Path Integrator Contributions to

**Hippocampal Map Formation** 

David S. Touretzky

Computer Science Department

Carnegie Mellon University

Abstract

Skaggs and McNaughton's (1998) experiment in which rats shuttled between two

visually identical boxes provided a clear demonstration of stable partial remapping

in hippocampal place cells. Any satisfactory theory of hippocampal map formation

must account for both the degree of remapping and the stability of the partially

remapped state. We suggest that a weak path integrator is reponsible for the remap-

ping observed in this experiment, but the PI also helps to stabilize the remapped

state. These ideas are tested in a computer simulation.

Key words: hippocampus, cognitive map, place cells, path integrator

Email address: dst@cs.cmu.edu (David S. Touretzky).

## 1 Introduction

Skaggs and McNaughton [1] reported that rats shuttling between two visually identical boxes connected by a corridor showed similar place codes in each box; see Figure 1. The degree of "partial remapping" between boxes varied somewhat from day to day, but with no discernable trend. This poses a problem for place cell theories in which the hippocampal map is a crossproduct of extrinsic (local view) and intrinsic (path integrator, or PI) variables, and failure to properly reset the PI upen reentry into an environment is what leads to remapping [2,3]. If the PI is not being reset upon each box entry, why isn't the remapping complete? On the other hand, if the PI is reset to an identical value upon each box entry, why is there any remapping at all? One possibility is that the rat remembers whether it entered the current box via a left turn (north box) or a right turn (south box), but it would have to retain that information throughout its visit. Here we consider a different explanation.

We suggest that the PI is not reset upon box entry, but exerts a relatively weak influence on map selection when visual cues are identical in two environments. The primary contribution of the PI to map formation is to help constrain the nascent place code to a proper 2D manifold. But the discrepancy in PI coordinates in the two boxes does induce some differences, hence partial remapping. Hebbian learning would then cause cells with fields in both boxes to become bound to two sets of path integrator coordinates. The PI thus both causes

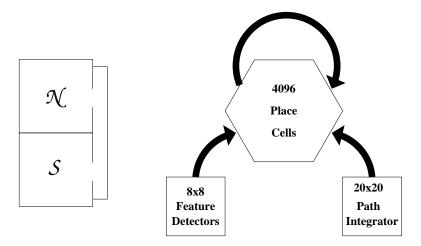


Fig. 1. Left: the two-box apparatus of Skaggs and McNaughton (1998): two visually identical boxes connected by a corridor. Right: structure of our place cell model. partial remapping and preserves it. We explore this hypothesis with computer simulations.

#### 2 Structure of the Model

We use a simple model in which place cells are driven by a combination of local view (feature detector) inputs, path integrator inputs, and recurrent connections. Pseuo-random wiring is used, so the model does not assume any built-in structure beyond sparse connectivity. In other words, there are no pre-wired maps.

A box is represented as a  $5 \times 5$  grid, giving 25 unique locations, and contains eight unique landmarks: four box corners and four wall midpoints. Since the two boxes are visually identical, the same landmarks are used in both. Each landmark has a set of eight Gaussian feature detectors tuned to specific

distances from that landmark. Thus, there are a total of  $8 \times 8 = 64$  feature detectors.

The path integrator is represented as a  $20 \times 20$  array of units with toroidal topology to eliminate edge effects. The simulated rat's current position is indicated by the peak of a Guassian activity bump. The two adjacent  $5 \times 5$  boxes are laid down on this grid, so each of the 50 box locations has unique path integrator coordinates.

The model uses 4096 place cells, each of which randomly samples 16 feature detectors (2 of 8 for each of 8 landmarks), 40 path integrator units (out of 400 total), and 400 place cells (10% recurrent connectivity.) Instead of an explicit recurrent inhibition term, we calculate a dynamic threshold to maintan a 3% activity rate over the place cell population, giving 122 active place cells. Figure 2 shows the contribution of each input type (feature detectors, PI, and and recurrent connections) for each of the 122 active place cells when the simulated rat is in the center of the north box.

In order to simulate the effects of map learning as the rat forages randomly in the box, we use the following procedure. Each location in the box is visited once, in random order. The location determines both the visual feature detector activity pattern and the path integrator activity pattern. These inputs drive the place cells, which are allowed to settle to a stable state. Then, we (1) increase the strengths of all synapses from active PI cells onto active

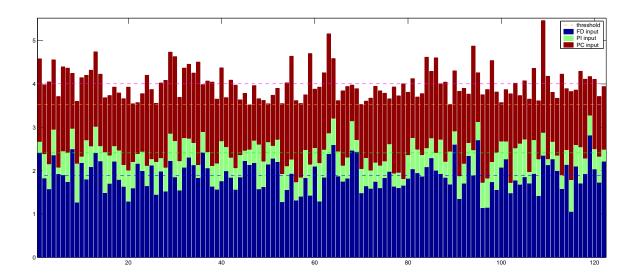


Fig. 2. Contributions of feature detector (FD), path integrator (PI), and recurrent place cell inputs (PC) to each of the 122 place cells active when the simulated rat is in the center of the north box. [Figure is in color but will be converted to b/w for final draft.] place cells, (2) slightly decrease the strengths of all recurrent synapses, but (3)

increase the strengths of recurrent synapses between active place cells. Then we pick another box location and continue the learning process. After all locations in the current box have been visited, we switch boxes and repeat the procedure, then return to the first box, etc. Each box is visited in this way several times, simulating the multiple days of exposure provided by Skaggs and McNaughton in their experiment.

## 3 Results

Treating the  $5 \times 5$  array of box locations as a 25 element vector, each plot in Figure 3 shows activity pattern cross-correlations for 50 locations, 25 per box.

These are organized in an L shape consisting of box 1-1, box 1-2, and box 2-2 correlations. The bottom left plot shows that prior to learning, there is a weak correlation in activity patterns for adjacent locations, and due to the PI input, an even weaker correlation between the two boxes. After learning (bottom right), the correlation between adjacent locations in a box is higher (i.e., the place code is more robust), and the correlation between boxes is also increased (less remapping). But remapping has not been eliminated.

Examining the PI afferents of active place cells shows that many cells are bound to two sets of PI coordinates. In other words, while the learning procedure made the maps more similar by potentiating the recurrent connections, it also helped to cement the partially remapped state by strengthening both sets of PI connections.

## 4 Discussion

We assume that the learning described in this model takes place during the initial formation of place fields as the rat explores the boxes for the first time. Whether learning acts to increase or decrease remapping depends on the relative rates of potentiation of place cells' recurrent connections vs. their path integrator connections. The model predicts that LTP-deficient rats should show greater amounts of remapping in the two-box experiment than normal rats, because the lack of potentiation of recurrent synapses that would help

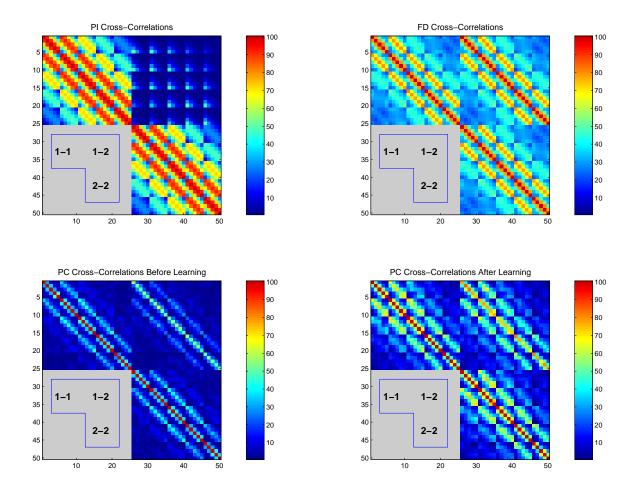


Fig. 3. Top left: cross-correlations of PI activity patterns for 50 locations, 25 per box.

Top right: cross-correlation of feature detector activity patterns; since the boxes are visually identical, the across-box correlations are the same as the within-box correlations. Bottom: place cell activity pattern cross-correlations, (left) prior to any learning, and (right) after learning. [Figure is in color but will be converted to b/w for final draft.]

to pull the maps together.

Skaggs and McNaughton found that the extent of remapping in their experiment showed no increasing or decreasing trend [1]. Other experiments have shown remapping that increases with increased exposure to the two envi-

ronments, either abruptly after a delay [4], or gradually [5]. An increase in remapping with experience would appear to involve mechanisms other than Hebbian learning [6].

In Knierim's double cue rotation experiment [7], place code dissociation, individual place field splitting, and partial remapping are attributed to weak attractor dynamics. But they also imply a weak but nonzero PI influence. When some cells appear to follow local landmarks while others follow distal, the weak PI may remain bound to the distal reference frame and compete with local landmarks for control of place fields.

# Acknowledgments

Supported by NIH MH59932. Thanks to Mark Fuhs for helpful discussions.

#### References

- [1] W. E. Skaggs, B. L. McNaughton, Spatial firing properties of hippcampal CA1 populations in an environment containing two visually identical regions, Journal of Neuroscience 18 (20) (1998) 8455–8466.
- [2] A. D. Redish, D. S. Touretzky, Cognitive maps beyond the hippocampus, Hippocampus 7 (1) (1997) 15–35.
- [3] A. D. Redish, D. S. Touretzky, Separating hippocampal maps, in: N. Burgess,

- K. Jeffery, J. O'Keefe (Eds.), The Hippocampal and Parietal Foundations of Spatial Cognition, Oxford University Press, New York, 1999, pp. 203–219.
- [4] E. Bostock, R. U. Muller, J. L. Kubie, Experience-dependent modifications of hippocampal place cell firing, Hippocampus 1 (2) (1991) 193–206.
- [5] H. Tanila, M. L. Shapiro, H. Eichenbaum, Discordance of spatial representation in ensmbles of hippocampal place cells, Hippocampus 7 (6) (1997) 613–623.
- [6] M. C. Fuhs, D. S. Touretzky, Synaptic learning models of map separation in hippocampus, Neurocomputing 32 (2000) 379–384.
- [7] J. J. Knierim, Dynamic interactions between local surface cues, distal landmarks, and intrinsic circuitry in hippocampal place cells, Journal of Neuroscience 22 (14) (2002) 6254–6264.