Conformal Isometry of Lie Group Representation in Recurrent Network of Grid Cells

Dehong Xu* XUDEHONG1996@UCLA.EDU
Department of Statistics, UCLA

Ruiqi Gao* RUIQIG@GOOGLE.COM
Google Research, Brain Team

Wen-Hao Zhang WENHAO.ZHANG@UTSOUTHWESTERN.EDU
Lyda Hill Department of Bioinformatics and O'Donell Brain Institute, UT Southwestern Medical Center

Xue-Xin Wei WEIXX@UTEXAS.EDU
Departments of Neuroscience and Psychology, Center for Perceptual Systems, Center for Theoretical and Computational Neuroscience, UT Austin

Ying Nian Wu YWU@STAT.UCLA.EDU
Department of Statistics, UCLA

Editors: Sophia Sanborn, Christian Shewmake, Simone Azeglio, Arianna Di Bernardo, Nina Miolane

Abstract

The activity of the grid cell population in the medial entorhinal cortex (MEC) of the mammalian brain forms a vector representation of the self-position of the animal. Recurrent neural networks have been proposed to explain the properties of the grid cells by updating the neural activity vector based on the velocity input of the animal. In doing so, the grid cell system effectively performs path integration. In this paper, we investigate the algebraic, geometric, and topological properties of grid cells using recurrent network models. Algebraically, we study the Lie group and Lie algebra of the recurrent transformation as a representation of self-motion. Geometrically, we study the conformal isometry of the Lie group representation where the local displacement of the activity vector in the neural space is proportional to the local displacement of the agent in the 2D physical space. Topologically, the compact abelian Lie group representation automatically leads to the torus topology commonly assumed and observed in neuroscience. We then focus on a simple non-linear recurrent model that underlies the continuous attractor neural networks of grid cells. Our numerical experiments show that conformal isometry leads to hexagon periodic patterns in the grid cell response and our model is capable of accurate path integration. [1]

Keywords: Grid cells, Conformal isometry, Lie group representation, Lie algebra, Peter-Weyl theory, Flat torus.

1. Introduction

Grid cells (Hafting et al., 2005; Fyhn et al., 2008; Yartsev et al., 2011; Killian et al., 2012; Jacobs et al., 2013; Doeller et al., 2010) in the mammalian dorsal medial entorhinal cortex (MEC) exhibit striking hexagon grid patterns when the agent (e.g., a rodent) navigates in 2-D open environments (Fyhn et al., 2004; Hafting et al., 2005; Fuhs and Touretzky, 2006; Burak and Fiete, 2009; Sreenivasan and Fiete, 2011; Blair et al., 2007; Couey et al., 2013; de Almeida et al., 2009; Pastoll et al., 2013;

* Equal contribution
1. Code is available at https://github.com/DehongXu/grid-cell-rnn

© 2022 D. Xu, R. Gao, W.-H. Zhang, X.-X. Wei & Y.N. Wu.

Agmon and Burak, 2020). It has been hypothesized that grid cell system performs path integration (Darwin, 1873; Etienne and Jeffery, 2004; Hafting et al., 2005; Fiete et al., 2008; McNaughton et al., 2006; Gil et al., 2018; Ridler et al., 2019; Horner et al., 2016). That is, the grid cells integrate self-motion of the animal over time to keep track of the animal's own location in space. This can be implemented by a recurrent neural network that takes the velocity of the self-motion as input, and transforms the activities of the grid cells based on the velocity inputs. The animal's self-position can then be decoded from the activities of the grid cells.

Collectively, the activities of the grid cell population form a vector in the high-dimensional neural activity space. This provides a representation of the self-position of the agent in space. The recurrent network transforms the activity vector based on the movement velocity of the agent, so that the transformation is a representation of self-motion, when considered from the perspective of representational learning. The vector and the transformation together form a representation of the 2D Euclidean group, which is an abelian additive Lie group.

In a recent paper, Gao et al. (2021) studied the group representation property and the isotropic scaling or conformal isometry property for general transformation model. In the context of linear transformation models, they connected this property to the hexagon periodic patterns of the grid cell response maps. With the conformal isometry property of the transformation of the recurrent neural network, the change of the activity vector in the neural space is proportional to the input velocity of the self-motion in the 2D physical space. Gao et al. (2021) justified this condition in terms of robustness to errors or noises in the neurons. Although Gao et al. (2021) studied general transformation model theoretically, they focused on a prototype model of linear recurrent network numerically, which has an explicit algebraic and geometric structure in the form of a matrix group of rotations.

In this paper, we study conformal isometry in the context of the non-linear recurrent model that underlies the hand-crafted continuous attractor neural network (CANN) (Burak and Fiete, 2009; Couey et al., 2013; Pastoll et al., 2013; Agmon and Burak, 2020). In particular, we will focus on the vanilla version of the recurrent network that is linear in the vector representation of self-position and is additive in the input velocity, followed by an element-wise non-linear rectification (such as ReLU nonlinearity). This model has the simplicity that it is additive in input velocity before rectification. We also explore more complex variants for non-linear recurrent network, such as long short-term memory network (LSTM) (Hochreiter and Schmidhuber, 1997). Such models have been studied in recent work (Cueva and Wei, 2018; Banino et al., 2018; Sorscher et al., 2019; Cueva et al., 2020).

Our numerical experiments show that our conformal isometry condition is able to learn highly structured multi-scale hexagon grid code, consistent with the properties of experimentally observed grid cells from rodents. In addition, our learned model is capable of accurate path integration over a long distance. Our results generalize previous results of linear network models in Gao et al. (2021) to an important class of non-linear neural network models in theoretical neuroscience that are more physiologically realistic.

The main contributions of our paper are as follows. (1) We investigate the algebraic, geometric, and topological properties of the transformation models of grid cells. (2) We study a simple non-linear recurrent network that underlies the hand-crafted continuous attractor networks for grid cells, and our numerical experiments suggest that conformal isometry is linked to hexagonal periodic patterns of grid cells.

2. Lie group representation and conformal isometry

2.1. Representations of self-position and self-motion

We start by introducing the basic components of our model. $\boldsymbol{x} = (x_1, x_2) \in \mathbb{R}^2$ denotes the position of the agent. Let $\Delta\boldsymbol{x} = (\Delta x_1, \Delta x_2)$ be the input velocity of the self-motion, i.e., displacement of the agent within a unit time, so that the agent moves from $\boldsymbol{x}$ to $\boldsymbol{x} + \Delta\boldsymbol{x}$ after the unit time.

We assume $\boldsymbol{v}(\boldsymbol{x}) = (v_i(\boldsymbol{x}), i = 1, ..., D)$ to be the vector representation of self-position $\boldsymbol{x}$, where each element $v_i(\boldsymbol{x})$ can be interpreted as the activity of a grid cell when the agent is at position $\boldsymbol{x}$. $(v_i(\boldsymbol{x}), \forall \boldsymbol{x})$ corresponds to the response map of grid cell i. D is the dimensionality of $\boldsymbol{v}$, i.e., the number of grid cells. We refer to the space of $\boldsymbol{v}$ as the "neural space". We normalize $\|\boldsymbol{v}(\boldsymbol{x})\| = 1$ in our experiments.

The set $(\boldsymbol{v}(\boldsymbol{x}), \boldsymbol{x} \in \mathbb{R}^2)$ forms a 2D manifold, or an embedding of $\mathbb{R}^2$, in the D-dimensional neural space. We will refer to $(\boldsymbol{v}(\boldsymbol{x}), \boldsymbol{x} \in \mathbb{R}^2)$ as the "coding manifold".

With self-motion $\Delta\boldsymbol{x}$, the vector representation $\boldsymbol{v}(\boldsymbol{x})$ is transformed to $\boldsymbol{v}(\boldsymbol{x} + \Delta\boldsymbol{x})$ by a general transformation model:

$$\boldsymbol{v}(\boldsymbol{x} + \Delta\boldsymbol{x}) = F(\boldsymbol{v}(\boldsymbol{x}), \Delta\boldsymbol{x}) = F_{\Delta\boldsymbol{x}}(\boldsymbol{v}(\boldsymbol{x})), \tag{1}$$

where by simplifying $F(\cdot, \Delta\boldsymbol{x})$ as $F_{\Delta\boldsymbol{x}}(\cdot)$ in notation, we emphasize that the transformation F is dependent on $\Delta\boldsymbol{x}$. While $\boldsymbol{v}(\boldsymbol{x})$ is a representation of $\boldsymbol{x}$, $F_{\Delta\boldsymbol{x}}$ is a representation of $\Delta\boldsymbol{x}$. $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$ and $(F_{\Delta\boldsymbol{x}}(\cdot), \forall \Delta\boldsymbol{x})$ together form a representation of the 2D additive Euclidean group $\mathbb{R}^2$, which is an abelian Lie group. Specifically, we have the following group representation condition for the transformation model:

Condition 1. *(Algebraic condition on Lie group representation). For any $\boldsymbol{x}$, we have (1) $F_0(\boldsymbol{v}(\boldsymbol{x})) = \boldsymbol{v}(\boldsymbol{x})$, and (2) $F_{\Delta\boldsymbol{x}_1 + \Delta\boldsymbol{x}_2}(\boldsymbol{v}(\boldsymbol{x})) = F_{\Delta\boldsymbol{x}_2}(F_{\Delta\boldsymbol{x}_1}(\boldsymbol{v}(\boldsymbol{x}))) = F_{\Delta\boldsymbol{x}_1}(F_{\Delta\boldsymbol{x}_2}(\boldsymbol{v}(\boldsymbol{x})))$ for any $\Delta\boldsymbol{x}_1$ and $\Delta\boldsymbol{x}_2$.*

Condition 1(1) requires that the coding manifold $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$ are fixed points of F_0 with $\Delta\boldsymbol{x} = 0$. If F_0 is further an contraction off the coding manifold, then $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$ are the attractor points of F_0. Condition 1(2) requires that moving in one step with displacement $\Delta\boldsymbol{x}_1 + \Delta\boldsymbol{x}_2$ should be the same as moving in two steps with displacements $\Delta\boldsymbol{x}_1$ and $\Delta\boldsymbol{x}_2$ respectively. The group representation condition is the necessary condition for any valid transformation model (Equation (1)) of grid cells.

Group representation is a central theme in modern mathematics and physics (Zee, 2016). However, most of the transformations studied in mathematics and physics are linear transformations that form matrix groups, and the coding manifold $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$ is often made implicit. Gao et al. (2019) focused on matrix groups, with $F_{\Delta\boldsymbol{x}}(\boldsymbol{v}(\boldsymbol{x})) = \boldsymbol{M}(\Delta\boldsymbol{x})\boldsymbol{v}(\boldsymbol{x})$, so that $\boldsymbol{M}(\Delta\boldsymbol{x}_1 + \Delta\boldsymbol{x}_2)\boldsymbol{v}(\boldsymbol{x}) = \boldsymbol{M}(\Delta\boldsymbol{x}_1)\boldsymbol{M}(\Delta\boldsymbol{x}_2)\boldsymbol{v}(\boldsymbol{x}) = \boldsymbol{M}(\Delta\boldsymbol{x}_2)\boldsymbol{M}(\Delta\boldsymbol{x}_1)\boldsymbol{v}(\boldsymbol{x})$. Gao et al. (2021) studied general transformation model theoretically, but then focused on linear transformation model in their numerical experiments. Since the transformations in the recurrent neural networks are usually non-linear, we will focus on non-linear transformation models in this paper.

2.2. Conformal embedding and conformal isometry

For an infinitesimal self-motion $\delta\boldsymbol{x}$, it is straightforward to derive a first-order Taylor expansion of the transformation model in Equation (1) with respect to $\delta\boldsymbol{x}$

$$\begin{aligned} \boldsymbol{v}(\boldsymbol{x} + \delta\boldsymbol{x}) &= F_{\boldsymbol{0}}(\boldsymbol{v}(\boldsymbol{x})) + F'_{\boldsymbol{0}}(\boldsymbol{v}(\boldsymbol{x}))\delta\boldsymbol{x} + o(|\delta\boldsymbol{x}|) \\ &= \boldsymbol{v}(\boldsymbol{x}) + f(\boldsymbol{v}(\boldsymbol{x}))\delta\boldsymbol{x} + o(|\delta\boldsymbol{x}|), \end{aligned} \tag{2}$$

where $f(\boldsymbol{v}(\boldsymbol{x})) = \frac{\partial F_{\Delta \boldsymbol{x}}}{\partial \Delta \boldsymbol{x}^\top}(\boldsymbol{v}(\boldsymbol{x}))\,|_{\Delta \boldsymbol{x}=0}$ is a $D \times 2$ matrix.

While $(F_{\Delta \boldsymbol{x}}, \forall \Delta \boldsymbol{x} \in \mathbb{R}^2)$ forms an abelian Lie group, its derivative of $\Delta \boldsymbol{x}$ at 0, i.e., f, spans its Lie algebra. Both $F_{\Delta \boldsymbol{x}}$ and f are transformations acting on the coding manifold $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$.

We identify the conformal isometry condition of the Lie group representation as follows:

Condition 2. *(Geometric condition on conformal embedding and conformal isometry).*

$$f(\boldsymbol{v}(\boldsymbol{x}))^\top f(\boldsymbol{v}(\boldsymbol{x})) = s^2 \boldsymbol{I}_2, \ \forall \boldsymbol{x}, \tag{3}$$

where $\boldsymbol{I}_2$ is the 2-dimensional identity matrix. That is, the two column vectors of $f(\boldsymbol{v}(\boldsymbol{x}))$ are of equal norm s, and are orthogonal to each other.

Under the condition above, $\boldsymbol{v}(\boldsymbol{x}+\delta\boldsymbol{x}) - \boldsymbol{v}(\boldsymbol{x}) \approx f(\boldsymbol{v}(\boldsymbol{x}))\delta\boldsymbol{x}$ is conformal to $\delta\boldsymbol{x}$, i.e., the 2D local Euclidean space of $(\delta\boldsymbol{x})$ in the physical space is embedded conformally as another 2D local Euclidean space $f(\boldsymbol{v}(\boldsymbol{x}))\delta\boldsymbol{x}$ in the neural activity space. We only need to replace the two orthogonal axes for $\delta\boldsymbol{x}$ in the 2D physical space by the two column vectors of $f(\boldsymbol{v}(\boldsymbol{x}))$ in the neural activity space.

An equivalent statement for the above condition is

$$\|\boldsymbol{v}(\boldsymbol{x}+\delta\boldsymbol{x}) - \boldsymbol{v}(\boldsymbol{x})\| = s\|\delta\boldsymbol{x}\| + o(\|\delta\boldsymbol{x}\|), \forall \boldsymbol{x}, \delta\boldsymbol{x}. \tag{4}$$

That is, the displacement in the neural space is proportional to the displacement in the 2D physical space.

Note that since our analysis is local, s may depend on $\boldsymbol{x}$. If s is a global constant, then the coding manifold $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$ has a flat intrinsic geometry (imagining folding a piece of paper without stretching it).

Gao et al. (2021) studied the conformal isometry property in a local polar coordinate system. In our definition above, we use 2D cartesian coordinate system, which is more convenient for the non-linear recurrent model where Δx enters the model additively. In the context of linear recurrent network, Gao et al. (2021) linked the conformal isometry to the hexagon grid patterns of grid cells.

2.3. 2D torus, 2D periodicity, and hexagon grid patterns

The 2D torus topology is commonly assumed *a priori* in the continuous attractor neural networks (CANN) for grid cells (Burak and Fiete, 2009; Couey et al., 2013; Pastoll et al., 2013; Agmon and Burak, 2020). The torus topology has been recently supported by analyzing data from population of simultaneously recorded grid cells (Gardner et al., 2022). Within our framework, we find that such a topology is in fact a theoretical consequence of the group representation condition (Condition 1) due to a theorem in Lie group theory.

Specifically, since the elements of $\boldsymbol{v}(\boldsymbol{x})$ represent neuron activities, which are bounded biologically, the coding manifold $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$ is bounded and compact, and therefore the group of $(F_{\Delta \boldsymbol{x}}, \forall \Delta \boldsymbol{x})$ is also compact. The Lie group $(F_{\Delta \boldsymbol{x}}, \forall \Delta \boldsymbol{x})$ is also connected since the 2D environment is connected. According to Lie group theory (Dwyer and Wilkerson, 1998), if the abelian Lie group $(F_{\Delta \boldsymbol{x}}, \forall \Delta \boldsymbol{x})$ is compact and connected, then it has a topology of 2D torus, i.e., it is isomorphic to $\mathbb{S}_1 \times \mathbb{S}_1$, where $\mathbb{S}_1$ is a circle. Thus $(\boldsymbol{v}(\boldsymbol{x}) = F_{\boldsymbol{x}}(\boldsymbol{v}(0)), \forall \boldsymbol{x} \in \mathbb{R}^2)$ also forms a 2D torus.

While the group representation condition only gives us an algebraic structure, the conformal isometry condition (Condition 2) further fixes the geometry. Under the conformal isometry condition,

if we further assume that the scaling factor s is a constant globally for all $\boldsymbol{x}$, then the intrinsic geometry of the coding manifold $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$ remains Euclidean, and the coding manifold is a flat torus. That is, the coding manifold is not only isomorphic to $\mathbb{S}_1 \times \mathbb{S}_1$, but is also conformally isometric to $\mathbb{S}_1 \times \mathbb{S}_1$. While isomorphism is defined by mapping between two spaces, isometry concerns about the metric properties. This leads to the periodic pattern in $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$ over $\boldsymbol{x}$

According to the theory of 2D Bravais lattice for 2D periodic patterns (Ashcroft et al., 1976), we can find two primitive vectors $\Delta\boldsymbol{x}_1$ and $\Delta\boldsymbol{x}_2$, with $\|\Delta\boldsymbol{x}_1\| = \|\Delta\boldsymbol{x}_2\|$, and $\boldsymbol{v}(\boldsymbol{x} + k_1\Delta\boldsymbol{x}_1 + k_2\Delta\boldsymbol{x}_2) = \boldsymbol{v}(\boldsymbol{x})$ for arbitrary integers k_1 and k_2. Along each primitive vector, for each period, $(\boldsymbol{v}(\boldsymbol{x} + c\Delta\boldsymbol{x}_i), c \in [0,1])$ traces out a circle in the neural space for $i = 1, 2$, causing the periodicity in $\boldsymbol{v}(\boldsymbol{x})$. According to the theory of 2D Bravais lattice, the angle between $\Delta\boldsymbol{x}_1$ and $\Delta\boldsymbol{x}_2$ can either be $\pi/2$ for square lattice or $2\pi/3$ for hexagon lattice. It is likely that the hexagon periodicity provides a better fit to place cells, in that the hexagon lattice provides denser packing of discrete Fourier components. While this seems to be intuitive, currently we have not been able to prove this point rigorously. It seems that hexagonal periodicity emerges under the general conditions of group representation and conformal isometry, independent of a specific form of the transformation model, as we observe empirically in our numerical experiments (Section 5).

3. Non-linear recurrent neural network

3.1. Model

In this paper, we mainly focus on studying transformations $F_{\Delta\boldsymbol{x}}(\cdot)$ that are locally approximated by non-linear recurrent neural networks, as studied in recent work (Cueva and Wei, 2018; Banino et al., 2018; Sorscher et al., 2019; Cueva et al., 2020). We start by assuming the following vanilla version of non-linear recurrent network:

$$\boldsymbol{v}(\boldsymbol{x} + \Delta\boldsymbol{x}) = \text{ReLU}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) + \boldsymbol{U}\Delta\boldsymbol{x}), \tag{5}$$

where $\text{ReLU}(a) = \max(0, a)$ is applied element-wise, $\boldsymbol{W}$ is a $D \times D$ weight matrix of recurrent connections, $\boldsymbol{U}$ is a $D \times 2$ matrix, and the self-motion $\Delta\boldsymbol{x} = (\Delta x_1, \Delta x_2)^\top$ is treated as 2×1 vector. Note that the above model is an accurate approximation to $F_{\Delta\boldsymbol{x}}(\cdot)$ only for small $\Delta\boldsymbol{x}$, as it may not satisfy Condition 1 in general for large $\Delta\boldsymbol{x}$. Following Condition 1(1), for $\Delta\boldsymbol{x} = 0$ we have $\boldsymbol{v}(\boldsymbol{x}) = \text{ReLU}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}))$. That is, the coding manifold $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$ consists of the fixed points of $F_0(\cdot)$.

Compared to the matrix group in Gao et al. (2019) where $F_{\Delta\boldsymbol{x}}(\boldsymbol{v}(\boldsymbol{x})) = \boldsymbol{M}(\Delta\boldsymbol{x})\boldsymbol{v}(\boldsymbol{x})$, and $\boldsymbol{M}(\Delta\boldsymbol{x})$ is further derived in Gao et al. (2021) as an exponential map that is highly non-linear in $\Delta\boldsymbol{x}$, the above model (5) is much simpler and more biologically plausible in that, before ReLU, we have a single recurrent weight matrix $\boldsymbol{W}$ that is independent of $\Delta\boldsymbol{x}$, and $\Delta\boldsymbol{x}$ enters the equation additively. The ReLU rectification plays a critical role for the overall non-linear effect of $\Delta\boldsymbol{x}$. The non-linear rectification in neural networks makes the transformations much more expressive than matrix representations in modern math and physics.

For this model, we can derive $f(\boldsymbol{v}(\boldsymbol{x}))$ as

$$f(\boldsymbol{v}(\boldsymbol{x})) = \mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) > \mathbf{0}) \odot \boldsymbol{U}, \tag{6}$$

where $\mathbf{1}(\cdot)$ is a vector of binary indicators calculated element-wise, and $\odot$ is row-wise product. The indicator vector $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) > \mathbf{0})$ changes as $\boldsymbol{x}$ changes, and it controls the change of $\boldsymbol{v}$ in the neural space according to $\Delta\boldsymbol{x}$.

Since $\mathbf{1}(\cdot)$ is not differentiable, we propose to define the loss function to enforce the conformal isometry condition based on the equivalent statement in Equation (4), as discussed in Section 4.3. In Appendix C, we also discuss a possible mechanism where conform isometry can be automatically satisfied by design.

Modules. Since it is well-known that biological grid cells are organized in discrete modules with different spatial scales (Stensola et al., 2012; Barry et al., 2007), we assume that in our model, the vector representations $\boldsymbol{v}(\boldsymbol{x})$ are divided into sub-vectors analogous to modules. Accordingly, we assume that the transformation is also module-wise by assuming $\boldsymbol{W}$ is block-diagonal and $\boldsymbol{U}$ is divided into sub-blocks by row. To address the point that the module-wise transformation construction is just to keep the consistency with isometry assumption but not the reason of the emergence of hexagonal periodicity, we have performed an ablation study (see Appendix A.3).

More complex transformations. We also explore the Long Short-Term Memory network (LSTM) (Hochreiter and Schmidhuber, 1997), a more complex variant of recurrent network, as the local transformation model. In the experiments, we observed that hexagonal grid patterns can emerge in both types of transformation models, implying that the emergence of hexagonal periodicity under the conformal isometry condition is not specific to the particular form of the transformation model, at least empirically.

3.2. Eigen analysis

Next, we will deepen our theoretical understanding of the model (5) by conducting eigen analysis under the conformal isometry condition.

Theorem 1. *Under conformal isometry condition, for every $\boldsymbol{x}$, the two columns of $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) > \mathbf{0}) \odot \boldsymbol{U}$ are orthogonal, and they are eigenvectors of $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) > \mathbf{0}) \odot \boldsymbol{W}$ with eigenvalue 1.*

See the Appendix for proof. If the magnitudes of all the other eigenvalues of $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) > \mathbf{0}) \odot \boldsymbol{W}$ are less than 1, the recurrent network is a contraction off the coding manifold, which is similar to the eigen structures of general continuous attractor neural networks as shown in Fung et al. (2010).

4. Reconstructing place cells and learning by numerical optimization

4.1. Place cells and decoding

For open fields, we can model place cells (O'Keefe, 1979) by Gaussian kernels and connect them to grid cells by the following basis expansion model (Dordek et al., 2016; Sorscher et al., 2019):

$$A(\boldsymbol{x}, \boldsymbol{p}) = \exp(-\|\boldsymbol{x} - \boldsymbol{p}\|^2/2\sigma^2) = \langle \boldsymbol{v}(\boldsymbol{x}), \boldsymbol{q}(\boldsymbol{p}) \rangle, \tag{7}$$

where $A(\boldsymbol{x}, \boldsymbol{p})$ represent the place field centered at position $\boldsymbol{p}$. $A(\boldsymbol{x}, \boldsymbol{p})$ measures the adjacency of $\boldsymbol{x}$ to $\boldsymbol{p}$. $\boldsymbol{q}(\boldsymbol{p})$ is the query vector of the place cell $\boldsymbol{p}$, which can be interpreted as connection weights between the grid cells and the place cell $\boldsymbol{p}$. For a vector $\boldsymbol{v}$, we can decode its position $\hat{\boldsymbol{x}}$ by

$$\hat{\boldsymbol{x}} = \arg\max_{\boldsymbol{p}} \langle \boldsymbol{v}, \boldsymbol{q}(\boldsymbol{p}) \rangle, \tag{8}$$

i.e., we choose the position $\boldsymbol{p}$ that is closest to the position encoded by $\boldsymbol{v}$.

In our experiments, we learn the query vectors $(\boldsymbol{q}(\boldsymbol{p}), \forall \boldsymbol{p})$ together with $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$, $\boldsymbol{W}$, $\boldsymbol{U}$.

4.2. Peter-Weyl theory

In model defined by Eq. (7), $\boldsymbol{v}(\boldsymbol{x})$ generated by the transformation model serves as a set of basis functions to reconstruct the set of functions $A(\boldsymbol{x},\boldsymbol{p}), \forall \boldsymbol{p}$. This is related to the Peter-Weyl theory (Taylor, 2002), which generalizes the Fourier analysis to Lie group and shows that the basis functions arise from Lie group representation. $\boldsymbol{v}(\boldsymbol{x})$ can be used to reconstruct and interpolate value functions of $\boldsymbol{x}$ in general. Peter-Weyl theory is about matrix Lie groups and irreducible representations. In our work, we focus on non-linear transformation groups which can still generate basis functions.

Peter-Weyl theory naturally connects two roles of grid cells: (1) path integration, and (2) basis expansion. It can be interesting to generalize Peter-Weyl theory to non-linear transformations.

4.3. Loss function

Assuming the kernel $A(\boldsymbol{x},\boldsymbol{p})$ is given as in Equation (7), we can learn the model by minimizing the following loss term:

$$L_1 = \sum_{t=1}^{T} \sum_{\boldsymbol{p}} \mathbb{E}_{\boldsymbol{x},\Delta\boldsymbol{x}} [A(\boldsymbol{x} + \Delta\boldsymbol{x}_1 + ... + \Delta\boldsymbol{x}_t, \boldsymbol{p}) - \langle F_{\Delta\boldsymbol{x}_t} ... F_{\Delta\boldsymbol{x}_1}(\boldsymbol{v}(\boldsymbol{x})), \boldsymbol{q}(\boldsymbol{p})\rangle]^2. \tag{9}$$

The learnable parameters includes $(\boldsymbol{v}(\boldsymbol{x}), \forall \boldsymbol{x})$, $(\boldsymbol{q}(\boldsymbol{p}), \forall \boldsymbol{p})$, and parameters in $F_{\Delta\boldsymbol{x}}$. The expectations are estimated by Monte Carlo samples from simulated trajectories. $A(\boldsymbol{x},\boldsymbol{p})$ are Gaussian kernels with predefined σ. In practice, we add an additional zero-step version of L_1, i.e. the expectation term changes to $[A(\boldsymbol{x},\boldsymbol{p}) - \langle \boldsymbol{v}(\boldsymbol{x}), \boldsymbol{q}(\boldsymbol{p})\rangle]^2$.

To ensure that the conformal isometry condition is satisfied, we add an extra loss term based on the equivalent statement of conformal isometry. Following Equation (4), for simplicity, we first denote $\boldsymbol{s}(\boldsymbol{x},\Delta\boldsymbol{x}) = (\|\boldsymbol{v}(\boldsymbol{x}) - \boldsymbol{v}(\boldsymbol{x}+\Delta\boldsymbol{x})\| / \|\Delta\boldsymbol{x}\|)^2$. Then we propose a conformal isometry loss:

$$L_2 = \mathbb{E}_{\boldsymbol{x},\Delta\boldsymbol{x}_1,\Delta\boldsymbol{x}_2} [\boldsymbol{s}(\boldsymbol{x},\Delta\boldsymbol{x}_1) - \boldsymbol{s}(\boldsymbol{x},\Delta\boldsymbol{x}_2)]^2, \tag{10}$$

where $\Delta\boldsymbol{x}_1$ and $\Delta\boldsymbol{x}_2$ are sampled such that they have the same length Δr but with different directions, i.e. $\Delta\boldsymbol{x}_1 = (\Delta r \cos\theta_1, \Delta r \sin\theta_1)$, $\Delta\boldsymbol{x}_2 = (\Delta r \cos\theta_2, \Delta r \sin\theta_2)$. Moreover, we add another regularization term to penalize $\|\boldsymbol{q}(\boldsymbol{p})\|^2$.

5. Experiments

We optimized the model using simulated trajectories as training data. The environment was assumed to be a 1m $\times$ 1m squared open field, discretized into a 40×40 lattice. $\boldsymbol{v}(\boldsymbol{x})$ is of 1800 dimensions, which was partitioned into 150 modules with module size 12. For $A(\boldsymbol{x},\boldsymbol{p})$, we used a Gaussian adjacency kernel with $\sigma = 0.07$. We trained a 10-step recurrent network as the transformation model, i.e., $T = 10$ in the loss term L_1.

For L_1, the displacement of $\Delta\boldsymbol{x}_t$ was restricted to be smaller than 3 grids. For the range of Δr in L_2, we hypothesize that it can be proportional to the scale of the module (i.e., s in Equation (4)), which is also reflected as the scale of the learned hexagon patterns. Thus we adaptively adjusted the upper bound of Δr for each module during training, based on the scale of the learned hexagon patterns at the current training stage. We averaged the scales of the learned patterns within each module to represent the scale of that module. We set the upper bound of Δr for the module with the largest average scale to be 15 grids. The ranges of Δr for the remaining modules are adjusted according to their scales.

5.1. Hexagon patterns

Figure 1 shows the learned firing patterns of $\boldsymbol{v}(\boldsymbol{x}) = (v_i(\boldsymbol{x}), i = 1, ..., d)$ over the 40×40 lattice of $\boldsymbol{x}$. We randomly selected 16 modules out of 150 modules for visualization purposes. Each image corresponds to the response map of a grid cell. Every row shows the learned units that belong to the same module. The hexagonal patterns in the emerging activity patterns are evident. We found that the loss term for imposing the conformal isometry was critical. Without it, the learned response maps showed stripe-like patterns (see Appendix for ablation results).

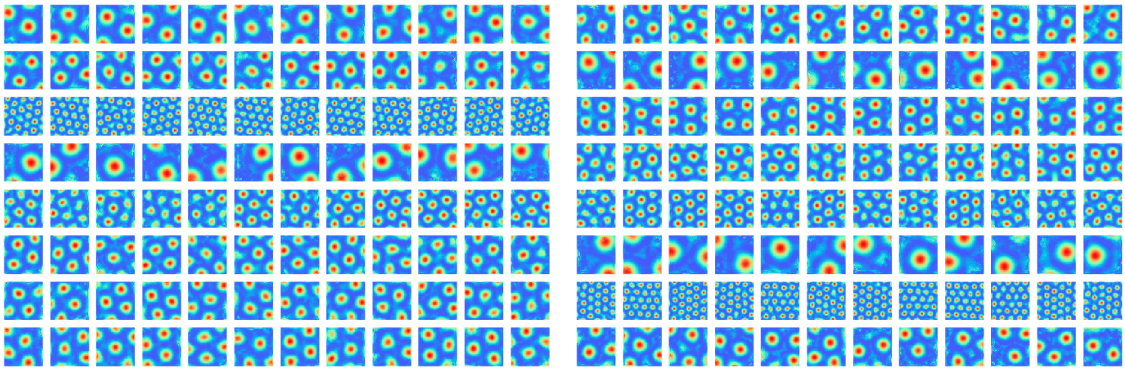


Figure 1: Hexagon grid firing patterns emerge in the learned $\boldsymbol{v}(\boldsymbol{x})$. Each row represents the firing patterns of all the cells in the same module. Each unit shows the learned neuron activity over the whole 2D squared environment. The figure shows patterns from 16 randomly selected modules.

To quantitatively evaluate whether the learned patterns match regular hexagon grids, in Table 1, we report the gridness scores (Langston et al. (2010); Sargolini et al. (2006)) that are adopted from the literature of grid cells, as well as the valid percentage of grid cells with gridness score > 0.37 being the criteria. Figure 2 shows the histogram of the spatial scales of the learned hexagon patterns. The multi-modal distribution is fitted by a mixture of three Gaussians, which are centered at 0.41, 0.64 and 0.86 respectively. The ratios between adjacent centers are 1.55 and 1.35, which are in the range of the data from rodent grid cells (Stensola et al., 2012).

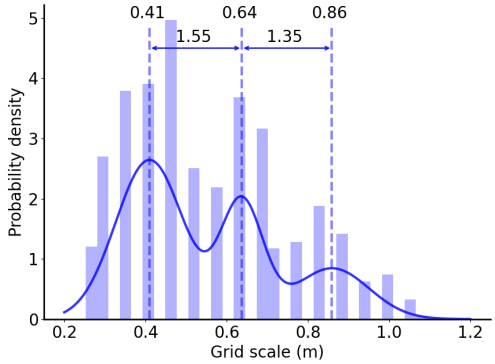


Figure 2: Histogram of grid scales of the learned model.

Table 1: Gridness scores and valid rates of grid cells of learned models.

Model	Gridness score (↑)	% of grid cells (↑)
Banino et al. (2018) (LSTM)	0.18	25.20
Sorscher et al. (2019) (RNN)	0.48	56.10
Ours (RNN)	**0.77**	**72.5**
Ours (LSTM)	**0.73**	**68.8**

5.2. Path integration

We further evaluate whether the learned model is capable of accurate path integration. We perform path integration in two scenarios. First, for path integration with re-encoding, we decode $\boldsymbol{v}_t \to \boldsymbol{x}_t$ to physical space and then apply encoder $\boldsymbol{v}(\boldsymbol{x}_t) \to \boldsymbol{v}'_t$ back to neuron space every few steps. This re-encoding strategy helps correct the errors accumulated in the neural space along the transformation. In the without re-encoding case, we apply transformation purely using neuron vector $\boldsymbol{v}_t$. As shown in the left panel of Figure 3, the model can perform near exact path integration for 30 steps (short distance) without re-encoding. For long-distance path integration, we train a 20-step recurrent network model, and evaluate the model for 500 steps over 1000 trajectories. As shown in the right subfigure of Figure 3, if we re-encode every 20 steps, the path integration error for the last step is 0.028, while the average error over the 500 steps trajectory is 0.017. Without re-encoding, the error is relatively larger, where the average error is around 0.03 along the whole trajectory, and 0.08 for the last step.

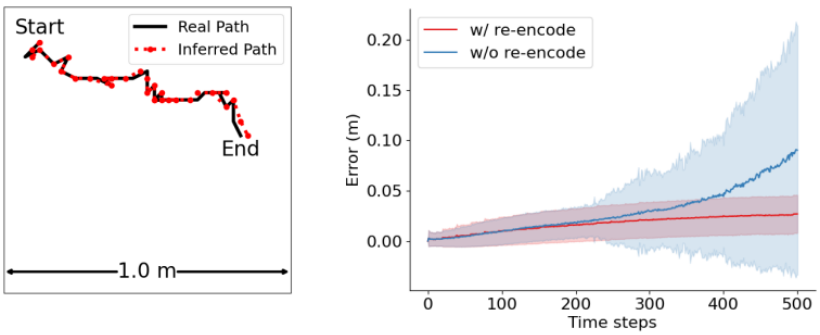


Figure 3: The learned model can perform accurate path integration. *Left*: path integration for 30 steps without re-encoding. Black: ground truth. Red: the inferred path by the learned model. *Right*: long distance (500-step) path integration error with (red) and without (blue) re-encoding by a learned 20-step RNN model over time steps. The average error and standard deviation are evaluated over 1000 trajectories.

To check if the model can apply precise encoding and decoding between physical space and neuron space, we also examine the fixed point condition by applying $\boldsymbol{v}(\boldsymbol{x}) \to \boldsymbol{v}_t \to \boldsymbol{x}'$. Ideally, the learned model can figure out the physical location $\boldsymbol{x}_t$ purely from $\boldsymbol{v}_t$. The L_2 error between $\boldsymbol{x}$ and $\boldsymbol{x}'$ is nearly zero (< 0.005).

5.3. Model with LSTM units

In this section, we evaluate the LSTM transformation model. The model is still trained by the same loss functions as the vanilla RNN model. Meanwhile, we force $\boldsymbol{q}(\boldsymbol{p}) > 0$ in training.

Examples of the learned patterns are visualized in Figure 4. Clear hexagon patterns are also evident. Different from the learned units from the vanilla recurrent network which are all non-negative, the learned $\boldsymbol{v}(\boldsymbol{x})$ from the LSTM model can be either positive or negative, resulting in the color shift of the learned patterns. As shown in Table 1, the average gridness score for LSTM model is 0.73, and 68.8% of the model units are classified as grid cells. We also evaluated the LSTM model on path integration still using 1000 trajectories and each trajecture is 500 steps long. With re-encoding every 10 steps, the average decoding error over the whole trajectory was 0.027, and the error at the 500th step still remained as low as 0.037.

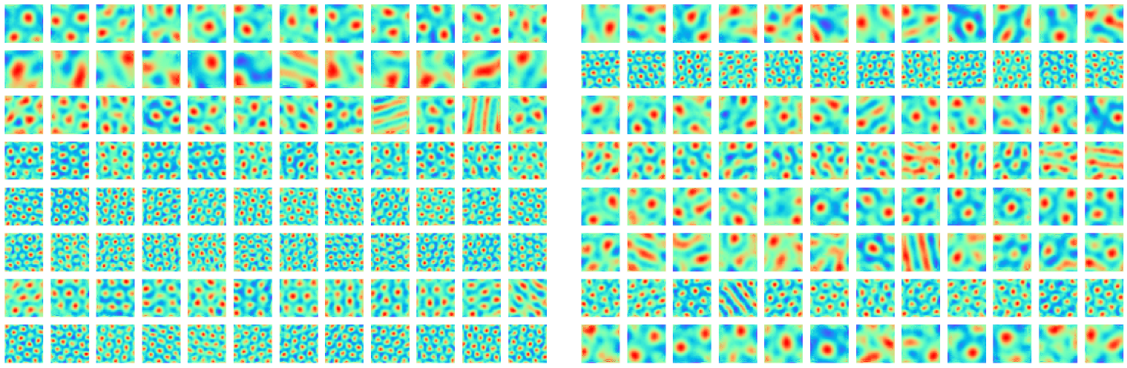


Figure 4: Learned hexagon grid patterns of $\boldsymbol{v}(\boldsymbol{x})$, which is the hidden state vector in the LSTM transformation model. For each row, it shows all the cells in the same module and 16 modules are randomly selected and visualized.

6. Conclusion and discussion

This paper investigates the algebraic, geometric, and topological properties of the generic transformation model of grid cells. We focus on non-linear recurrent neural networks under the conformal isometry condition. Our numerical experiments demonstrated that hexagon periodic patterns emerged in the response maps of grid cells under the conformal isometry condition. Our experiments also showed that the learned model was capable of accurate path integration.

The conformal isometry property seems to be related to the difference of Gaussian kernel assumed for place cells in Dordek et al. (2016); Sorscher et al. (2019), which constrains the frequency components within a ring in the frequency domain. It remains to be determined whether the hexagon patterns of grid cells were caused by the recurrent network itself or by interaction with place cells, an important issue that should be investigated further.

A limitation of our work is that we did not pursue specially designed recurrent neural networks where conformal isometry is automatically satisfied. We leave it to future investigation. In the Appendix, we provide a discussion of a possible design inspired by the hand-crafted continuous attractor neural networks of grid cells where conformal isometry is satisfied.

Acknowledgments

The work was supported by NSF DMS-2015577 and XSEDE grant ASC170063. We thank the reviewers for their valuable comments and suggestions.

References

Haggai Agmon and Yoram Burak. A theory of joint attractor dynamics in the hippocampus and the entorhinal cortex accounts for artificial remapping and grid cell field-to-field variability. *eLife*, 9: e56894, 2020.

Neil W Ashcroft, N David Mermin, et al. Solid state physics, 1976.

Andrea Banino, Caswell Barry, Benigno Uria, Charles Blundell, Timothy Lillicrap, Piotr Mirowski, Alexander Pritzel, Martin J Chadwick, Thomas Degris, Joseph Modayil, et al. Vector-based navigation using grid-like representations in artificial agents. *Nature*, 557(7705):429, 2018.

Caswell Barry, Robin Hayman, Neil Burgess, and Kathryn J Jeffery. Experience-dependent rescaling of entorhinal grids. *Nature neuroscience*, 10(6):682–684, 2007.

Hugh T Blair, Adam C Welday, and Kechen Zhang. Scale-invariant memory representations emerge from moire interference between grid fields that produce theta oscillations: a computational model. *Journal of Neuroscience*, 27(12):3211–3229, 2007.

Yoram Burak and Ila R Fiete. Accurate path integration in continuous attractor network models of grid cells. *PLoS computational biology*, 5(2):e1000291, 2009.

Jonathan J Couey, Aree Witoelar, Sheng-Jia Zhang, Kang Zheng, Jing Ye, Benjamin Dunn, Rafal Czajkowski, May-Britt Moser, Edvard I Moser, Yasser Roudi, et al. Recurrent inhibitory circuitry as a mechanism for grid formation. *Nature neuroscience*, 16(3):318–324, 2013.

Christopher J Cueva and Xue-Xin Wei. Emergence of grid-like representations by training recurrent neural networks to perform spatial localization. *arXiv preprint arXiv:1803.07770*, 2018.

Christopher J Cueva, Peter Y Wang, Matthew Chin, and Xue-Xin Wei. Emergence of functional and structural properties of the head direction system by optimization of recurrent neural networks. *International Conferences on Learning Representations (ICLR)*, 2020.

Charles Darwin. Origin of certain instincts, 1873.

Licurgo de Almeida, Marco Idiart, and John E Lisman. The input–output transformation of the hippocampal granule cells: from grid cells to place fields. *Journal of Neuroscience*, 29(23): 7504–7512, 2009.

Christian F Doeller, Caswell Barry, and Neil Burgess. Evidence for grid cells in a human memory network. *Nature*, 463(7281):657, 2010.

Yedidyah Dordek, Daniel Soudry, Ron Meir, and Dori Derdikman. Extracting grid cell characteristics from place cell inputs using non-negative principal component analysis. *Elife*, 5:e10094, 2016.

William Gerard Dwyer and CW Wilkerson. The elementary geometric structure of compact lie groups. *Bulletin of the London Mathematical Society*, 30(4):337–364, 1998.

Ariane S Etienne and Kathryn J Jeffery. Path integration in mammals. *Hippocampus*, 14(2):180–192, 2004.

Ila R Fiete, Yoram Burak, and Ted Brookings. What grid cells convey about rat location. *Journal of Neuroscience*, 28(27):6858–6871, 2008.

Mark C Fuhs and David S Touretzky. A spin glass model of path integration in rat medial entorhinal cortex. *Journal of Neuroscience*, 26(16):4266–4276, 2006.

CC Alan Fung, KY Michael Wong, and Si Wu. A moving bump in a continuous manifold: a comprehensive study of the tracking dynamics of continuous attractor neural networks. *Neural Computation*, 22(3):752–792, 2010.

Marianne Fyhn, Sturla Molden, Menno P Witter, Edvard I Moser, and May-Britt Moser. Spatial representation in the entorhinal cortex. *Science*, 305(5688):1258–1264, 2004.

Marianne Fyhn, Torkel Hafting, Menno P Witter, Edvard I Moser, and May-Britt Moser. Grid cells in mice. *Hippocampus*, 18(12):1230–1238, 2008.

Ruiqi Gao, Jianwen Xie, Song-Chun Zhu, and Ying Nian Wu. Learning grid cells as vector representation of self-position coupled with matrix representation of self-motion. In *International Conference on Learning Representations*, 2019.

Ruiqi Gao, Jianwen Xie, Xue-Xin Wei, Song-Chun Zhu, and Ying Nian Wu. On path integration of grid cells: group representation and isotropic scaling. In *Neural Information Processing Systems*, 2021.

Richard J Gardner, Erik Hermansen, Marius Pachitariu, Yoram Burak, Nils A Baas, Benjamin A Dunn, May-Britt Moser, and Edvard I Moser. Toroidal topology of population activity in grid cells. *Nature*, 602(7895):123–128, 2022.

Mariana Gil, Mihai Ancau, Magdalene I Schlesiger, Angela Neitz, Kevin Allen, Rodrigo J De Marco, and Hannah Monyer. Impaired path integration in mice with disrupted grid cell firing. *Nature neuroscience*, 21(1):81–91, 2018.

Torkel Hafting, Marianne Fyhn, Sturla Molden, May-Britt Moser, and Edvard I Moser. Microstructure of a spatial map in the entorhinal cortex. *Nature*, 436(7052):801, 2005.

Sepp Hochreiter and Jürgen Schmidhuber. Long short-term memory. *Neural computation*, 9(8): 1735–1780, 1997.

Aidan J Horner, James A Bisby, Ewa Zotow, Daniel Bush, and Neil Burgess. Grid-like processing of imagined navigation. *Current Biology*, 26(6):842–847, 2016.

Joshua Jacobs, Christoph T Weidemann, Jonathan F Miller, Alec Solway, John F Burke, Xue-Xin Wei, Nanthia Suthana, Michael R Sperling, Ashwini D Sharan, Itzhak Fried, et al. Direct recordings of grid-like neuronal activity in human spatial navigation. *Nature neuroscience*, 16(9):1188, 2013.

Nathaniel J Killian, Michael J Jutras, and Elizabeth A Buffalo. A map of visual space in the primate entorhinal cortex. *Nature*, 491(7426):761, 2012.

Diederik P Kingma and Jimmy Ba. Adam: A method for stochastic optimization. *arXiv preprint arXiv:1412.6980*, 2014.

Rosamund F Langston, James A Ainge, Jonathan J Couey, Cathrin B Canto, Tale L Bjerknes, Menno P Witter, Edvard I Moser, and May-Britt Moser. Development of the spatial representation system in the rat. *Science*, 328(5985):1576–1580, 2010.

Bruce L McNaughton, Francesco P Battaglia, Ole Jensen, Edvard I Moser, and May-Britt Moser. Path integration and the neural basis of the'cognitive map'. *Nature Reviews Neuroscience*, 7(8): 663, 2006.

John O'Keefe. A review of the hippocampal place cells. *Progress in neurobiology*, 13(4):419–439, 1979.

Hugh Pastoll, Lukas Solanka, Mark CW van Rossum, and Matthew F Nolan. Feedback inhibition enables theta-nested gamma oscillations and grid firing fields. *Neuron*, 77(1):141–154, 2013.

Thomas Ridler, Jonathan Witton, Keith G Phillips, Andrew D Randall, and Jonathan T Brown. Impaired speed encoding is associated with reduced grid cell periodicity in a mouse model of tauopathy. *bioRxiv*, page 595652, 2019.

Francesca Sargolini, Marianne Fyhn, Torkel Hafting, Bruce L McNaughton, Menno P Witter, May-Britt Moser, and Edvard I Moser. Conjunctive representation of position, direction, and velocity in entorhinal cortex. *Science*, 312(5774):758–762, 2006.

Ben Sorscher, Gabriel Mel, Surya Ganguli, and Samuel A Ocko. A unified theory for the origin of grid cells through the lens of pattern formation. 2019.

Sameet Sreenivasan and Ila Fiete. Grid cells generate an analog error-correcting code for singularly precise neural computation. *Nature neuroscience*, 14(10):1330, 2011.

Hanne Stensola, Tor Stensola, Trygve Solstad, Kristian Frøland, May-Britt Moser, and Edvard I Moser. The entorhinal grid map is discretized. *Nature*, 492(7427):72, 2012.

Michael Taylor. Lectures on lie groups. *Lecture Notes, available at http://www. unc. edu/math/Faculty/met/lieg. html*, 2002.

Michael M Yartsev, Menno P Witter, and Nachum Ulanovsky. Grid cells without theta oscillations in the entorhinal cortex of bats. *Nature*, 479(7371):103, 2011.

Anthony Zee. *Group theory in a nutshell for physicists*. Princeton University Press, 2016.

Appendix A. More experimental details

A.1. Training details

We train the model for $200,000$ iterations and learn the model by minimizing $L_1 + \lambda L_2$, where $\lambda = 0.05$. For the extra zero-step version of L_1, the weight is set as 10. The regularization of $\|\boldsymbol{q}(\boldsymbol{p})\|^2$ is added to the loss in the first $10,000$ iterations, where we linearly decay the weight of the regularization from 0.1 to 0. During training, the normalization of $\|\boldsymbol{v}(\boldsymbol{x})\|$ is done for every position $\boldsymbol{x}$ at the end of each epoch. For each module, we normalize the value of $\|\boldsymbol{v}(\boldsymbol{x})\|$ to be $1/\sqrt{\boldsymbol{m}}$, where $\boldsymbol{m}$ is the number of modules. For isometry loss (L_2), we fix the upper bounds of displacements as 15 grids for all modules for the first $10,000$ iterations, and start to adaptively adjust the upper bounds afterwards every 2000 iterations. All the learned parameters are updated by Adam (Kingma and Ba, 2014) optimizer. The learning rate is linearly decayed from 0.006 to 0.0003 for the first $10,000$ iterations, fixed as 0.0003 until $120,000$ iterations, and then linearly decayed to 0 afterwards. For batch sizes, we use 8000 for zero-step transformation loss and isometry loss (L_1), and 100 for multi-step transformation loss. We trained all the models on a single 2080 Ti GPU.

A.2. Learned patterns

In Figure 5, we show the autocorrelograms of the learned grid patterns from the vanilla recurrent network.

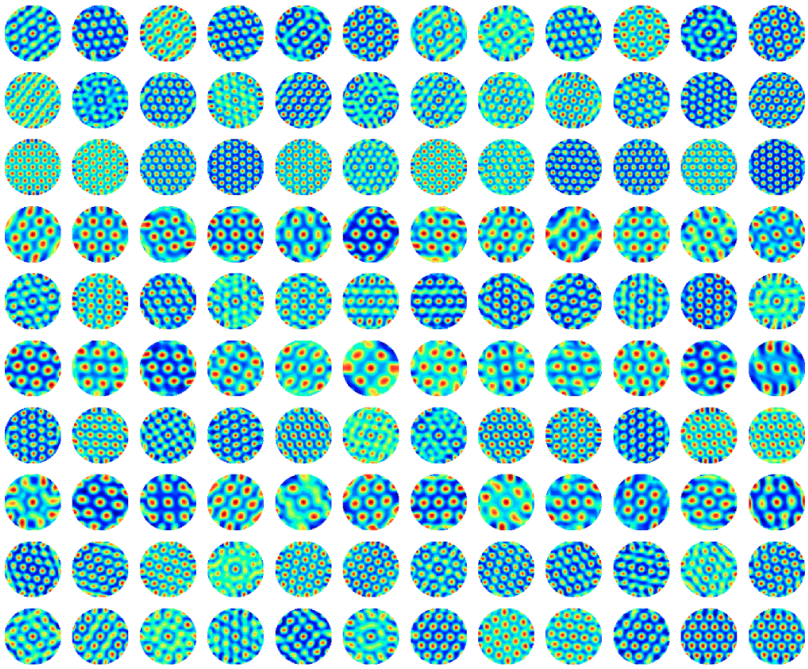


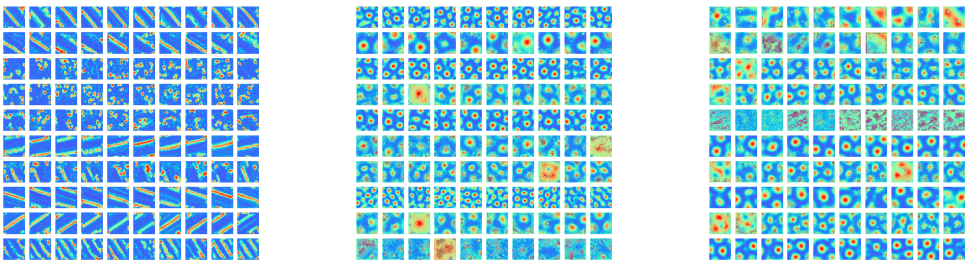
Figure 5: Autocorrelograms of the learned patterns.

A.3. Ablation studies

In this section, we present part of our ablation results to examine whether certain components of our model are empirically important for the emergence of hexagon grid patterns. Here are some key observations: (1) hexagon patterns do not emerge without conformal isometry loss. That is, the conformal isometry condition is crucial for the emergence of grid-like patterns. (2) Regularization of $\|\boldsymbol{q}(\boldsymbol{p})\|^2$ is not necessary, but the patterns are less clear without it. (3) Without zero-step transformation loss, the learned model is unable to path integrate accurately, although hexagon grid patterns still emerge.

We further try different module size. Figure 8 visualizes the learned patterns when we fix the total number of grid cells and change the module size to 24. It shows hexagonal grid firing patterns can emerge with larger module size. In Figure 9, we show the path integration error based on different lengths (5-step, 10-step, 15-step and 30-step) of RNN models. Path integration for 30-step trajectories is performed for settings with and without re-encoding.

Finally, we try to remove the block-diagonal assumption in the transformation model, i.e. we change $\boldsymbol{W}$ to be a full matrix instead of a block-diagonal one. But we still impose conformal isometry on the modules. In this setting, we can still learn multi-scale patterns as in Figure 7, while the path integration also works well. For a learned 10-step vanilla RNN without block-diagonal construction, the average error of 500 step path integration is around 0.02 with re-encoding every 10 steps and 0.046 without re-encoding over the whole trajectory. This suggests that the emergence of the hexagon patterns is not from block-diagonal construction but the isometry condition.



(a) Without isometry loss (b) Without regularization of $\boldsymbol{u}$ (c) Without zero-step loss

Figure 6: Results of ablation on certain components of the training loss. (a) Learned patterns without conformal isometry loss. (b) Learned patterns without the regularization of $\|\boldsymbol{q}(\boldsymbol{p})\|^2$. (c) Learned patterns without zero-step transformation loss.

Appendix B. Eigen analysis

Proof With $\Delta\boldsymbol{x} = 0$, we have the following fixed point property:

$$\boldsymbol{v}(\boldsymbol{x}) = \mathrm{ReLU}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x})), \tag{11}$$

$$\boldsymbol{v}(\boldsymbol{x}+\delta\boldsymbol{x}) = \mathrm{ReLU}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}+\delta\boldsymbol{x})). \tag{12}$$

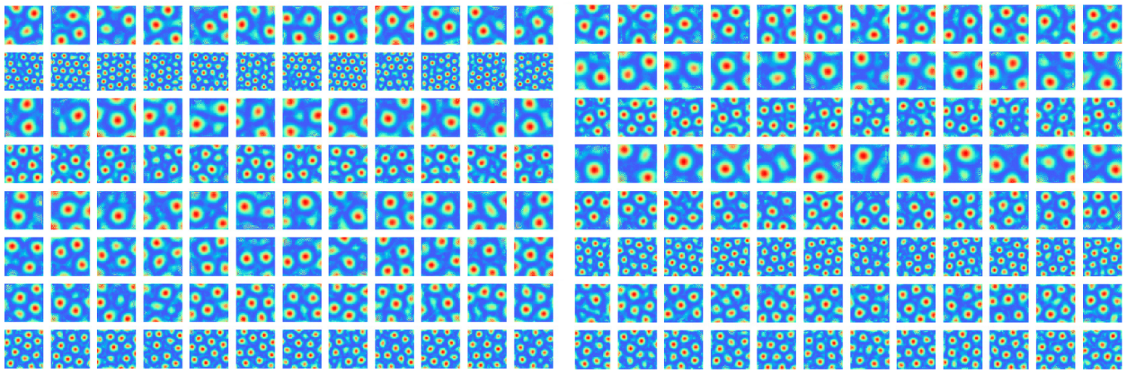


Figure 7: Hexagonal patterns emerged from the transformation model without block-diagonal assumption.

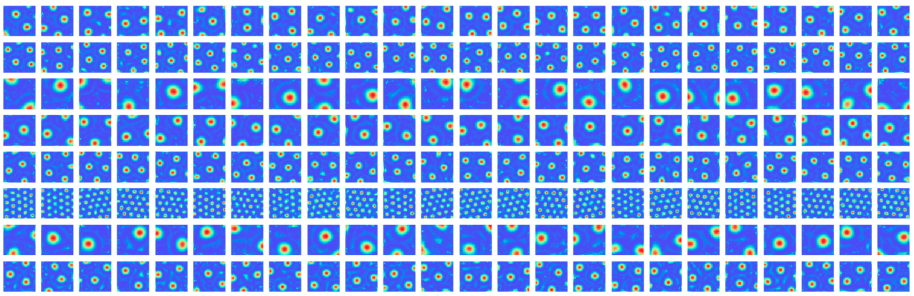


Figure 8: Learned patterns with module size 24.

Thus

$$\boldsymbol{v}(\boldsymbol{x}+\delta\boldsymbol{x})-\boldsymbol{v}(\boldsymbol{x}) = \text{ReLU}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}+\delta\boldsymbol{x}))-\text{ReLU}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x})) \tag{13}$$

$$= \mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x})>\mathbf{0})\odot\boldsymbol{W}(\boldsymbol{v}(\boldsymbol{x}+\delta\boldsymbol{x})-\boldsymbol{v}(\boldsymbol{x}))+o(|\delta\boldsymbol{x}|). \tag{14}$$

Thus for any $\boldsymbol{x}$, the matrix $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x})>\mathbf{0})\odot\boldsymbol{W}$ has an eigenvalue 1 with geometric multiplicity 2. Meanwhile,

$$\boldsymbol{v}(\boldsymbol{x}+\delta\boldsymbol{x})-\boldsymbol{v}(\boldsymbol{x}) = \text{ReLU}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x})+\boldsymbol{U}\delta\boldsymbol{x})-\text{ReLU}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x})) \tag{15}$$

$$= \mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x})>\mathbf{0})\odot\boldsymbol{U}\delta\boldsymbol{x}+o(|\delta\boldsymbol{x}|). \tag{16}$$

Under conformal isometry, the two column vectors of $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x})>\mathbf{0})\odot\boldsymbol{U}$ are orthogonal with equal norm, and they span the eigen-subspace of $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x})>\mathbf{0})\odot\boldsymbol{W}$.

Appendix C. Constructive conformal isometry

In our experiment, we impose conformal isomtry with a loss term based on Equation (4). In this section, we discuss a special case of the recurrent network that satisfies Condition 2 by design.

Specifically, we divide $\boldsymbol{v}$ into low-dimensional sub-vectors, $\boldsymbol{v}=(\boldsymbol{v}_k, k=1,...,K)$, where each $\boldsymbol{v}_k$ is a sub-vector of dimension d, e.g., $d=4$ so that each sub-vector consists of 4 neurons. We call each sub-vector a mini-block. Then the total dimension $D=Kd$. Such mini-blocks are commonly

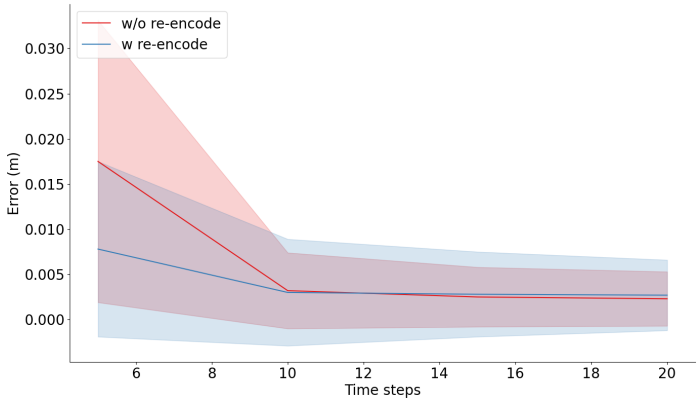


Figure 9: Path integration error over different lengths of RNN.

assumed in handcrafted continuous attractor neural networks (Burak and Fiete, 2009; Couey et al., 2013; Pastoll et al., 2013; Agmon and Burak, 2020). Correspondingly, we divide the $D \times 2$ matrix $\boldsymbol{U}$ into K mini-blocks, with each being $d \times 2$, so that $\boldsymbol{U} = (\boldsymbol{U}_k, k = 1, ..., K)$. The indicator vector $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) > \mathbf{0})$ is also divided into K mini-blocks, each of dimension d.

We assume the following two conditions for the mini-blocks:

Condition 3. *(Orthogonality condition). Each $\boldsymbol{U}_k$ has two column vectors that are of equal norm s_k and are orthogonal to each other, i.e., $\boldsymbol{U}_k^\top \boldsymbol{U}_k = s_k^2 \boldsymbol{I}_2$.*

The above condition can be easily satisfied, e.g., for $d = 4$, we let the first column of $\boldsymbol{U}_k$ be $[1, 0, -1, 0]^\top$ and the second column be $[0, 1, 0, -1]^\top$. The idea is that within each mini-block, the cells are sensitive to different directions of self-motion.

Condition 4. *(Synchronicity condition). Each mini-block of $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) > \mathbf{0})$ is synchronized, i.e., all its elements are either all 0 or all 1. For mini-block k, define $a_k = 1$ if all the elements of $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) > \mathbf{0})$ are 1, and $a_k = 0$ if all the elements of $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) > \mathbf{0})$ are 0.*

We call the mini-blocks that satisfy the above two conditions the conformal mini-blocks. Given those two conditions, we have the following result.

Theorem 2. *Under the orthogonality condition and the synchronicity condition, the mini-block-wise recurrent network satisfies the conformal isometry condition (Condition 2).*

Proof *The change of the vector*

$$\boldsymbol{v}(\boldsymbol{x} + \delta\boldsymbol{x}) - \boldsymbol{v}(\boldsymbol{x}) = \mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) > \mathbf{0}) \odot \boldsymbol{U}\delta\boldsymbol{x} + o(|\delta\boldsymbol{x}|). \tag{17}$$

Thus

$$\|\boldsymbol{v}(\boldsymbol{x} + \delta\boldsymbol{x}) - \boldsymbol{v}(\boldsymbol{x})\|^2 = \sum_k a_k s_k^2 \|\delta\boldsymbol{x}\|^2 + o(|\delta\boldsymbol{x}|^2) = s^2 \|\delta\boldsymbol{x}\|^2 + o(\|\delta\boldsymbol{x}\|^2), \tag{18}$$

where $s^2 = \sum_k a_k s_k^2$.

Define the activation pattern of $\mathbf{1}(\boldsymbol{W}\boldsymbol{v}(\boldsymbol{x}) > \mathbf{0})$ be $\boldsymbol{a}(\boldsymbol{x}) = (a_k, k = 1, ..., K)$. $\boldsymbol{a}(\boldsymbol{x})$ controls the change of the vector $\boldsymbol{v}(\boldsymbol{x} + \boldsymbol{\delta x}) - \boldsymbol{v}(\boldsymbol{x})$. At different $\boldsymbol{x}$, $\boldsymbol{a}(\boldsymbol{x})$ are different. Thus the direction of the change $\boldsymbol{v}(\boldsymbol{x} + \boldsymbol{\delta x}) - \boldsymbol{v}(\boldsymbol{x})$ depends on $\boldsymbol{x}$, and it is possible for the model to create rotation of the vector $\boldsymbol{v}(\boldsymbol{x})$.

It is an interesting problem to construct simple recurrent networks that satisfy conformal isometry automatically.