely related to the fundamental issue of basic relationships between spiking activity, oscillatory power, and the BOLD signal.

### Common Grounds for Multilevel Spatial Representations?

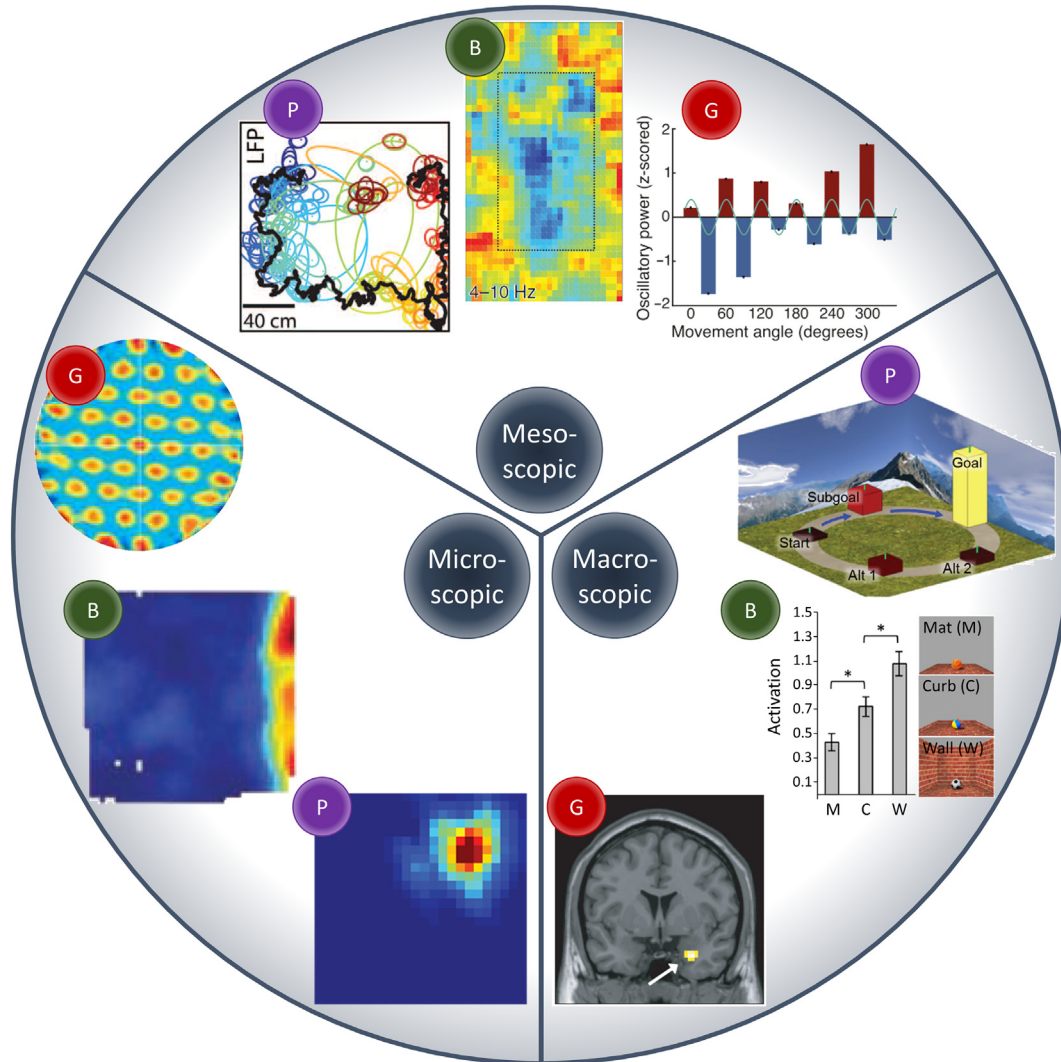
#### Basic Relationships between Spiking Activity, LFPs, and BOLD

Although it is widely agreed that links between spiking activity, LFP power, and the BOLD signal exist (Figure 2), the details of these interrelations remain elusive. Here, we summarize what is currently known and propose that deepening this knowledge will boost our understanding of how spatial codes can coexist at multiple levels of brain organization.

Various studies of the relationship between LFP power and spiking activity revealed that spiking activity correlates positively with broadband power and gamma power, in both human [78] and monkey neocortex [79,80], with high gamma (>80 Hz) power being particularly tightly correlated with spiking activity in some studies [80,81]. Of note, neural synchrony may drive increases in gamma power to a greater extent than firing rate [80]. By contrast, low-frequency oscillations often have a negative relationship to spiking activity, as demonstrated for delta, theta, and alpha power in human and monkey neocortex [78,79]. In certain brain regions, including the hippocampal area, the relationship between LFP power and spiking activity is even more complex. For example, in some studies, spiking activity did not correlate with either broadband, theta-band, or gamma-band LFP power in human hippocampus and EC [82,83]. Such uncoupling between spiking activity and LFPs may be explained by multiple factors, including reduced spike synchronization [82,84], pronounced recurrent processing, and high interneuron density [85,86]. Cell morphology and orientation, topological organization, and stimulus characteristics constitute additional determinants of the relationship between spiking activity and LFP power [87–89]. Notably, the spatial extent of gamma power might sometimes be too local [89] to establish indirect relationships between spiking activity and BOLD (given that gamma power is also associated with BOLD; see later).

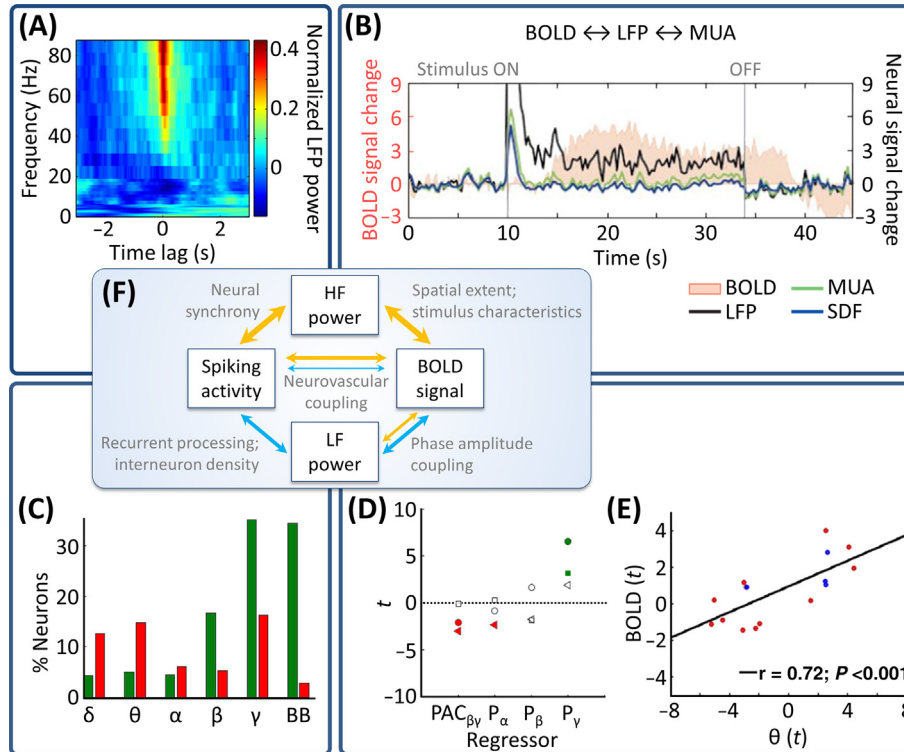
## Key Figure

## Towards a Multilevel Neural Code of Spatial Navigation



Trends In Cognitive Sciences

**Figure 1.** Each third of the pie chart depicts exemplary neural representations of spatial locations (places, P), boundaries (B), and grids (G) at the microscopic, mesoscopic, and macroscopic level of brain organization, respectively. The microscopic (i.e., single-neuron) level shows activation maps of a place cell (P; adapted from [150]); a border cell (B; adapted from [5]); and a grid cell (G; adapted from [6]). Warmer colors correspond to higher firing rates (P and B) or higher autocorrelations (G), respectively. The mesoscopic level depicts hippocampal theta oscillations, the spatiotemporal structure of which allows the prediction (colored eclipses) of the location of a rodent (black line) (P; adapted from [53]); theta power in human subiculum encoding the proximity of goal locations to a spatial boundary (B; adapted from [63]); and entorhinal theta power exhibiting hexadirectional modulation by movement direction (i.e., mesoscopic grid-like representations; G; adapted from [69]). Examples of spatial representations at the macroscopic level include multivariate hippocampal blood oxygen level-dependent (BOLD) patterns that allow the decoding of goal locations (3D bars indicate classifier accuracy; P; adapted from [8]); parahippocampal BOLD signals that encode the presence of boundaries in a spatial scene (B; adapted from [13]); and entorhinal BOLD signals exhibiting hexadirectional modulation by movement direction (i.e., macroscopic grid-like representations; G; adapted from [14]).



Trends in Cognitive Sciences

**Figure 2. Relationships between Spiking Activity, Local Field Potentials (LFPs), and the Blood Oxygen Level-Dependent (BOLD) signal.** (A) Spike-triggered average time-frequency power plot showing that spiking activity is accompanied by increases in high-frequency ('HF') and decreases in low-frequency ('LF') power. Power is normalized across time. (B) Close relationship between the (delayed) BOLD response and LFP HF power, and weaker relationship between BOLD and multiunit activity (MUA): whereas MUA shows a transient response, BOLD and LFPs exhibit a sustained response throughout stimulus presentation. SDF, spike-density function. (C) Spiking activity (in widespread regions including frontal, parietal, and occipital cortices, amygdala, hippocampus, and parahippocampal region) is positively related to broadband ('BB') and HF (gamma) power, but negatively related to LF (delta, theta) power. Green/red bars, percentage of neurons showing a positive/negative relationship, respectively, between spiking activity and LFP power. (D) While beta-gamma phase-amplitude coupling (PAC) and alpha power are negative BOLD predictors, gamma power relates positively to BOLD. Red, significantly negative; green, significantly positive. Each marker indicates one patient. (E) Positive relationship between theta power and parahippocampal BOLD signal. (F) Highly simplified model of the relationship between spiking activity, LFP power, and the BOLD signal. Orange arrows indicate positive relationships, whereas light-blue arrows indicate negative relationships. Thicker lines indicate stronger evidence. Deviations from this model can arise due to various modulating factors (see text annotations for nonexhaustive examples and the main text for details). Adapted from [79] (A), [98] (B), [78] (C), [104] (D), and [83] (E).

Spiking activity is not only associated with oscillatory power, but is also modulated by the phase of neural oscillations, as demonstrated via the phenomena of **phase locking** [90] and **phase precession** [91]. Phase locking reflects the higher excitability of neurons during phase ranges that correspond to more depolarized membrane states. Since both theta [92] and alpha [44] oscillations putatively reflect rhythmically changing levels of inhibition, phase-locked neurons tend to fire during oscillatory troughs, which correspond to lower levels of inhibition. During rodent navigation, phase locking occurs in place cells [93], speed cells [33], grid cells [94], and less strongly in border cells [95]. In human virtual navigation, phase locking to oscillations in the delta, theta, and gamma frequency bands has been observed [96].

Phase precession describes the phenomenon in which a neuron fires at successively earlier phases of an ongoing oscillation, possibly resulting from an interaction of (theta) oscillatory



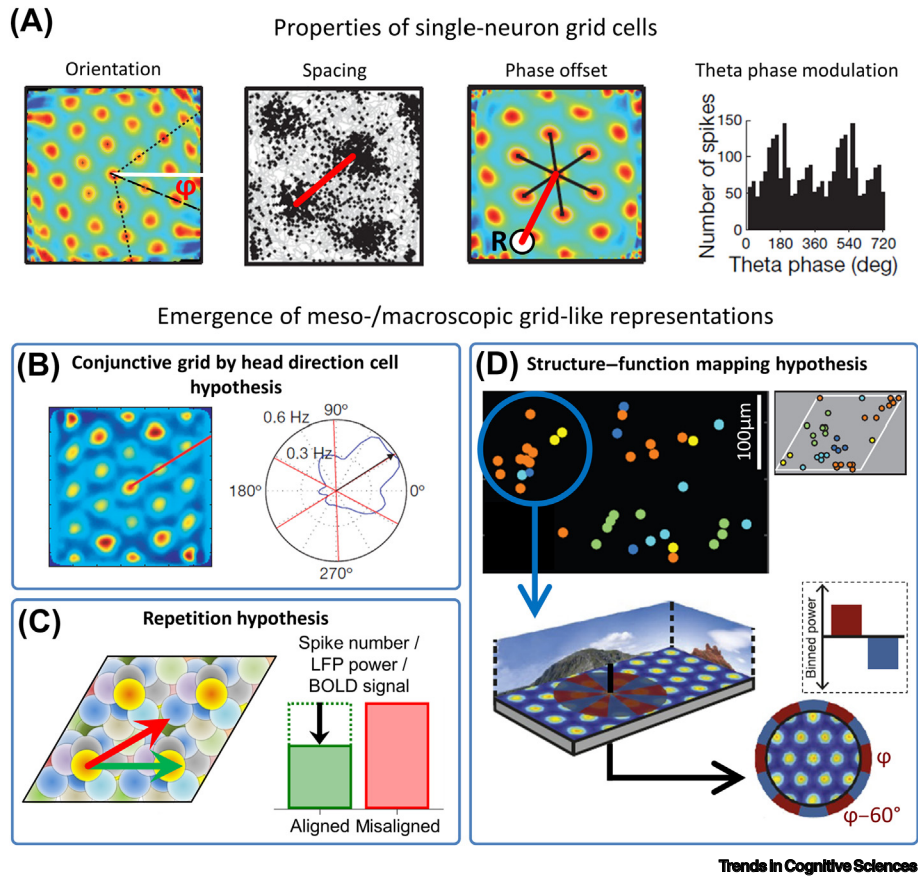
inhibition and an (asymmetric) rate code, which can be induced, for example, when moving through the firing field of a place cell [92]. In rodents, both place [91] and grid cells [94] exhibit phase precession, while evidence for phase precession in humans is still missing. Various functions have been suggested for phase precession, including information processing via a phase code in which information is represented by the oscillatory phase at which a cell or cellular assembly activates [29,92].

Relating electrophysiology and BOLD via simultaneous iEEG/fMRI recordings is challenging for practical and safety reasons [97]. Regarding the relationship between spiking activity and BOLD, simultaneous recordings in sensory neocortex showed that spiking activity and BOLD correlate less tightly than mean LFPs (in particular, gamma-band power) and BOLD [98]. Consecutive and parallel studies in both neocortex and hippocampus revealed variable results ranging from no correlation [83,99] and low degrees of concordance [100] to clear positive relationships between spiking activity and BOLD [101]. Again, divergent findings may be explained by region-specific recurrent processing and interneuron activity that cause dissociations between spiking activity and BOLD. In addition, region-specific neurovascular coupling [85,86] may account for inverse relationships between spiking activity and BOLD in situations when the ratio between blood flow and oxygen metabolism is atypically low [85].

Hence, given these caveats, the BOLD signal is generally thought to correlate more strongly with LFPs than with spiking activity [85,102]: Examining the relationship between high-frequency oscillations and BOLD, simultaneous recordings in monkey visual cortex showed pronounced positive correlations between BOLD and gamma power ([98]; for similar results in various regions and species, see for example, [103,104]). These results were corroborated by consecutive recordings in human neocortex [105], as well as by parallel experiments in the medial temporal lobe of patients with epilepsy and noninvasively in healthy participants [106]. Regarding low-frequency oscillations, simultaneous LFP/BOLD recordings showed negative relationships between BOLD and delta power in rat striatum [107], and between BOLD and alpha/beta power in human motor cortex [104]. Consecutive iEEG/fMRI studies showed similar negative associations between BOLD and beta power, although with pronounced differences between brain regions [108], again in line with the notion that the local circuitry and region-specific neurovascular coupling are modulating factors [85]. In particular, the investigation of theta power and BOLD in the hippocampal formation may lead to variable results, perhaps because this relationship is mediated by the presence or absence of theta/gamma-coupling [104], which depends on ongoing cognitive demands [109]. For example, theta (4–8 Hz) power was found to correlate positively with BOLD in the parahippocampal region (including EC) but not in hippocampus [83].

#### Emergence of Meso- and Macroscopic Spatial Representations

Due to the complex associations between spiking activity, LFPs, and BOLD, it is under intense debate how meso- and macroscopic representations of space relate to the activity of spatially selective single neurons. Potential answers to this question have been suggested mainly for grid cells in the context of spatial representations. Apart from their main feature of firing fields being arranged with sixfold rotational symmetry, these cells exhibit unique functional and anatomical properties (Figure 3A) that may facilitate the emergence of meso-/macroscopic grid-like representations. Specifically, grid cells are organized in a small number of modules [110], with grid cells of the same module exhibiting similar orientation, spacing, and theta modulation [6,110]. In the following paragraphs, we describe four potential explanations for how the features of grid cell-spiking activity relate to hexadirectionally modulated meso-/macroscopic neural signals. All hypotheses assume that, across the grid cell population, the same number of firing fields (centers and peripheries) are traversed along paths of similar distance at any direction.



**Figure 3. Grid Cells and Grid-Like Representations.** (A) Key characteristics of grid cells comprise orientation  $\phi$ , spacing, and phase offset from a reference location (R; adapted from [110]) as well as theta phase modulation (adapted from [94]). (B) Grid-like representations may emerge from firing of conjunctive grid by head-direction cells that exhibit increased firing when moving aligned as compared to misaligned with the grid axes (reprinted from [14]). (C) In a grid cell population with similar orientation and spacing (but with random phases), aligned movements lead to more frequent activation (shorter distance between firing fields) of a smaller number of different grid cells, whereas misaligned movements lead to less frequent activation (larger distance between firing fields) of a higher number of different grid cells. Thus, aligned movements may lead to more pronounced repetition suppression (black arrow) at the level of spiking activity and/or LFP power and/or BOLD signal (fMRI adaptation), resulting in meso- and/or macroscopic grid-like representations. Shaded circles represent firing fields of different grid cells with similar orientation and spacing, but different phase offsets. Each color represents one grid cell. Green arrow indicates aligned movement; red arrow, indicates misaligned movement. (D) Since anatomically adjacent grid cells exhibit similar phase offsets (upper panels adapted from [113]; similar color indicates similar phase offset) in addition to similar orientation and spacing, recording from a limited number of grid cells with a nonrandom distribution of phase offsets leads to grid-like representations (lower panels adapted from [69]).

The ‘conjunctive grid by head-direction cell hypothesis’ (Figure 3B) builds upon prior research that suggested that the firing of conjunctive grid by head-direction cells located in deeper layers of the EC and in pre- and parasubiculum [71] is aligned with the grid axes ([14]; but see [111]). Assuming that the directional tuning width of these conjunctive grid by head-direction cells is not too broad, movements aligned with the grid axes (compared with misaligned movements) will result in increased spiking activity of the conjunctive grid by head-direction cell population, thus causing the appearance of meso-/macroscopic grid-like representations [14]. Of note, (i) the hexadirectional modulation of oscillatory power or BOLD reflects the activity of the population of conjunctive grid by head-direction cells (rather than the grid cells themselves) in this scenario; (ii) such hexadirectional modulation may also occur when the conjunctive grid by

head-direction cells are aligned with the grid axes plus a consistent angular offset; and (iii) pure head-direction cells, the preferred firing directions of which are 60° offset from each other, may also result in hexadirectional modulation of oscillatory power or BOLD.

The ‘repetition hypothesis’ (Figure 3C) is based on the assumption that the phenomenon of repetition suppression (i.e., neural activity being reduced for repeated stimuli [112]) holds true in the context of grid cells [14,72]. Relatively fewer different grid cells are activated more often during aligned movements and relatively more different grid cells are activated less often during misaligned movements. Thus, during aligned movements, there would be a higher degree of repetition suppression at the level of spiking activity (which to the best of our knowledge has yet to be confirmed) and/or LFP power and/or BOLD signal (i.e., fMRI adaptation) compared with misaligned movements, resulting in grid-like representations. Repetition suppression at the level of spiking activity may lead to hexadirectional modulation of LFP power in the theta band, since grid cells exhibit phase locking and phase precession with respect to theta oscillations (see Text S2 and Figure S1 in the supplemental information online).

Regarding the ‘structure–function mapping hypothesis’ (Figure 3D), recent evidence demonstrates that the firing fields of anatomically adjacent grid cells not only have similar spacing and orientation [110], but also a similar phase offset to a reference location [113]. Therefore, recordings from a small area of the EC may sample grid cells with similar firing fields. Such a grid cell population shows higher average firing rates during aligned movements (since more firing fields are traversed) versus misaligned movements, possibly resulting in meso-/macroscopic grid-like representations. Of note, the anatomical area over which adjacent grid cells share a common phase offset may be markedly smaller than the area that is sampled by iEEG or standard fMRI, suggesting that it could be useful to examine fMRI grid-like representations at the high spatial resolution achievable with high-field (7 Tesla) fMRI [114]. Nevertheless, recording from a restricted area of the EC may sample grid cells that are biased towards similar phase offsets, enabling the detection of grid-like representations at lower resolution.

The previous three hypotheses assume consistent (positive or negative) relationships between spiking activity, LFP theta power, and BOLD in EC, an assumption for which there is currently only limited evidence (Figure 2E [83]) and that should be further clarified in future studies (particularly via simultaneous iEEG/fMRI recordings). By contrast, the ‘independence hypothesis’ states that sixfold rotational symmetry occurs independently at different levels of brain organization. This lack of a common neural basis does not exclude a biological and behavioral relevance of meso-/macroscopic grid-like representations, and removes the need for explaining how a limited number of grid cells in medial EC (~10–20% [33,65], see also [115]) can result in a meso-/macroscopically visible neural signal. Of note, all of these hypotheses imply that meso- and macroscopic grid-like representations may also be observable in animal models. Furthermore, combinations of two or more of the hypotheses presented above may hold true, either amplifying or attenuating each other.

#### Redundant or Complementary Information Systems?

Do mesoscopic representations simply mirror the spatial information of single-neuron representations or do they provide independent information? Findings in rodents and humans favor the latter option, but it currently remains elusive whether mesoscopic spatial representations are indeed part of the neural code for spatial navigation or whether they are epiphenomena.

For example, the accuracy of estimating the position of a rat can be improved when information about the LFP phases at which place cells fire is added to the information obtained from place cell firing alone [51]. Similarly, the LFP phases at which a place cell fires and its firing rate can independently represent the location of the animal within the place field and its movement speed through

the field, respectively [52]. Additionally, spike sequences of place cells within theta cycles can be used to predict movement direction [116]. Regarding speed, single-neuron representations and mesoscopic representations may contribute independent information, because they are dissociable from each other analytically [32] and physiologically, via inactivation of the medial septum [35]. Furthermore, phase-specific neuronal firing encodes information about a patient's prospective navigational goal in the absence of firing rate changes [117], and human single-neuron representations of specific passengers or landmarks in a virtual navigation task were shown to be uncorrelated with theta- and gamma-band LFP representations of these stimuli [82]. These studies suggest that micro- and mesoscopic brain signals may carry (partially) independent information and propose that different levels of brain organization make independent contributions to the neural code of spatial navigation. Specifically, as shaped out in this review, mesoscopic theta oscillations may constitute an important partner of microscopic spatial representations. Since several different features relevant to spatial navigation are all represented by theta power, they may jointly shape the functional role of theta for representing and integrating multiple pieces of information.

### Behavioral and Clinical Relevance of Multilevel Spatial Representations

Having described how features of navigation and space relate to micro-, meso-, and macroscopic brain signals, examining their relevance for behavioral performance is important. At the microscopic level, experimentally induced shifts of place fields were associated with impaired spatial behavior [118], increased place field stability correlated with better task performance [119], and disrupting grid cell activity reduced the accuracy of path integration [67]. At the mesoscopic level, abolition of the hippocampal theta rhythm impaired spatial memory [120] and stronger speed modulation of hippocampal theta frequency correlated with better performance [121]. The strength of mesoscopic grid-like representations correlated positively with spatial memory [69] and increased after learning [68]. Similarly, macroscopic grid-like representations in EC and medial prefrontal cortex showed positive associations with memory performance [14–16,70]. Moreover, stronger goal direction signals in human EC correlated with better task accuracy [11]. At the purely behavioral level, grid cell properties predicted biases in human path integration [122]. However, no study demonstrated sixfold rotational symmetry (in contrast to fourfold symmetry [123]) at the behavioral level to date. Future studies are needed to corroborate these findings on how multilevel spatial codes support specific aspects of spatial memory performance (e.g., via optogenetics or deep brain stimulation).

Behavioral relationships between neural representations of space and spatial memory performance suggest that our ability to understand and treat neurological and psychiatric diseases involving spatial disorientation will benefit if we have an improved understanding of the neural representations underlying spatial navigation and how they are disrupted from neuropathologies. Particularly, Alzheimer's disease (AD) may lead to subtle changes in navigational strategies and altered neural representations of space already in very early disease stages [124]. Specific types of spatial representation may be impaired at consecutive AD stages, following either the spreading of neurofibrillary tangles or the development of amyloid  $\beta$  deposition, possibly enabling a 'neurocognitive Braak staging' [125]. AD mouse models have already revealed the adverse impact of tau pathology on grid cell functioning [126] and the degradation of place cell firing due to amyloid plaque burden [127], both pathological changes being accompanied by spatial memory deficits. In addition, AD mouse models identified disrupted phase locking of place cells to hippocampal theta and gamma oscillations [128] as well as reduced cross-frequency coupling of gamma to theta oscillations in medial EC [129]. The investigation of the associations between AD neuropathology and multilevel spatial codes will both enable mechanistic explanations of symptoms and allow the establishment of early biomarkers for AD [124,130,131], such as

reduced grid-like representations [15]. In this respect, establishing meso- and macroscopic representations of space also has practical relevance, because single-neuron spatial representations can be measured from the human brain only under rare circumstances.

### Concluding Remarks

Mesoscopic representations of space based on electrophysiological recordings in humans point towards a multilevel neural code for spatial navigation. These mesoscopic representations exhibit complex connections to spatial representations based on single neurons or on macroscopic fMRI patterns, and may yet mediate the relationship between these two other levels of brain organization. Thus, follow-up studies should further clarify the emergence of meso-/macroscopic spatial representations and show how their information content relates to single-neuron activity. Future research may go beyond the 2D spatial domain and examine to what extent mesoscopic brain signals represent 3D spatial information [132–134], support navigation through other  $n$ -dimensional ‘spaces’ [66,70,77,135–137], and provide biomarkers for neurological and psychiatric diseases (see Outstanding Questions).

### Acknowledgments

L.K. was supported by the Federal Ministry of Education and Research (BMBF; 01GQ1705A), National Science Foundation (NSF) grant BCS-1724243, and National Institutes of Health (NIH) grant 563386. S.M. and J.J. were supported by NIH grants MH061975 and MH104606, and the NSF (BCS-1724243). L.W. was supported by the Strategic Priority Research Program of Chinese Academy of Science (XDB32010300), the Beijing Municipal Science and Technology Commission (Z171100000117014), CAS Interdisciplinary Innovation Team (JCTD-2018-07), and the Natural Science Foundation of China (81422024, 31771255). N.A. received funding by the Deutsche Forschungsgemeinschaft (DFG, German Research Foundation) – Projektnummer 316803389 – SFB 1280 as well as via Projektnummer 122679504 – SFB 874.

### Supplemental Information

Supplemental information associated with this article can be found online at <https://doi.org/10.1016/j.tics.2019.04.011>.

### References

- Chersi, F. and Burgess, N. (2015) The cognitive architecture of spatial navigation: hippocampal and striatal contributions. *Neuron* 88, 64–77
- O’Keefe, J. and Dostrovsky, J. (1971) The hippocampus as a spatial map. Preliminary evidence from unit activity in the freely-moving rat. *Brain Res.* 34, 171–175
- Ekstrom, A.D. et al. (2003) Cellular networks underlying human spatial navigation. *Nature* 425, 184–188
- Taube, J.S. et al. (1990) Head-direction cells recorded from the postsubiculum in freely moving rats. I. Description and quantitative analysis. *J. Neurosci.* 10, 420–435
- Solstad, T. et al. (2008) Representation of geometric borders in the entorhinal cortex. *Science* 322, 1865–1868
- Hafting, T. et al. (2005) Microstructure of a spatial map in the entorhinal cortex. *Nature* 436, 801–806
- Jacobs, J. et al. (2013) Direct recordings of grid-like neuronal activity in human spatial navigation. *Nat. Neurosci.* 16, 1188–1190
- Brown, T.I. et al. (2016) Prospective representation of navigational goals in the human hippocampus. *Science* 352, 1323–1326
- Kyle, C.T. et al. (2015) Successful retrieval of competing spatial environments in humans involves hippocampal pattern separation mechanisms. *eLife* 4, e10499
- Marchette, S.A. et al. (2014) Anchoring the neural compass: coding of local spatial reference frames in human medial parietal lobe. *Nat. Neurosci.* 17, 1598–1606
- Chadwick, M.J. et al. (2015) A goal direction signal in the human entorhinal/subicular region. *Curr. Biol.* 25, 87–92
- Shine, J.P. et al. (2016) The human retrosplenial cortex and thalamus code head direction in a global reference frame. *J. Neurosci.* 36, 6371–6381
- Ferrara, K. and Park, S. (2016) Neural representation of scene boundaries. *Neuropsychologia* 89, 180–190
- Doeller, C.F. et al. (2010) Evidence for grid cells in a human memory network. *Nature* 463, 657–661
- Kunz, L. et al. (2015) Reduced grid-cell-like representations in adults at genetic risk for Alzheimer’s disease. *Science* 350, 430–433
- Stangl, M. et al. (2018) Compromised grid-cell-like representations in old age as a key mechanism to explain age-related navigational deficits. *Curr. Biol.* 28, 1108–1115
- Buzsáki, G. (2002) Theta oscillations in the hippocampus. *Neuron* 33, 325–340
- Ekstrom, A.D. et al. (2005) Human hippocampal theta activity during virtual navigation. *Hippocampus* 15, 881–889
- Watrous, A.J. et al. (2013) A comparative study of human and rat hippocampal low-frequency oscillations during spatial navigation. *Hippocampus* 23, 656–661
- Jacobs, J. (2014) Hippocampal theta oscillations are slower in humans than in rodents: implications for models of spatial navigation and memory. *Philos. Trans. R. Soc. B Biol. Sci.* 369, 20130304
- Buzsáki, G. and Draguhn, A. (2004) Neuronal oscillations in cortical networks. *Science* 304, 1926–1929
- Bohbot, V.D. et al. (2013) Low-frequency theta oscillations in the human hippocampus during real-world and virtual navigation. *Nat. Commun.* 8, 14415
- M Aghajani, Z. et al. (2017) Theta oscillations in the human medial temporal lobe during real-world ambulatory movement. *Curr. Biol.* 27, 3743–3751
- Goyal, A. et al. (2018) Functionally distinct high and low theta oscillations in the human hippocampus. *bioRxiv*. Published online November 17, 2018. <https://doi.org/10.1101/498055>
- Buzsáki, G. et al. (2003) Hippocampal network patterns of activity in the mouse. *Neuroscience* 116, 201–211
- Jacobs, J. et al. (2010) Right-lateralized brain oscillations in human spatial navigation. *J. Cogn. Neurosci.* 22, 824–836

### Outstanding Questions

Can we identify further types of mesoscopic representations of space, such as mesoscopic representations of head direction, borders, or 3D spatial information?

What is the brain-wide distribution of mesoscopic spatial representations, in particular of grid-like representations? Are mesoscopic spatial representations only observable in topographically organized brain regions?

What type of mesoscopic brain signals (which frequencies and which types of features) are causally most relevant for neural representations of spatial navigation?

How do representations of spatial navigation at multiple levels of brain organization relate to each other?

Do neural representations of spatial navigation at different levels of brain organization carry redundant or complementary information?

Will it be possible to simultaneously record LFPs and BOLD in the hippocampus of rodents, nonhuman primates, or patients with epilepsy to reveal the BOLD correlate of hippocampal theta oscillations?

What will new recording techniques, such as wireless EEG, wearable MEG, and chronic invasive implants (which enable neural recordings during naturalistic spatial navigation tasks), contribute to our understanding of neural spatial representations in humans?

Does the disruption of mesoscopic spatial representations lead to impaired spatial memory accuracy? Conversely, can we enhance these representations via neuromodulation to achieve better spatial memory?

Does behavioral spatial navigation performance show sixfold rotational symmetry (e.g., in path integration paradigms)?

How can mesoscopic spatial representations improve our understanding of the interrelations between spatial navigation and visual exploration?



27. Colgin, L.L. *et al.* (2009) Frequency of gamma oscillations routes flow of information in the hippocampus. *Nature* 462, 353–357
28. Canolty, R.T. and Knight, R.T. (2010) The functional role of cross-frequency coupling. *Trends Cogn. Sci.* 14, 506–515
29. Watrous, A.J. *et al.* (2015) More than spikes: common oscillatory mechanisms for content specific neural representations during perception and memory. *Curr. Opin. Neurobiol.* 31, 33–39
30. McFarland, W.L. *et al.* (1975) Relationship between hippocampal theta activity and running speed in the rat. *J. Comp. Physiol. Psychol.* 88, 324–328
31. Ślawińska, U. and Kasicki, S. (1998) The frequency of rat's hippocampal theta rhythm is related to the speed of locomotion. *Brain Res.* 796, 327–331
32. Góis, Z.H.T.D. and Tort, A.B.L. (2018) Characterizing speed cells in the rat hippocampus. *Cell Rep.* 25, 1872–1884.e4
33. Kropff, E. *et al.* (2015) Speed cells in the medial entorhinal cortex. *Nature* 523, 419–424
34. Fuhrmann, F. *et al.* (2015) Locomotion, theta oscillations, and the speed-correlated firing of hippocampal neurons are controlled by a medial septal glutamatergic circuit. *Neuron* 86, 1253–1264
35. Hinman, J.R. *et al.* (2016) Multiple running speed signals in medial entorhinal cortex. *Neuron* 91, 666–679
36. Kraus, B.J. *et al.* (2013) Hippocampal 'time cells': time versus path integration. *Neuron* 78, 1090–1101
37. MacDonald, C.J. *et al.* (2011) Hippocampal 'time cells' bridge the gap in memory for discontinuous events. *Neuron* 71, 737–749
38. Tsao, A. *et al.* (2018) Integrating time from experience in the lateral entorhinal cortex. *Nature* 561, 57–62
39. Thavabalasingam, S. *et al.* (2018) Multivoxel pattern similarity suggests the integration of temporal duration in hippocampal event sequence representations. *Neuroimage* 178, 136–146
40. Montchal, M.E. *et al.* (2019) Precise temporal memories are supported by the lateral entorhinal cortex in humans. *Nat. Neurosci.* 22, 284–288
41. Nakazono, T. *et al.* (2015) Theta oscillation and neuronal activity in rat hippocampus are involved in temporal discrimination of time in seconds. *Front. Syst. Neurosci.* 9, 95
42. Vass, L.K. *et al.* (2016) Oscillations go the distance: low-frequency human hippocampal oscillations code spatial distance in the absence of sensory cues during teleportation. *Neuron* 89, 1180–1186
43. Bush, D. *et al.* (2017) Human hippocampal theta power indicates movement onset and distance travelled. *Proc. Natl. Acad. Sci.* 114, 12297–12302
44. Jensen, O. *et al.* (2014) Temporal coding organized by coupled alpha and gamma oscillations prioritize visual processing. *Trends Neurosci.* 37, 357–369
45. Morgan, L.K. *et al.* (2011) Distances between real-world locations are represented in the human hippocampus. *J. Neurosci.* 31, 1238–1245
46. Howard, L.R. *et al.* (2014) The hippocampus and entorhinal cortex encode the path and euclidean distances to goals during navigation. *Curr. Biol.* 24, 1331–1340
47. Ekstrom, A.D. *et al.* (2017) Interacting networks of brain regions underlie human spatial navigation: a review and novel synthesis of the literature. *J. Neurophysiol.* 118, 3328–3344
48. Diba, K. and Buzsáki, G. (2007) Forward and reverse hippocampal place-cell sequences during ripples. *Nat. Neurosci.* 10, 1241–1242
49. Pfeiffer, B.E. and Foster, D.J. (2013) Hippocampal place-cell sequences depict future paths to remembered goals. *Nature* 497, 74–79
50. Wikenheiser, A.M. and Redish, A.D. (2015) Hippocampal theta sequences reflect current goals. *Nat. Neurosci.* 18, 289–294
51. Jensen, O. and Lisman, J.E. (2000) Position reconstruction from an ensemble of hippocampal place cells: contribution of theta phase coding. *J. Neurophysiol.* 83, 2602–2609
52. Huxter, J. *et al.* (2003) Independent rate and temporal coding in hippocampal pyramidal cells. *Nature* 425, 828–832
53. Aganwal, G. *et al.* (2014) Spatially distributed local fields in the hippocampus encode rat position. *Science* 344, 626–630
54. Hassabis, D. *et al.* (2009) Decoding neuronal ensembles in the human hippocampus. *Curr. Biol.* 19, 546–554
55. Nolan, C.R. *et al.* (2018) Evidence against the detectability of a hippocampal place code using functional magnetic resonance imaging. *eNeuro* 5, ENEURO.0177-18.2018
56. Redish, A.D. *et al.* (2001) Independence of firing correlates of anatomically proximate hippocampal pyramidal cells. *J. Neurosci.* 21, RC134
57. Dombeck, D.A. *et al.* (2010) Functional imaging of hippocampal place cells at cellular resolution during virtual navigation. *Nat. Neurosci.* 13, 1433–1440
58. Redish, A.D. and Ekstrom, A. (2013) Hippocampus and related areas: what the place cell literature tells us about cognitive maps in rats and humans. In *Handbook of Spatial Cognition* (Waller, D. and Nadel, L., eds), pp. 15–34, American Psychological Association
59. Lever, C. *et al.* (2009) Boundary vector cells in the subiculum of the hippocampal formation. *J. Neurosci.* 29, 9771–9777
60. O'Keefe, J. and Burgess, N. (1996) Geometric determinants of the place fields of hippocampal neurons. *Nature* 381, 425–428
61. Hardcastle, K. *et al.* (2015) Environmental boundaries as an error correction mechanism for grid cells. *Neuron* 86, 827–839
62. Julian, J.B. *et al.* (2016) The occipital place area is causally involved in representing environmental boundaries during navigation. *Curr. Biol.* 26, 1104–1109
63. Lee, S.A. *et al.* (2018) Electrophysiological signatures of spatial boundaries in the human subiculum. *J. Neurosci.* 38, 3265–3272
64. Omer, D.B. *et al.* (2018) Social place-cells in the bat hippocampus. *Science* 359, 218–224
65. Diehl, G.W. *et al.* (2017) Grid and nongrid cells in medial entorhinal cortex represent spatial location and environmental features with complementary coding schemes. *Neuron* 94, 83–92
66. Behrens, T.E.J. *et al.* (2018) What is a cognitive map? organizing knowledge for flexible behavior. *Neuron* 100, 490–509
67. Gil, M. *et al.* (2018) Impaired path integration in mice with disrupted grid cell firing. *Nat. Neurosci.* 21, 81–91
68. Chen, D. *et al.* (2018) Hexadirectional modulation of theta power in human entorhinal cortex during spatial navigation. *Curr. Biol.* 28, 3310–3315
69. Maidenbaum, S. *et al.* (2018) Grid-like hexadirectional modulation of human entorhinal theta oscillations. *Proc. Natl. Acad. Sci.* 115, 10798–10803
70. Constantinescu, A.O. *et al.* (2016) Organizing conceptual knowledge in humans with a gridlike code. *Science* 352, 1464–1468
71. Sargolini, F. *et al.* (2006) Conjunctive representation of position, direction, and velocity in entorhinal cortex. *Science* 312, 758–762
72. Killian, N.J. *et al.* (2012) A map of visual space in the primate entorhinal cortex. *Nature* 491, 761–764
73. Julian, J.B. *et al.* (2018) Human entorhinal cortex represents visual space using a boundary-anchored grid. *Nat. Neurosci.* 21, 191–194
74. Nau, M. *et al.* (2018) Hexadirectional coding of visual space in human entorhinal cortex. *Nat. Neurosci.* 21, 188–190
75. Staudigl, T. *et al.* (2018) Hexadirectional modulation of high-frequency electrophysiological activity in the human anterior medial temporal lobe maps visual space. *Curr. Biol.* 28, 3325–3329
76. Kahana, M.J. *et al.* (1999) Human theta oscillations exhibit task dependence during virtual maze navigation. *Nature* 399, 781–784
77. Bellmund, J.L.S. *et al.* (2018) Navigating cognition: spatial codes for human thinking. *Science* 362, eaat6766
78. Manning, J.R. *et al.* (2009) Broadband shifts in local field potential power spectra are correlated with single-neuron spiking in humans. *J. Neurosci.* 29, 13613–13620
79. Rasch, M.J. *et al.* (2008) Inferring spike trains from local field potentials. *J. Neurophysiol.* 99, 1461–1476
80. Ray, S. *et al.* (2008) Neural correlates of high-gamma oscillations (60–200 Hz) in macaque local field potentials and their potential implications in electrocorticography. *J. Neurosci.* 28, 11526–11536

Do mesoscopic brain signals that represent features related to spatial navigation also have a role in 'navigating' *n*-dimensional cognitive 'spaces'?

Can mesoscopic representations of space be used as biomarkers for neurological and psychiatric diseases?

81. Ray, S. and Maunsell, J.H.R. (2011) Different origins of gamma rhythm and high-gamma activity in macaque visual cortex. *PLoS Biol.* 9, e1000610
82. Ekstrom, A. *et al.* (2007) Contrasting roles of neural firing rate and local field potentials in human memory. *Hippocampus* 17, 606–617
83. Ekstrom, A. *et al.* (2009) Correlation between BOLD fMRI and theta-band local field potentials in the human hippocampal area. *J. Neurophysiol.* 101, 2668–2678
84. Nir, Y. *et al.* (2007) Coupling between neuronal firing rate, gamma LFP, and BOLD fMRI is related to interneuronal correlations. *Curr. Biol.* 17, 1275–1285
85. Ekstrom, A. (2010) How and when the fMRI BOLD signal relates to underlying neural activity: The danger in dissociation. *Brain Res. Rev.* 62, 233–244
86. Logothetis, N.K. (2008) What we can do and what we cannot do with fMRI. *Nature* 453, 869–878
87. Buzsáki, G. *et al.* (2012) The origin of extracellular fields and currents — EEG, ECoG, LFP and spikes. *Nat. Rev. Neurosci.* 13, 407–420
88. Pesaran, B. *et al.* (2018) Investigating large-scale brain dynamics using field potential recordings: analysis and interpretation. *Nat. Neurosci.* 21, 903–919
89. Jia, X. *et al.* (2011) Stimulus selectivity and spatial coherence of gamma components of the local field potential. *J. Neurosci.* 31, 9390–9403
90. Buzsáki, G. and Eidelberg, E. (1983) Phase relations of hippocampal projection cells and interneurons to theta activity in the anesthetized rat. *Brain Res.* 266, 334–339
91. O’Keefe, J. and Recce, M.L. (1993) Phase relationship between hippocampal place units and the EEG theta rhythm. *Hippocampus* 3, 317–330
92. Mehta, M.R. *et al.* (2002) Role of experience and oscillations in transforming a rate code into a temporal code. *Nature* 417, 741–746
93. Souza, B.C. and Tort, A.B.L. (2017) Asymmetry of the temporal code for space by hippocampal place cells. *Sci. Rep.* 7, 8507
94. Hafting, T. *et al.* (2008) Hippocampus-independent phase precession in entorhinal grid cells. *Nature* 453, 1248–1252
95. Tang, Q. *et al.* (2014) Pyramidal and stellate cell specificity of grid and border representations in layer 2 of medial entorhinal cortex. *Neuron* 84, 1191–1197
96. Jacobs, J. *et al.* (2007) Brain oscillations control timing of single-neuron activity in humans. *J. Neurosci.* 27, 3839–3844
97. Murta, T. *et al.* (2015) Electrophysiological correlates of the BOLD signal for EEG-informed fMRI. *Hum. Brain Mapp.* 36, 391–414
98. Logothetis, N.K. *et al.* (2001) Neurophysiological investigation of the basis of the fMRI signal. *Nature* 412, 150–157
99. Ojemann, G.A. *et al.* (2013) Relation between functional magnetic resonance imaging (fMRI) and single neuron, local field potential (LFP) and electrocorticography (ECoG) activity in human cortex. *Front. Hum. Neurosci.* 7, 34
100. Kayser, C. *et al.* (2004) A comparison of hemodynamic and neural responses in cat visual cortex using complex stimuli. *Cereb. Cortex* 14, 881–891
101. Mukamel, R. *et al.* (2005) Coupling between neuronal firing, field potentials, and fMRI in human auditory cortex. *Science* 309, 951–954
102. Rauch, A. *et al.* (2008) The effect of a serotonin-induced dissociation between spiking and perisynaptic activity on BOLD functional MRI. *Proc. Natl. Acad. Sci. U. S. A.* 105, 6759–6764
103. Niessing, J. *et al.* (2005) Hemodynamic signals correlate tightly with synchronized gamma oscillations. *Science* 309, 948–951
104. Murta, T. *et al.* (2017) Phase–amplitude coupling and the BOLD signal: a simultaneous intracranial EEG (icEEG) - fMRI study in humans performing a finger-tapping task. *Neuroimage* 146, 438–451
105. Khursheed, F. *et al.* (2011) Frequency-specific electrocorticographic correlates of working memory delay period fMRI activity. *Neuroimage* 56, 1773–1782
106. Axmacher, N. *et al.* (2007) Sustained neural activity patterns during working memory in the human medial temporal lobe. *J. Neurosci.* 27, 7807–7816
107. Jaime, S. *et al.* (2019) Delta rhythm orchestrates the neural activity underlying the resting state BOLD signal via phase–amplitude coupling. *Cereb. Cortex* 29, 119–133
108. Conner, C.R. *et al.* (2011) Variability of the relationship between electrophysiology and BOLD-fMRI across cortical regions in humans. *J. Neurosci.* 31, 12855–12865
109. Axmacher, N. *et al.* (2010) Cross-frequency coupling supports multi-item working memory in the human hippocampus. *Proc. Natl. Acad. Sci. U. S. A.* 107, 3228–3233
110. Stensola, H. *et al.* (2012) The entorhinal grid map is discretized. *Nature* 492, 72–78
111. Keinath, A.T. (2016) The preferred directions of conjunctive grid x head direction cells in the medial entorhinal cortex are periodically organized. *PLoS ONE* 11, e0152041
112. Grill-Spector, K. *et al.* (2006) Repetition and the brain: neural models of stimulus-specific effects. *Trends Cogn. Sci.* 10, 14–23
113. Gu, Y. *et al.* (2018) A map-like micro-organization of grid cells in the medial entorhinal cortex. *Cell* 175, 736–750.e30
114. Schroeder, T.N. *et al.* (2018) Entorhinal cortex minimises uncertainty for optimal behaviour. *bioRxiv*. Published online July 27, 2018. <https://doi.org/10.1101/166306>
115. Boccara, C.N. *et al.* (2010) Grid cells in pre- and parasubiculum. *Nat. Neurosci.* 13, 987–994
116. Huxter, J.R. *et al.* (2008) Theta phase-specific codes for two-dimensional position, trajectory and heading in the hippocampus. *Nat. Neurosci.* 11, 587–594
117. Watrous, A.J. *et al.* (2018) Phase-tuned neuronal firing encodes human contextual representations for navigational goals. *eLife* 7, e32554
118. Lenck-Santini, P.-P. *et al.* (2001) Evidence for a relationship between place-cell spatial firing and spatial memory performance. *Hippocampus* 11, 377–390
119. Kentros, C.G. *et al.* (2004) Increased attention to spatial context increases both place field stability and spatial memory. *Neuron* 42, 283–295
120. Winson, J. (1978) Loss of hippocampal theta rhythm results in spatial memory deficit in the rat. *Science* 201, 160–163
121. Richard, G.R. *et al.* (2013) Speed modulation of hippocampal theta frequency correlates with spatial memory performance. *Hippocampus* 23, 1269–1279
122. Chen, X. *et al.* (2015) Bias in human path integration is predicted by properties of grid cells. *Curr. Biol.* 25, 1771–1776
123. McNamara, T.P. *et al.* (2003) Egocentric and geocentric frames of reference in memory of large-scale space. *Psychon. Bull. Rev.* 10, 589–595
124. Coughlan, G. *et al.* (2018) Spatial navigation deficits — overlooked cognitive marker for preclinical Alzheimer disease? *Nat. Rev. Neurol.* 14, 496–506
125. Braak, H. and Braak, E. (1995) Staging of Alzheimer’s disease-related neurofibrillary changes. *Neurobiol. Aging* 16, 271–278
126. Fu, H. *et al.* (2017) Tau pathology induces excitatory neuron loss, grid cell dysfunction, and spatial memory deficits reminiscent of early Alzheimer’s disease. *Neuron* 93, 533–541
127. Cacucci, F. *et al.* (2008) Place cell firing correlates with memory deficits and amyloid plaque burden in Tg2576 Alzheimer mouse model. *Proc. Natl. Acad. Sci. U. S. A.* 105, 7863–7868
128. Mably, A.J. *et al.* (2017) Impairments in spatial representations and rhythmic coordination of place cells in the 3xTg mouse model of Alzheimer’s disease. *Hippocampus* 27, 378–392
129. Nakazono, T. *et al.* (2017) Impaired in vivo gamma oscillations in the medial entorhinal cortex of knock-in Alzheimer model. *Front. Syst. Neurosci.* 11, 48
130. Vitek, K. and Laczó, J. (2014) Neural correlates of spatial navigation changes in mild cognitive impairment and Alzheimer’s disease. *Front. Behav. Neurosci.* 8, 89
131. Lester, A.W. *et al.* (2017) The aging navigational system. *Neuron* 95, 1019–1035
132. Hayman, R. *et al.* (2011) Anisotropic encoding of three-dimensional space by place cells and grid cells. *Nat. Neurosci.* 14, 1182–1188
133. Yartsev, M.M. and Ulanovsky, N. (2013) Representation of three-dimensional space in the hippocampus of flying bats. *Science* 340, 367–372

134. Kim, M. *et al.* (2017) Multivoxel pattern analysis reveals 3D place information in the human hippocampus. *J. Neurosci.* 37, 4270–4279
135. Theves, S. *et al.* (2019) The hippocampus encodes distances in multidimensional feature space. *Curr. Biol.* 29, 1226–1231
136. Aronov, D. *et al.* (2017) Mapping of a non-spatial dimension by the hippocampal–entorhinal circuit. *Nature* 543, 719–722
137. Bicanski, A. and Burgess, N. (2019) A computational model of visual recognition memory via grid cells. *Curr. Biol.* 29, 979–990
138. Lachaux, J.P. *et al.* (2003) Intracranial EEG and human brain mapping. *J. Physiol.* 97, 613–628
139. Jacobs, J. and Kahana, M.J. (2010) Direct brain recordings fuel advances in cognitive electrophysiology. *Trends Cogn. Sci.* 14, 162–171
140. Norman, K.A. *et al.* (2006) Beyond mind-reading: multi-voxel pattern analysis of fMRI data. *Trends Cogn. Sci.* 10, 424–430
141. Kriegeskorte, N. *et al.* (2008) Representational similarity analysis – connecting the branches of systems neuroscience. *Front. Syst. Neurosci.* 2, 4
142. Miller, J. *et al.* (2018) Lateralized hippocampal oscillations underlie distinct aspects of human spatial memory and navigation. *Nat. Commun.* 9, 2423
143. Colom, L.V. *et al.* (2005) Characterization of medial septal glutamatergic neurons and their projection to the hippocampus. *Synapse* 58, 151–164
144. Frotscher, M. and Léránth, C. (1985) Cholinergic innervation of the rat hippocampus as revealed by choline acetyltransferase immunocytochemistry: a combined light and electron microscopic study. *J. Comp. Neurol.* 239, 237–246
145. Freund, T.F. and Buzsáki, G. (1998) Interneurons of the hippocampus. *Hippocampus* 6, 347–470
146. Jensen, O. (2005) Reading the hippocampal code by theta phase-locking. *Trends Cogn. Sci.* 9, 551–553
147. Lisman, J.E. and Idiart, M.A. (1995) Storage of 7 +/- 2 short-term memories in oscillatory subcycles. *Science* 267, 1512–1515
148. Buzsáki, G. (1989) Two-stage model of memory trace formation: a role for 'noisy' brain states. *Neuroscience* 31, 551–570
149. Huerta, P.T. and Lisman, J.E. (1993) Heightened synaptic plasticity of hippocampal CA1 neurons during a cholinergically induced rhythmic state. *Nature* 364, 723–725
150. Koenig, J. *et al.* (2011) The spatial periodicity of grid cells is not sustained during reduced theta oscillations. *Science* 332, 592–595