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A novel somatosensory spatial navigation system outside the hippocampal formation

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Spatially selective firing of place cells, grid cells, boundary vector/border cells and head direction cells constitutes the basic building blocks of a canonical spatial navigation system centered on the hippocampal-entorhinal complex. While head direction cells can be found throughout the brain, spatial tuning outside the hippocampal formation is often non-specific or conjunctive to other representations such as a reward. Although the precise mechanism of spatially selective firing activity is not understood, various studies show sensory inputs, particularly vision, heavily modulate spatial representation in the hippocampal-entorhinal circuit. To better understand the contribution of other sensory inputs in shaping spatial representation in the brain, we performed recording from the primary somatosensory cortex in foraging rats. To our surprise, we were able to detect the full complement of spatially selective firing patterns similar to that reported in the hippocampal-entorhinal network, namely, place cells, head direction cells, boundary vector/border cells, grid cells and conjunctive cells, in the somatosensory cortex. These newly identified somatosensory spatial cells form a spatial map outside the hippocampal formation and support the hypothesis that location information modulates body representation in the somatosensory cortex. Our findings provide transformative insights into our understanding of how spatial information is processed and integrated in the brain, as well as functional operations of the somatosensory cortex in the context of rehabilitation with brain-machine interfaces.

INTRODUCTION

Spatial representation in the brain has been largely associated with the hippocampus and the entorhinal cortex.[1,2] Place cells[3] were originally discovered in the hippocampus, with boundary vector/border cells,[4,5] grid cells[6] and conjunctive cells[7] discovered later in the entorhinal cortex. Unlike the location-specific spatial cells in the hippocampal-entorhinal circuit, head direction cells[8] can be found in areas ranging from the diencephalon to thalamic, striatal and cortical regions.[9]

Despite the focus on hippocampal-entorhinal contributions to spatial representation, location-dependent firing conjunctive to other sensory/cognitive (e.g., reward) information has been reported in sensory areas[10–12] and cortical regions heavily interconnected with the hippocampal-entorhinal system such as the septum, subicular complex, cingulate cortex, parietal cortex and retrosplenial cortex.[13–20] In rodents, the only extra-parahippocampal region exhibiting apparent location-dependent firing restricted to space alone is the anterior claustrum, where stable, visually-anchored place cell- and border cell-like activities can be recorded.[21] A preliminary report also suggests the dorsal geniculate nucleus may contain place cells similar to those found in the hippocampus.[22] In humans, hexadirectional grid-like signals have been observed in many brain regions outside the hippocampal formation including the posterior cingulate cortex, the medial prefrontal cortex, the retrosplenial cortex, the medial parietal cortex and the frontal cortex[23–26] while grid-cell-like neuronal representations were identified with intracranial electro-encephalography recordings on presurgical epilepsy patients and

fMRI studies in the human entorhinal cortex.[27–29] Computational modeling also predicted the existence of grid cell-like neurons in all cortical regions throughout the neocortex.[30] These results therefore indicate that spatial tuning is much more distributed into multiple cortical domains beyond the classical hippocampal formation than we previously thought.

The exact physiological mechanism of spatial selectivity in the hippocampal-entorhinal system is still incompletely understood. Lesions to either area severely affect, but do not abolish spatially selective activities in the other.[31,32] Past studies have shown manipulation of visual cues,[33,34] olfaction[35] and vibrissae inputs[36] all impact hippocampal place cell activity, yet none is necessary. Recent theoretical work has suggested the somatosensory area may contain higher-order presentations of the body itself beyond elementary sensory information, of which location presentation is integral in a "body simulation" model of somatosensation.[37] While there are place cell-like activities in the wheel chair-seated monkey sensorimotor cortices,[38] it is unclear if these spatial representations are comparable to their freely moving rodent equivalents. Based on the assumption that somatosensation should play a major role in sensory inputs shaping parahippocampal spatial activity[39,40] and spatial presentation has been theorized to be crucial for body representation,[37] we sought to identify spatially selective activities in the primary somatosensory cortex (S1). Remarkably, we were able to detect all spatial cell types reported in the hippocampal-entorhinal system[1,2] within the S1. We show these spatial cell types within this newly discovered somatosensory spatial representation system share similar

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properties to their parahippocampal counterparts in terms of stability and the response to environmental manipulations.[41,42]

RESULTS

Extracellular recording from the somatosensory cortex

We obtained neurophysiological recordings in freely behaving rats by using tetrodes mounted on movable micro-drives. We recorded 2025 putative single units from eight implanted rats across 287 recording sessions. Recordings were obtained mainly from layers IV to VI within the S1 area (Fig. 1a; Supplementary information, Fig. S1), as confirmed by the histological reconstruction. The tetrodes were generally lowered in steps of either 25 μm or 50 μm per day. The recording sessions lasted between 10 and 30 min to ensure sufficient coverage of the whole arena, which was facilitated by foraging food pellets intermittently thrown into the arena. Spike sorting was performed manually off-line using TINT (Axona, St. Albans, U.K.). The separation of the clusters and waveforms was quantitatively measured with an isolation distance and the L-ratio[43] (Supplementary information, Fig. S2).

We applied the same analysis described in previous work[7,17,44] on 2025 somatosensory cells and 390 met the criteria for various types (e.g., place, head direction, boundary vector/border and grid cells) of spatial tuning (Supplementary information, Fig. S3). The number of each functionally distinct somatosensory spatial cell type from individual implanted animals was summarized in the Supplementary information, Table S1. No significant differences were observed among spatial information, mean vector length and grid score across different animals (ANOVA test, $P = 0.38$, 0.44 and 0.18, respectively; Supplementary information, Fig. S4a, b, d). Both border score and grid spacing remained similar across individual animals except that they are slightly different in one of eight animals (ANOVA test, $*P < 0.05$; Supplementary information, Fig. S4c, d). The distribution of different spatial cell types across different animals suggested, at least between layers IV and VI, there was a tendency for head direction cells to avoid layer VI and the presence of spatially tuned somatosensory cells was sparse in superficial layer V (Supplementary information, Fig. S3b–f). Place cells appeared to predominate deeper layers (Supplementary information, Fig. S3a–c).
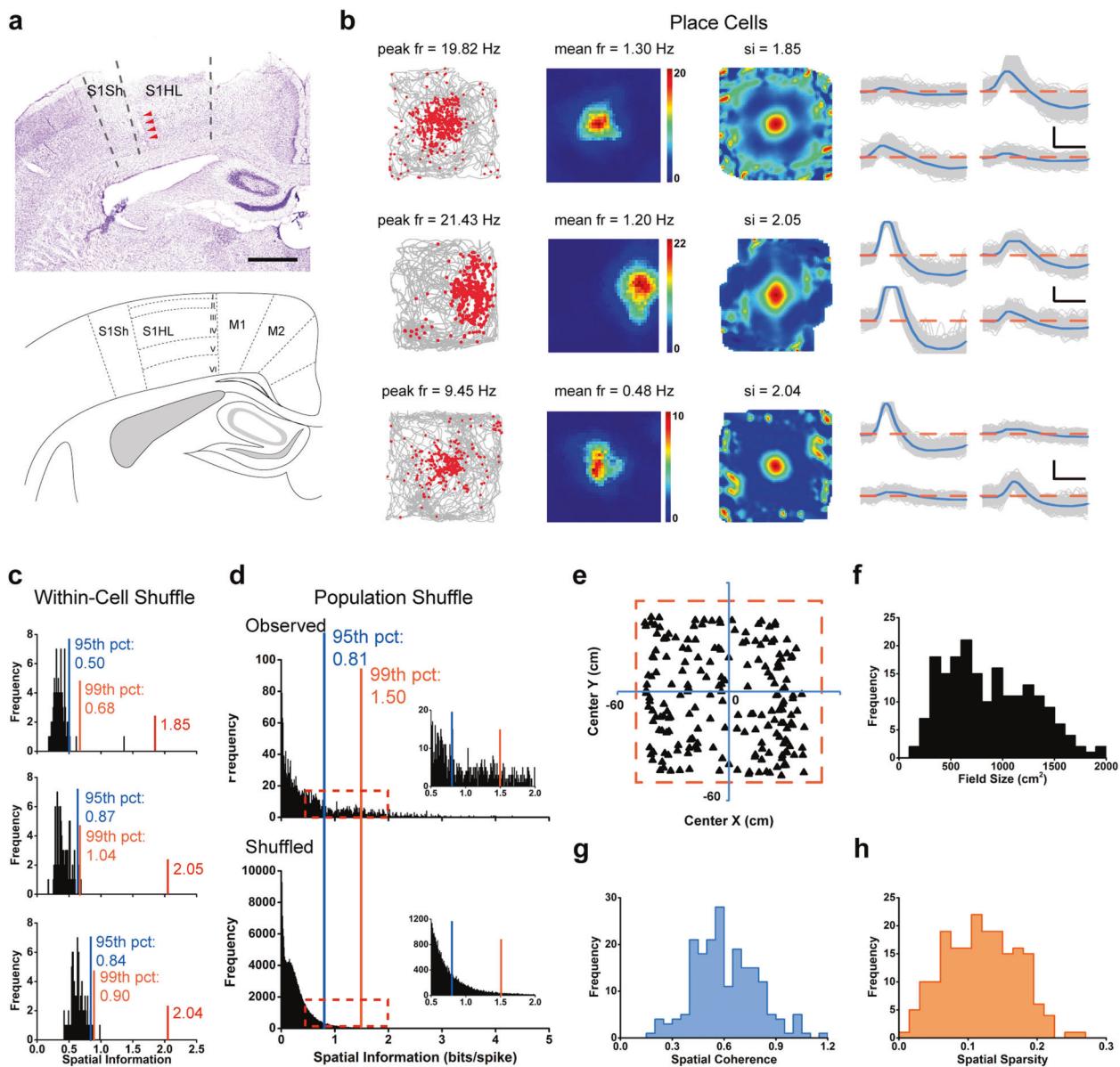
Place cells in the somatosensory cortex

As illustrated in Fig. 1, somatosensory cells possessing place cell-like properties were found the most abundant spatial cell type (Supplementary information, Fig. S3a–c). Spatially selective firing patterns similar to those originally described for hippocampal place cells[3] can be observed on them (Fig. 1b; Supplementary information, Fig. S5). These S1 place cells were classified if their spatial information (SI) scores[45] exceed a stringent 99th percentile threshold (SI > 1.50 cut-off) from the population-based distribution of shuffled rate maps (Fig. 1d). 195 out of the 2025 S1 neurons (9.63%) met criterion and thus are referred to as "place cells" from here on. These place cells also passed the spatial threshold for the within-cell shuffling (Fig. 1c; Supplementary information, Fig. S6a), which is less stringent than the population shuffling. The within-cell shuffling outcome is also true for other types of somatosensory spatial cells (Supplementary information, Fig. S6b–d). Accordingly, we used the population shuffling for defining all different types of somatosensory spatial cells in this study. Furthermore, we applied the maximum-likelihood approach[46,47] to evaluate inhomogeneous sampling biases on spatial tuning of S1 place cells. By employing the maximum-likelihood correction technique, we found that the locational information was increased by 7.11% on average (SD, 7.25%) (Supplementary information, Fig. S7d, e) among a total sample of 195 identified S1 place cells, which showed a robust locational effect and were not altered by the correction algorithm (Supplementary information, Fig. S7a–c). Such marginal increase in locational information after applying

maximum-likelihood approach was similar to that for hippocampal place cell reported by Burgess et al.[47] The average SI in bits per spike of all 195 classified S1 place cells was 2.19 ± 0.04 (all data are shown as means $\pm$ SEM unless otherwise indicated). This percentage was significantly higher than expected by random selection from the entire shuffled population (Fig. 1d; $Z = 39.03$, $P < 0.001$; binomial test with expected P_0 of 0.01 among large samples).

We noted that if a 95th percentile criterion was applied, 511 out of the 2025 S1 neurons (24.00%) would be classified as place cells (average SI of 1.59 ± 0.03; cut-off = 0.81). This percentage was substantially higher than expected with a random selection from the entire shuffled population (Fig. 1d; $Z = 109.61$, $P < 0.001$; binomial test with expected P_0 of 0.01 among large samples). We found that the percentage of identified S1 place cells is lower within the somatosensory cortex than that of hippocampal place cells.[48] The average peak firing rate and the mean firing rate of identified somatosensory location-specific place cells were 10.32 ± 0.37 Hz and 0.70 ± 0.04 Hz (Supplementary information, Fig. S8a), respectively. The histograms of peak-to-peak amplitude and peak-to-trough spike width were shown in Supplementary information, Fig. S8e, and both the average and the highest amplitudes of identified S1 place cells were higher than those of all the other spatial cell types identified from the somatosensory cortex (Supplementary information, Fig. S8i). Like a mixture of a single place field and multiple place fields identified from the hippocampal place cells,[49] we found that the proportion of multi-field place cells was 25.64%. The average somatosensory spatial firing field size was 891.82 ± 32.65 cm^2 (Fig. 1f). To quantify the smoothness of recorded somatosensory place fields, we calculated the spatial coherence by computing the mean correlation between the firing rate of each bin and the averaged firing rate of the 8 neighboring bins from unsmoothed spatial firing rate maps.[34] For 195 recorded somatosensory place cells, the average spatial coherence was 0.61 ± 0.01 (Fig. 1g). Furthermore, we also calculated spatial sparsity to evaluate the extent of spatial information in each spike discharged by S1 place cells.[45] The spatial sparsity measure of S1 place cells was 0.12 ± 0.01 (Fig. 1h). Besides, S1 place cells can be found at any position within the arena for the uniformity testing based on the Friedman-Rafsky's MST test (Fig. 1e, Friedman-Rafsky's MST test, $P = 0.42$, two-tailed z-test) as in the hippocampus.[50]

Place cell activity was defined not only by its location-specific firing but also by the stability of its spatial representation. To this end, we next evaluated the spatial stability of somatosensory place fields by computing the spatial correlations between single intra-trial behavioral sessions. The average spatial correlation for S1 place cells was 0.64 ± 0.01 (Supplementary information, Fig. S9d) between the two halves of a recording session (Supplementary information, Fig. S9a–c), which remain stable in extended recording sessions up to 30 min (Supplementary information, Fig. S10a–f). There was no significant change in both average firing rate and SI between 10 min and 30 min recording (Supplementary information, Fig. S10g, h; two-sided Wilcoxon signed rank test, $W = 13$, $n = 6$, $P = 0.60$ and $W = 8$, $n = 6$, $P = 0.60$, respectively), suggesting the firing field of S1 place cells were stable during the extended recordings. Hippocampal place cell activities were known to be heavily modulated by visual cues,[33,34] and were known to rotate with the positioning of visual cues.[34] We performed similar experiments to test if S1 place cell activities were modulated by visual cues by moving it 90° from its original position and back. Remarkably, the relative position of S1 place fields from four representative place cells did indeed rotate with the placement of the visual cue and remained stable after the cue was moved back to its original position (Supplementary information, Fig. S11a–d). Notably, four pairs of co-recorded border cells and place cells rotated concurrently with the external cue card (Supplementary information, Fig. S11a–d). Furthermore, as hippocampal place codes were reorganized to remap under different

Fig. 1 Place cells in the somatosensory cortex. a A Nissl-stained coronal section (top) showing tetrode tracking trajectory (arrowheads) through all of six layers across the rat primary somatosensory cortex. Dashed lines depict the boundaries of the hindlimb (S1HL) and shoulder (S1Sh) regions of the primary somatosensory cortex (S1). The bottom panel shows the schematic delimitation of the different layers and sub-regions of the primary somatosensory cortex. Scale bar, 1 mm. **b** Trajectory (gray line) with superimposed spike locations (red dots) (left column); heat maps of spatial firing rate (middle left column) and autocorrelation (middle right column) are color-coded with dark blue indicating minimal firing rate and dark red indicating maximal firing rate. The scale of the autocorrelation is twice the scale of the spatial firing rate maps. Peak firing rate (fr), mean firing rate (fr) and spatial information (si) for each representative cell are labeled on the top of the panels. Spike waveforms on four electrodes are shown on the right column. Scale bar, 150 µV, 300 µs. The zero microvolt horizontal baseline is drawn with the orange dashed lines for the spike waveforms on all four electrodes. **c** Distribution of within-cell shuffled spatial information for three representative somatosensory place cells. The orange and blue stippled lines mark the 99th and the 95th percentile significance level of each randomly shuffled distribution, respectively. The red line indicates the observed spatial information. **d** Distribution of spatial information for all isolated somatosensory units. The top panel shows the distribution for observed spatial information. The bottom panel shows the distribution for randomly shuffled data from the same sample. The orange and blue stippled lines mark the 99th and the 95th percentile significance level of each randomly shuffled distribution, respectively. A zoomed panel shows the magnification of the specified area marked by the red dashed rectangle. **e** The uniformly areal distribution of various place firing fields relative to the field center. **f–h** The population histograms of place firing field sizes (**f**), spatial coherence (**g**) and spatial sparsity (**h**) for all of 195 identified somatosensory place cells.

geometric environments,[51,52] we also found that S1 place fields remapped between a square box and a circular box in the same recording room with changed firing rate and firing location (Supplementary information, Fig. S12a–h). Spatial correlation between rate maps on the square trails versus the circle trial (S-C and C-S') were significantly decreased compared to that on two square trials (S-S') (Supplementary information, Fig. S12j; two-sided Wilcoxon signed rank test, $W = 36$, $n = 8$, $P = 0.012$). The average firing rate was also altered (Supplementary information, Fig. S12i; two-sided Wilcoxon signed rank test, $W = 36$, $n = 8$, $P = 0.012$), indicating significant remapping of S1 place cells occurred between different environments.

To measure the animal's running speed within and outside the firing fields of S1 place cells, we calculated the average "in-field"

and "out-field" running speed (>2.5 cm/s) for 195 identified place cells in S1. We quantified the average running speed across all the spatial bins within the firing field of the place cell ("in-field" running speed) as well as outside the firing field of the place cell ("out-field" running speed) (Supplementary information, Fig. S13a). We found that there was no significant difference between the average "in-field" and "out-field" running speed (14.58 ± 0.25 cm/s vs. 15.00 ± 0.18 cm/s, $n = 195$, $P = 0.10$, two-tailed paired t-test; Supplementary information, Fig. S13b). To evaluate the effect of the animal's running speed on spatial responses, we calculated the spatial information scores of S1 place cells for running speed above 2.5 cm/s and below 2.5 cm/s, respectively. We found that the place fields during inactive mobility are less prevalent with lower spatial information score than those for active exploring (Supplementary information, Fig. S14a–c). Similar results were observed for S1 border cells (Supplementary information, Fig. S27a–c) and S1 grid cells (Supplementary information, Fig. S30a–c). As for S1 head direction cells, it appeared that head directional responses were less influenced with immobility than other three S1 spatial cell types (Supplementary information, Fig. S21a–c).

Head direction cells in the somatosensory cortex

Head direction cells discharge only when the animal's head is at a particular angle respective to the environment.[8] To classify the preferred firing angle of each cell, the mean vector length was used to compute head direction tuning.[7] Cells were categorized as head direction cells if the mean vector length exceeded the 99th percentile of the total distribution of the shuffling procedure from the entire pool of putative single cells (Fig. 2c). 80 cells (3.95% of all recorded cells) met the criterion, showing significant modulation with the animal's head direction (Fig. 2a; Supplementary information, Fig. S15). To correct for the possible effects of direction and location for identified head direction cells, we applied the maximum-likelihood correction approach[46,47] and found that three representative somatosensory head directional responses remain unaltered from the corrected directional firing rate histograms (Supplementary information, Fig. S16a–c). Of those 80 identified S1 head direction cells, the directional information was decreased by 8.37% on average (SD, -1.8%) (Supplementary information, Fig. S6d, e). Consequently, 70 cells still passed the stringent 99th percentile threshold with the mean vector length being 0.4 and were thus further analyzed. The decrease in directional information after applying maximum-likelihood correction algorithm was consistent with that for pre-subicular head direction cells reported by Burgess et al.[47] The average mean vector length was found to be 0.58 ± 0.01 and the mean firing rate to be 0.92 ± 0.08 Hz (Supplementary information, Fig. S8b). The histograms of peak-to-peak amplitude and peak-to-trough spike width were displayed in Supplementary information, Fig. S8f. Besides, the within-cell shuffling for those three representative somatosensory head direction cells generated similar results to the population shuffling (Fig. 2b). This percentage is significantly higher than expected by random selection from the entire shuffled population (Fig. 2c; $Z = 11.11$, $P < 0.001$; binomial test with expected P_0 of 0.01 among large samples). Notably, 352 out of 2025 identified S1 neurons (17.38%) showed HD tuning at a lower (95th percentile cut-off) threshold and had the mean vector length being 0.26. As expected, changing the cut-off from the stringent 99th percentile to 95th percentile resulted in drastic increase in the number of S1 head direction cells.

To measure the sharpness of the head directional tuning curve, we quantified the average full width at half maximum (FWHM) of somatosensory head direction cells. The average FWHM of the angular distributions was 67.97 ± 3.36 degrees (means $\pm$ SEM) (Fig. 2d). Across all recorded S1 head direction cells, the distribution of their head direction responses were broadly clustered across 360° (Rayleigh test for uniformity, $P = 0.08$,
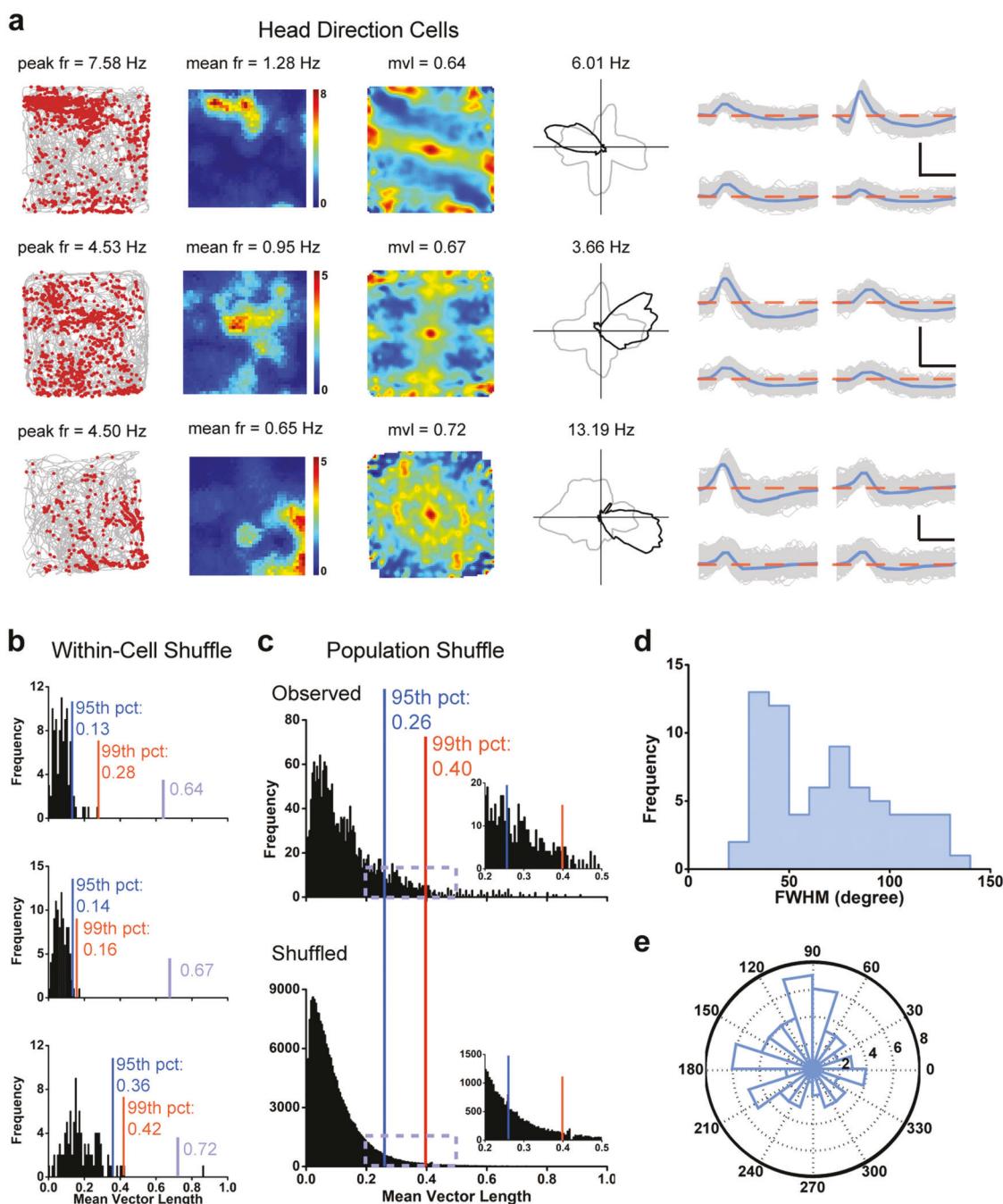
Fig. 2e). However, their preferred directions seemed to show a bias with a peak towards the north, where a white cue card pasted on the arena wall might contribute to such biased preferred head directions. Given a lower P value, it might be possible that with a larger recording dataset or under a different statistical test S1 head direction cells would have found uniformly clustered across 360°. We next evaluated the intra-trial directional stability by correlating the distributed firing rate across all directional bins between the first and second halves of individual recording trials. We found head direction-related firing to be stable within sessions (Supplementary information, Fig. S17a–c), with the average angular correlation coefficient for S1 head direction cells being 0.61 ± 0.02 (Supplementary information, Fig. S17d). Like somatosensory place cells, head direction cells also anchored to salient visual cues.[53] We further tested the specificity and stability of S1 head direction cells by rotating a salient visual cue 90° (R) from its original standard baseline (B) position in the enclosure and back to the second baseline (B') session (Supplementary information, Fig. S18, as in S1 place cells described above). As with co-recorded S1 place cells, S1 head direction cells also rotated with the movement of the salient visual cue (Supplementary information, Fig. S18a–d). We monitored 15 recorded head direction cells during a cue card rotation test. 9 out of 15 head direction cells rotated with the cue card taped to the inside wall of the running box. A 90° clockwise or counter-clockwise rotation for the cue card produced a near equal yet imperfect shift in the preferred direction (mean shift = $83.30 \pm 1.80°$ (B-R) and $85.28 \pm 3.49°$ (R-B'), two-sided Wilcoxon signed rank test, $W = 45$, $n = 9$, $P = 0.008$ and $W = 36$, $n = 9$, $P = 0.01$, respectively; Supplementary information, Fig. S18e), whereas 6 out of 15 tested S1 head direction cells did not follow the rotated cue card (mean shift = $12.39 \pm 3.79°$ (B-R) and $18.29 \pm 8.55°$ (R-B'), two-sided Wilcoxon signed rank test, $W = 9$, $n = 6$, $P = 0.75$ and $W = 10.5$, $n = 6$, $P = 0.41$, respectively; Supplementary information, Fig. S18f).

The animal's head turning velocity or angular head velocity, defined as the first derivative of head direction, was calculated as previously described.[54] To measure the difference of the animal's angular velocity within and outside preferred firing directions for those identified head direction cells in S1, we calculated the average angular velocity in the "preferred firing directions" and "non-preferred firing directions" for 70 identified head direction cells in S1 (Supplementary information, Fig. S19a). We found that there was no significant difference between the average angular velocity in the "preferred firing directions" and "non-preferred firing directions" for head direction cells in S1 (53.06 ± 1.18 degrees/s vs 54.17 ± 1.27 degrees/s, $n = 70$, $P = 0.47$, two-tailed paired t-test; Supplementary information, Fig. S19b).
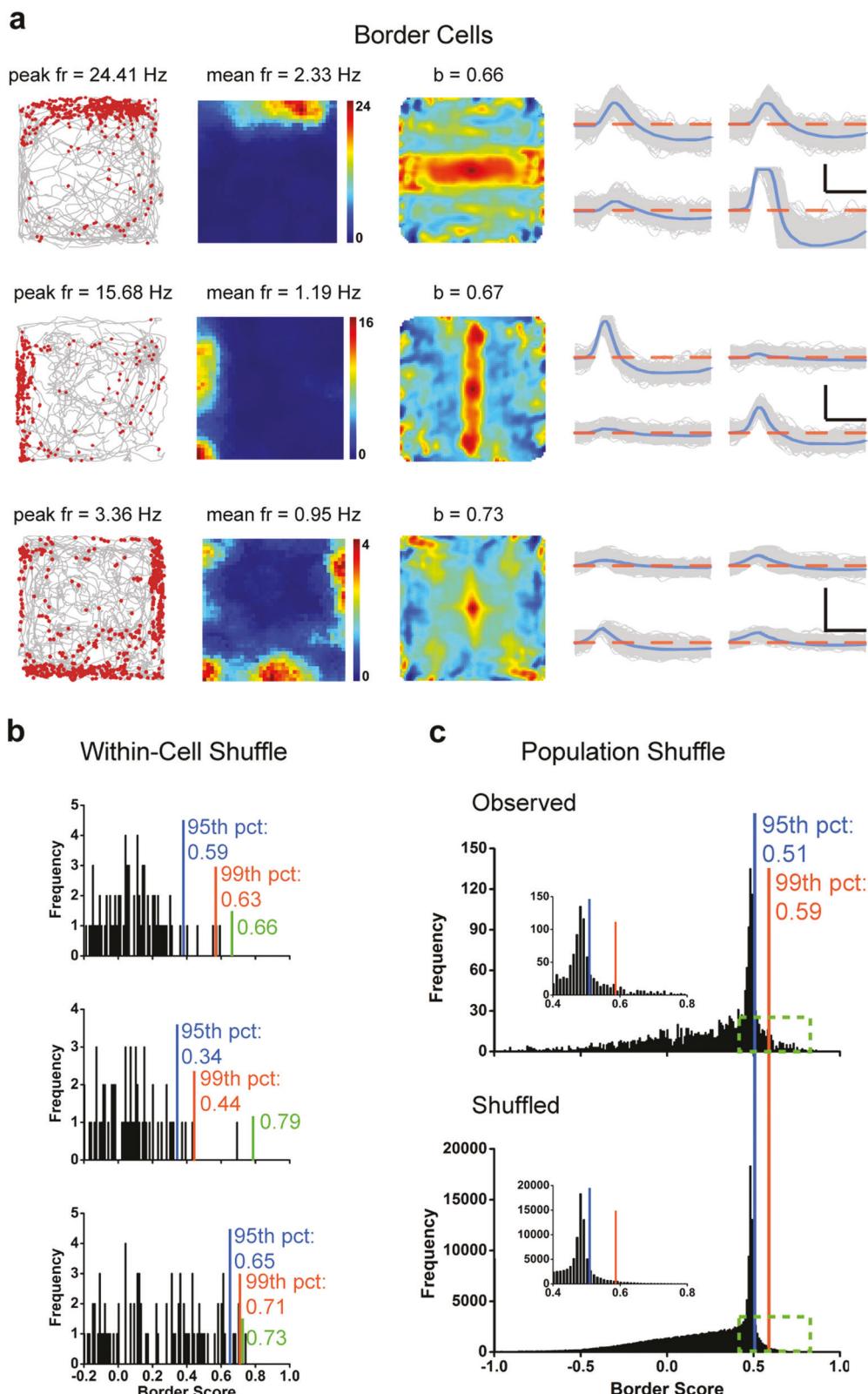
Although the previous study had shown that food chasing had little effect on head directional firing in the rat anterior thalamic nuclei,[54] we evaluated the firing patterns of head direction cells in the S1 without food pellet dispersion. We analyzed three pairs of co-recorded head direction cells and place cells from the same micro-drive in two consecutive secessions with (bottom panels in Supplementary information, Fig. S20a–c) and without (upper panels in Supplementary information, Fig. S20a–c) food pellets, respectively. No significant differences were observed, suggesting that food chasing did not modulate preferred firing angles in S1 head direction cells. Similar results were observed for place cells in the S1 without throwing food pellets within the running arena (Supplementary information, Fig. S20a–c).

Border/boundary vector coding in the somatosensory cortex

Boundary vector cells and border cells discharge exclusively when an animal is physically close, at a specific distance and direction, to one or several environmental boundaries, for example, the enclosure walls of the recording arena.[4,5,55] S1 cells were defined as border cells if border scores (see "Material and Methods") were larger than the 99th percentile of population shuffled scores

Fig. 2 Head direction cells recorded from the somatosensory cortex. a Three examples of somatosensory head direction cells. Trajectory (gray line) with superimposed spike locations (red dots) (left column); spatial firing rate maps (middle left column), autocorrelation diagrams (middle right column) and head direction tuning curves (black) plotted against dwell-time polar plot (gray) (right column). Firing rate is color-coded with dark blue indicating minimal firing rate and dark red indicating maximal firing rate. The scale of the autocorrelation is twice the scale of the spatial firing rate maps. Peak firing rate (fr), mean firing rate (fr), mean vector length (mvl) and angular peak rate for each representative head direction cell are labeled on the top of the panels. The directional plots show strong head direction tuning. Spike waveforms on four electrodes are shown on the right column. The zero microvolt horizontal baseline is drawn with the orange dashed lines for the spike waveforms on all four electrodes. Scale bar, 150 μV, 300 μs. **b** Distribution of within-cell shuffled mean vector length for three representative somatosensory head direction cells. The orange and blue stippled lines mark the 99th and the 95th percentile significance level of each randomly shuffled distribution, respectively. The purple line indicates the observed mean vector length. **c** Distribution of mean vector length for all identified somatosensory units. The top panel shows the distribution for observed values. The bottom panel shows the distribution for randomly shuffled data from the same sample. The orange and blue stippled lines mark the 99th and the 95th percentile significance level of each randomly shuffled distribution, respectively. The insets show the magnification of the specified area marked by the purple dashed rectangle. **d** Distribution of head directional tuning width. **e** Preferred direction of recorded head direction cells from S1. The polar plot shows the distribution of the peak firing direction of all identified somatosensory head direction cells. The preferred head direction exhibits a uniform distribution ($P = 0.08$, Rayleigh's test).

a

Border Cells

peak fr = 24.41 Hz   mean fr = 2.33 Hz   b = 0.66

peak fr = 15.68 Hz   mean fr = 1.19 Hz   b = 0.67

peak fr = 3.36 Hz   mean fr = 0.95 Hz   b = 0.73

b

Within-Cell Shuffle

95th pct: 0.59
99th pct: 0.63
0.66

95th pct: 0.34
99th pct: 0.44
0.79

95th pct: 0.65
99th pct: 0.71
0.73

c

Population Shuffle

Observed

95th pct: 0.51
99th pct: 0.59

Shuffled

(Fig. 3c). A total of 86/2025 (4.25%) S1 cells were classified as border cells, and this percentage was significantly higher than expected by random selection from the entire shuffled population (Fig. 3c; $Z = 14.68$, $P < 0.001$; binomial test with expected P_0 of 0.01 among large samples). Similar results were observed by performing cell-specific shuffling for those three representative somatosensory border cells (Fig. 3b). Most border cells ($n = 58$, 67.44%) fired along a single boundary while others were active along with two ($n = 15$, 17.44%), three ($n = 6$, 6.98%) or even four ($n = 7$, 8.14%) boundaries within the enclosure (Fig. 3a;

Fig. 3 Border cells recorded from the somatosensory cortex. a Three representative examples of somatosensory border cells. Trajectory (gray line) with superimposed spike locations (red dots) (left column); heat maps of spatial firing rate (middle column) and autocorrelation diagrams (right column). Firing rate is color-coded with dark blue indicating minimal firing rate and dark red indicating maximal firing rate. The scale of the autocorrelation is twice the scale of the spatial firing rate maps. Peak firing rate (fr), mean firing rate (fr) and border score (b) for each representative border cell are labeled on the top of the panels. Spike waveforms on four electrodes are shown on the right column. The zero microvolt horizontal baseline is drawn with the orange dashed lines for the spike waveforms on all four electrodes. Scale bar, 150 μV, 300 μs. **b** Distribution of within-cell shuffled border score for three representative somatosensory border cells. The orange and blue stippled lines mark the 99th and the 95th percentile significance level of each randomly shuffled distribution, respectively. The green line indicates the observed border score. **c** Distribution of border scores for pooled somatosensory cells. The top panel shows the distribution for observed values. The bottom panel shows the distribution for randomly shuffled versions from the same sample. The orange and blue stippled lines mark the 99th and the 95th percentile significance level of each randomly shuffled distribution, respectively. A zoomed panel shows the magnification of the specified area marked by the green dashed rectangle.

Supplementary information, Fig. S22). The average firing rate and the border score were 1.00 ± 0.09 Hz (Supplementary information, Fig. S8c) and 0.66 ± 0.01, respectively. The histograms of peak-to-peak amplitude and peak-to-trough spike width were summarized in Supplementary information, Fig. S8g. The number of S1 border cells seemed to be higher in layer V than in layer IV or layer VI (Supplementary information, Fig. S3b, e). At a lower 95th percentile criterion, 234/2025 (11.56%) somatosensory cells were classified as border cells.

Qualitatively, S1 border cells maintained their border-specific firing fields between circular and square enclosures (Supplementary information, Fig. S23a–g), comparable to boundary vector cells and border cells from the presubiculum, parasubiculum and the medial entorhinal cortex (MEC).[4,5,17,55] There was no significant change in both average firing rate and border scores for the same border cell recorded between running boxes of different geometric shapes (Supplementary information, Fig. S23h, i; two-sided Wilcoxon signed rank test, $n = 7$, $P > 0.05$). S1 border cells ($n = 6$) also responded to a new wall insert to a familiar enclosure (Supplementary information, Fig. S24a–f), as described in the MEC.[5,55] As observed in the MEC and the subiculum,[4,5] removal of walls from the enclosure on an elevated platform preserved similar "border" specificity as in the walled arena.[55] These results confirmed that somatosensory border cells continued to code for geometric boundaries instead of physical borders in the unwalled environments (Supplementary information, Fig. S25a–f). There was no significant change in average firing rate from the first standard square enclosure (S) with walls to the elevated platform without walls (E) and from the elevated platform without walls back to the second standard (S') walled square enclosure [Supplementary information, Fig. S25g; two-sided Wilcoxon signed rank test, $W = 21$, $n = 6$, $P = 0.11$ (S-E) and $W = 21$, $n = 6$, $P = 0.11$ (E-S'), respectively]. However, the border scores of the same recorded border cells on the elevated platform as those from the square enclosure showed a tendency of decrease [Supplementary information, Fig. S25h; two-sided Wilcoxon signed rank test, $W = 21$, $n = 6$, $P = 0.028$ (S-E) and $W = 21$, $n = 6$, $P = 0.028$ (E-S'), respectively]. Interestingly, we also detected a certain number of boundary vector cells[4,55] which discharged further away from the edge of the enclosure (Supplementary information, Fig. S26). Further quantitative characterization of somatosensory boundary vector cells combined with the previously described "perimeter" coding might reveal how somatosensory boundary vector cell closely resemble their counterparts in the subiculum.[1,4,55]
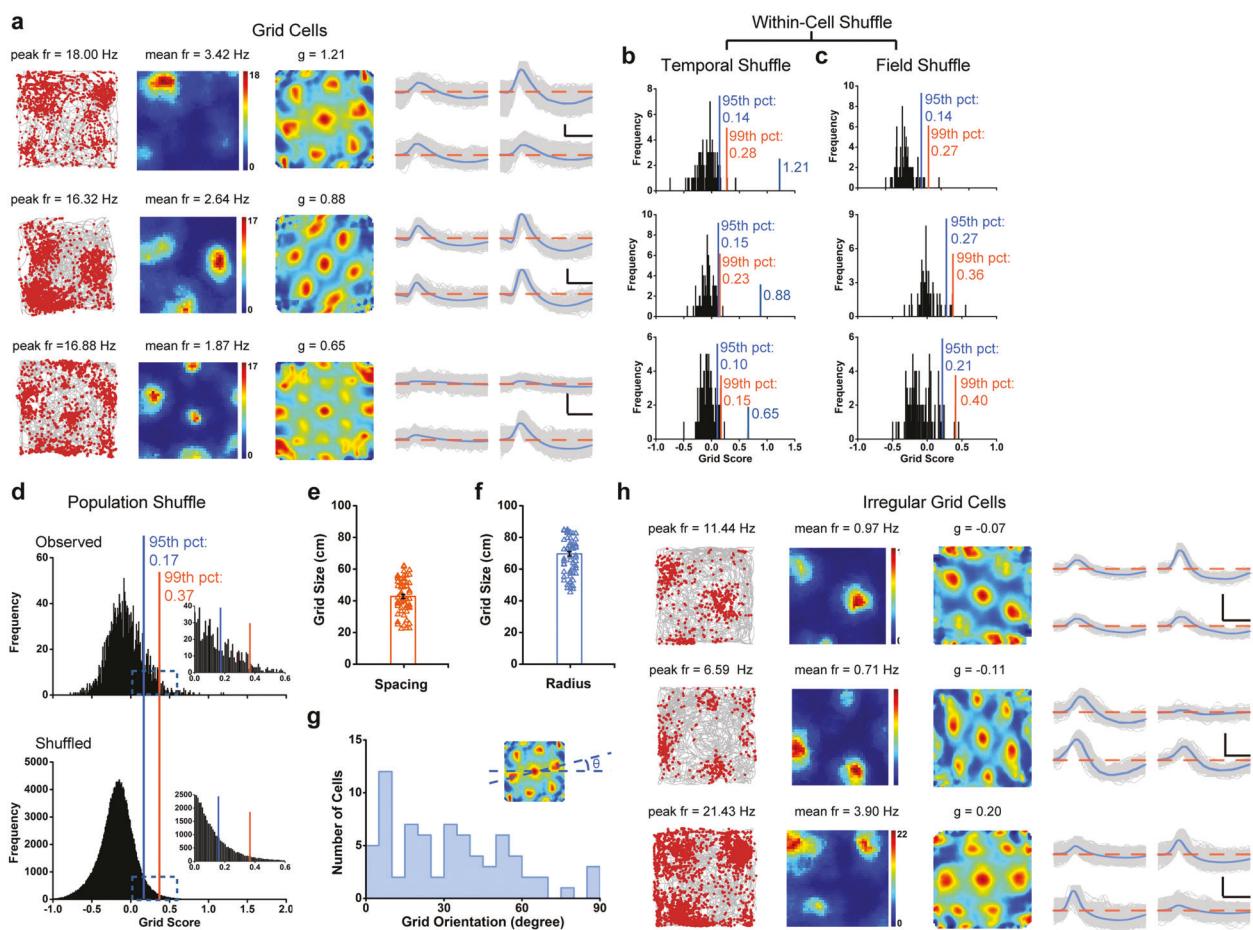
Grid cells in the somatosensory cortex
Hexagonal firing patterns identified in the MEC[6] can also be observed in the somatosensory cortex. To quantify the regularity of hexagonal firing patterns of the somatosensory cells (Fig. 4a; Supplementary information, Fig. S28), we calculated grid scores based on rotated autocorrelograms of rate maps. A cell was categorized as a grid cell if its grid score was higher than the 99th percentile of the shuffled data. 72/2025 cells (3.55%) met this criterion (Fig. 4d), and this percentage was significantly higher

than expected by random selection from the entire shuffled population (Fig. 4d; $Z = 12.90$, $P < 0.001$; binomial test with expected P_0 of 0.01 among large samples). Similar results were observed by performing within-cell shuffling (Fig. 4b). To eliminate the possible influence of inhomogeneous sampling on the multiple firing fields of somatosensory grid cells and verify S1 grid cells through spike shuffling, we performed additional "field shuffling"[56] and found that these exampled grid cells also passed the gridness threshold for the "field shuffling' procedure (Fig. 4c). The average firing rate and the grid score were 1.18 ± 0.12 Hz (Supplementary information, Fig. S8d) and 0.55 ± 0.02 (Supplementary information, Fig. S4d), respectively. The histograms of peak-to-peak amplitude and peak-to-trough spike width were presented in Supplementary information, Fig. S8h.

Grid spacing in somatosensory grid cells (Fig. 4e) was 42.72 ± 1.30 cm, and the average grid radius was 69.61 ± 0.16 cm (Fig. 4f). The fluctuation in somatosensory grid spacing is not a result of individual differences across different animals (Supplementary information, Fig. S4d). Moreover, the orientation of the somatosensory grid cells varied and was on the average found to be $32.63 \pm 2.64°$ (Fig. 4g). Again, with a lower threshold at the 95th percentile, 218/2025 S1 neurons (10.37%) were classified as grid cells, with the grid score threshold of 0.17 (blue line in Fig. 4d). When recorded in a larger environment (1.5 m × 1.5 m box), the discharging patterns of somatosensory grid cells showed similar multiple regular triangular structures in a larger enclosure to those within the smaller enclosure (1.0 m × 1.0 m) with a high grid score (Supplementary information, Fig. S29a–c), confirming that those identified grid cells were not false-positive grid cells due to accidental triangular node structures. It appeared that somatosensory grid cells were less prevalent and a little noisier than place cells, head direction cells and grid cells identified in the S1. It remains to be determined whether multisensory inputs and intrinsic oscillatory dynamics within the somatosensory cortex might generate different rate-coded grid computations from those within the hippocampal-entorhinal cortex. Taken together, the local configuration of grid cells in the somatosensory cortex exhibited similar features as those in MEC. Furthermore, we detected irregular grid cells with negative grid scores in the S1 along with regular grid cells (Fig. 4h; Supplementary information, Fig. S31). Since grid score calculated the 6-fold rotational symmetry of the cell's spatial autocorrelogram, a negative grid score means that the identified grid cells had a lower correlation for rotations of 60° and 120° than that for rotations of 30°, 60° and 120°.[7,17,57] Similar irregular grid cells such as elliptic grid cells were also reported in the MEC,[58,59] and both somatosensory regular and irregular grid cells were present side by side in the same recorded animals.

Conjunctive cells in the somatosensory cortex
To establish the existence of S1 neurons coding more than one spatial correlate, we assessed the strength of head direction tuning for all somatosensory place cells, border cells and grid cells.[7] A certain number of somatosensory neurons were found to

**Fig. 4 Grid cells in the somatosensory cortex. a** Three representative examples of somatosensory grid cells. Trajectory (gray line) with superimposed spike locations (red dots) (left column); heat maps of spatial firing rate (middle column) and autocorrelation diagrams (right column). Firing rate is color-coded with dark blue indicating minimal firing rate and dark red indicating maximal firing rate. The scale of the autocorrelation is twice the scale of the spatial firing rate maps. Peak firing rate (fr), mean firing rate (fr) and grid score (g) for each representative grid cell are labeled on the top of the panels. A crystal-like hexagonal firing pattern was observed. Spike waveforms on four electrodes are shown on the right column. The zero microvolt horizontal baseline is drawn with the orange dashed lines for the spike waveforms on all four electrodes. Scale bar, 100 µV, 300 µs. **b** Distribution of within-cell shuffled grid score for three representative somatosensory grid cells. The orange and blue stippled lines mark the 99th and the 95th percentile significance level of each randomly shuffled distribution, respectively. The cyan line indicates the observed grid score. **c** The same as (**b**) but for field shuffle. **d** Distribution of grid scores for somatosensory cells. The top panel shows the distribution for observed values. The bottom panel shows the distribution for randomly shuffled versions from the same sample. The orange and blue stippled lines mark the 99th and the 95th percentile significance level of each randomly shuffled distribution, respectively. Insets show the magnification of the specified area marked by the cyan dashed rectangle. **e, f** The raster plots show the distribution of grid spacing and grid size of the classified somatosensory grid cells. **g** The histogram of grid orientation from categorized somatosensory grid cells. **h** Three representative examples of irregular somatosensory grid cells. Spike waveforms on four electrodes are shown on the right column. Scale bar, 150 µV, 300 µs.

be conjunctive cells (Fig. 5; Supplementary information, Figs. S3a, S15b). Specifically, 17/195 place cells (Fig. 5a), 11/86 border cells (Fig. 5b) and 5/72 grid cells (Fig. 5c) were significantly tuned by head direction. Interestingly, we also detected irregular hexagonal grid cells modulated with head direction (Fig. 5d).
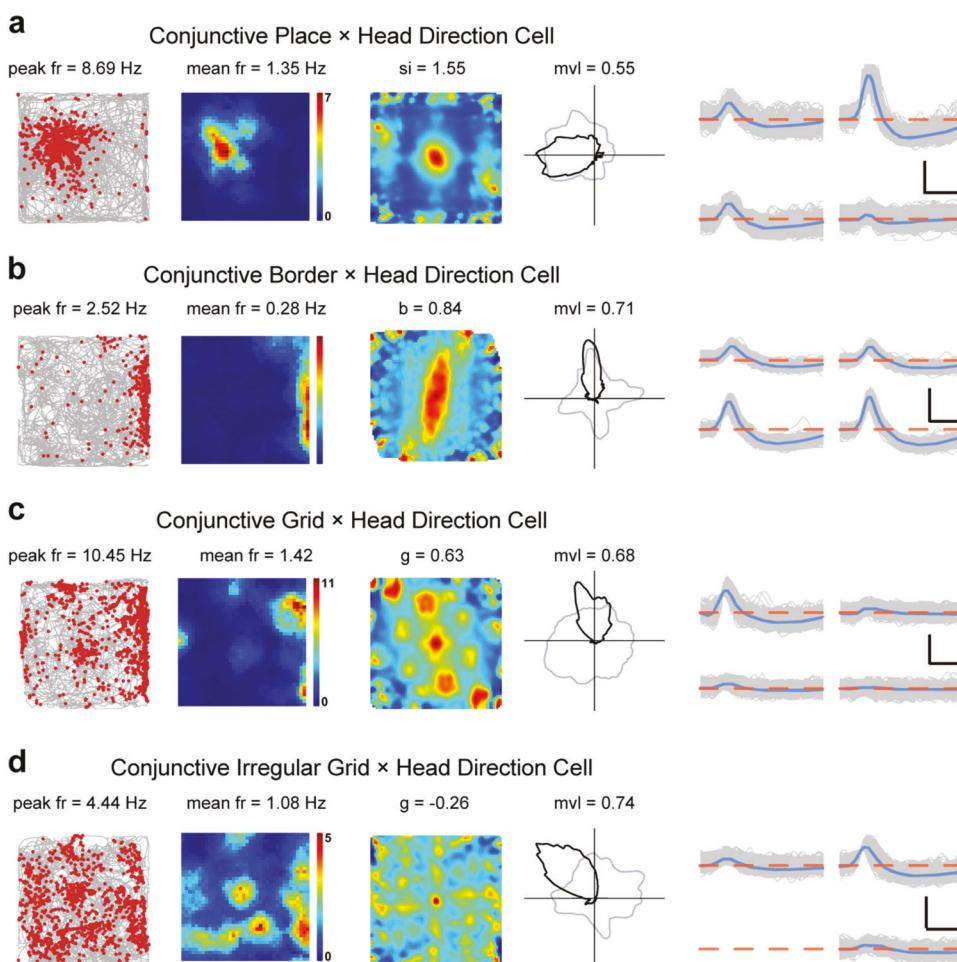
The degree of directionality in conjunctive cells showed no obvious difference from that of pure head direction cells (Watson's U^2-test, $P = 0.66$). These results suggested that the somatosensory cortex carried similar conjunctive place-by-direction, border-by-direction and gird-by-direction signal to those identified previously in the rat subiculum[46] and the rat MEC.[5,7]

Impact of the whisker trimming on the somatosensory spatial activities
Since rodents use their whiskers to actively explore the external environment for whisker-based navigation,[60,61] we implanted two

additional rats (Supplementary information, Fig. S1b) and recorded neuronal activity before and after repeated whisker trimming (Supplementary information, Fig. S32a, c) to gauge the involvement of the vibrissae. Two consecutive recording sessions from the same rat before and right after whisker trimming showed that the distribution of clusters and waveforms in the S1 was qualitatively similar (Supplementary information, Fig. S32b, d).

To further quantify the change of well-isolated spike clusters before and after whisker trimming, we calculated the isolation distance and L-ratio.[43] The median isolation distances were 122.35 and 132.9 before and after whisker trimming, respectively. There was no significant difference in cluster separation and L-ratio in the S1 before and right after whisker trimming ($n = 78$, $P = 0.73$ and 0.76, respectively, two-tailed paired t-test; Supplementary information, Fig. S32e, f). The average drift of center of mass for the total cell sample before and after whisker trimming was 0.13 ±

Fig. 5 Four different types of conjunctive cells in the somatosensory cortex. a A representative conjunctive place-by-head direction cell. **b** A representative conjunctive border-by-head direction cell. **c** A representative conjunctive grid-by-head direction cell. **d** A representative conjunctive irregular grid-by-head direction cell. Trajectory (gray line) with superimposed spike locations (red dots) (left column); heat maps of spatial firing rate (middle left column), autocorrelation diagrams (middle right column) and head direction tuning curves (black) plotted against dwell-time polar plot (gray) (right column). Firing rate is color-coded with dark blue indicating minimal firing rate and dark red indicating maximal firing rate. The scale of the autocorrelation is twice the scale of the spatial firing rate maps. Peak firing rate (fr), mean firing rate (fr), spatial information (si), border score (b), grid score (g) and mean vector length (mvl) for each representative cell are labeled on the top of the panels. Spike waveforms on four electrodes are shown on the right column. The zero microvolt horizontal baseline is drawn with the orange dashed lines for the spike waveforms on all four electrodes. Scale bar, 150 μV, 300 μs.

0.01, indicating there was minimal change in spike clusters before and after whisker trimming. Besides, both peak and average firing rates of the same neurons before and after whisker trimming did not change significantly ($n = 78$, $P = 0.75$ and 0.47, respectively, two-tailed paired t-test; Supplementary information, Fig. S32g, h).
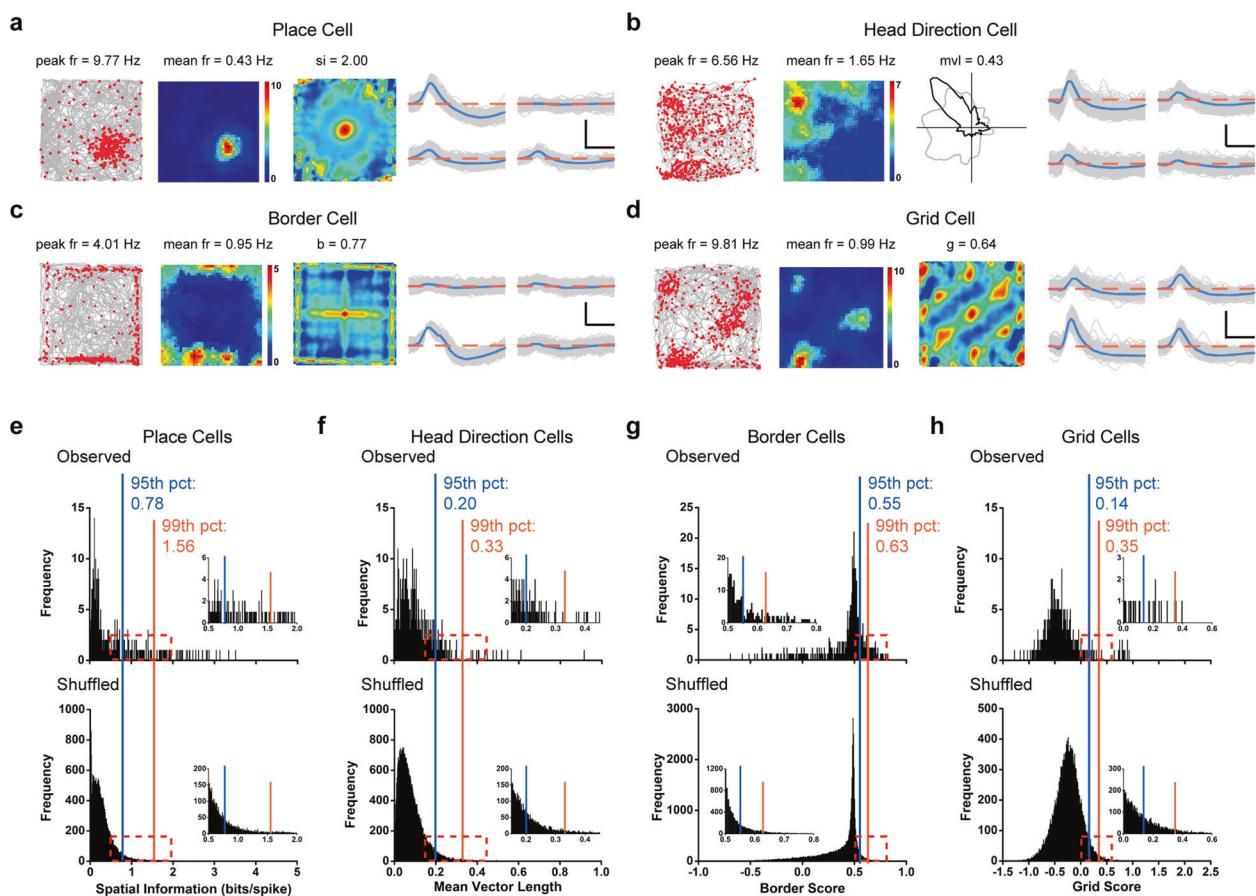
To further assess whether whisker-trimming affected the S1 spatial firing properties, we were able to continuously record the same S1 border cell before and after whisker trimming. The spatial firing patterns of the border cell persisted and responded to a new wall inserted into the square enclosure (Supplementary information, Fig. S33a, b). Overall, both the spatial responses and firing properties in the S1 were not significantly affected by the whisker trimming.

When we pooled and analyzed a total of 461 cells recorded after whisker trimming, we were still able to identify all different spatial cell types including 43 place cells (Fig. 6a), 15 head direction cells (Fig. 6b), 29 border cells (Fig. 6c) and 17 grid cells (Fig. 6d). The percentage of all recorded spatially tuned cell types after whisker-trimming were similar to those recorded from rats without whisker-trimming. Furthermore, the cut-offs for defining place cells (Fig. 6e), head direction cells (Fig. 6f), border cells

(Fig. 6g) and grid cells (Fig. 6h) after whisker-trimming highly resembled those for different somatosensory spatial cell types without whisker-trimming. Taken together, the persistence of recorded somatosensory spatial cells after whisker-trimming demonstrated that our identified spatial cells were not significantly influenced by the whisker-trimming.

DISCUSSION

This study provides the first demonstration and characterization of spatially tuned neuronal discharges in the primary somatosensory cortex. We have shown that place, grid, boundary vector/border, head direction and conjunctive cell activities, as reported in parahippocampal and associated areas,[39] co-exist in the single-domain primary somatosensory cortex. Given the current "blind" tetrode recording technique, we cannot be sure of the cellular sources of those distinct spatial signals described in the study. However, one clue is that the waveform amplitudes of identified somatosensory place cells were larger than those of other somatosensory grid, boundary vector/border, head direction and conjunctive cells, suggesting that somatosensory place cells may

Fig. 6 Somatosensory spatial response after whisker-trimming. a–d Four representative examples of somatosensory place cell (**a**), head direction cell (**b**), border cell (**c**) and grid cell (**d**) after whisker-trimming. Trajectory (gray line) with superimposed spike locations (red dots) (left column); spatial firing rate maps (middle left column), autocorrelation diagrams (middle right column) and head direction tuning curves (black) plotted against dwell-time polar plot (gray) (right column). Firing rate is color-coded with dark blue indicating minimal firing rate and dark red indicating maximal firing rate. The scale of the autocorrelation is twice the scale of the spatial firing rate maps. Peak firing rate (fr), mean firing rate (fr), spatial information (si), mean vector length (mvl), border score (b) or grid score (g) and angular peak rate for each representative cell are labeled on the top of the panels. Spike waveforms on four electrodes are shown on the right column. The zero microvolt horizontal baseline is drawn with the orange dashed lines for the spike waveforms on all four electrodes. Scale bar, 150 μV, 300 μs. **e–h** Distribution of spatial information (**e**), mean vector length (**f**), border score (**g**) or grid score (**h**) for somatosensory cells after whisker-trimming. The top panel shows the distribution for observed values. The bottom panel shows the distribution for randomly shuffled data from the same sample. The orange and blue stippled lines mark the 99th and the 95th percentile significance level of each randomly shuffled distribution, respectively. A zoomed panel shows the magnification of the specified area marked by the dark red dashed rectangle.

have larger cell bodies than those of other somatosensory spatial cells. These spatially tuned activities encoded within the newly identified somatosensory spatial navigation system appear to be specific, stable, anchored to external visual cues and robust against non-spatial environmental perturbations.

Based on the assumption that proprioception makes a significant contribution to spatial representation in the parahippocampal cortices,[39,40] we predicted spatially selective activities should exist in the somatosensory cortex. To our surprise, we were able to record the full complement of spatially selective cells (place, grid, boundary vector/border, head direction and conjunctive cells) in the primary somatosensory cortex. Given the paucity of direct projections from the parahippocampal areas to the S1 region,[62,63] we assume somatosensory spatial activities do not arise from the hippocampal-entorhinal circuit. However, we cannot completely eliminate the possibility that the somatosensory spatial positioning system may be at least partially dependent on the hippocampal-entorhinal system. Whether the newly found somatosensory spatial navigation system is independent of, parallel with or convergent onto the classical one within the hippocampal-entorhinal micro-circuit remains an interesting question and needs to be addressed

in future studies. The availability of single-cell transcriptomics and connectomics for the somatosensory cortex might help to unravel the interaction or interdependence between these two genetically pre-configured spatial maps.[64,65] Since there is very little overlap between septal projections to the hippocampal-entorhinal areas,[66] and that septal inactivation severely attenuates place and grid cell activities in both regions,[67–69] how S1 responds to septal inactivation may provide intriguing insights for hippocampal-entorhinal dependency.

The laminar organization of the S1 in relation to connectivity and function has been well-characterized, largely arising from studies on the vibrissae barrel fields.[70,71] There is insufficient information on the specific connectivity patterns of S1 in the rat, but mouse data suggest similar cortico-cortical, thalamic-cortical, cortical-thalamic, cortico-striatal and corticofugal connection patterns between S1 and other parts of the S1.[72] Our current study is roughly limited to sampling layers IV through VI. There is a pattern for head direction cells to be absent in layer VI and place cells to predominate in deeper layers (Supplementary information, Fig. S3c, d). From such distribution of spatial cell types across cortical depth, we could infer that head direction information is

Training and data collection

Behavioral training, tetrode turning and data recording started 1-week post-surgery. Rats were trained to run around in a 1 m × 1 m square box with a white cue card (297 mm × 210 mm) mounted on one side of the wall. Food pellets were scattered into the arena intermittently to encourage exploration.

Each recording session lasted between 10 and 30 min to facilitate full coverage of the testing arena. Tetrodes were lowered in steps of 25 or 50 μm daily until well-separated single units can be identified. Data were acquired by an Axona system (Axona Ltd., St. Albans, U.K.) at 48 kHz, band-passed between 0.8–6.7 kHz and a gain of × 5–18k. Spikes were digitized with 50 8-bit sample windows. Local field potentials were recorded from one of the electrodes with a low-pass filter (500 Hz).

Spike sorting, cell classification and rate map

Spike sorting was manually performed offline with TINT (Axona Ltd, St. Albans, U.K.), and the clustering was primarily based on features of the spike waveform (peak-to-trough amplitude and spike width), together with additional autocorrelations and cross-correlations.[45,87] During our manual cluster cutting, we always counted neurons with similar or identical waveform shapes only once whenever similar or identical individual neurons were recorded and tracked across two consecutive recording sessions. To confirm the quality of cluster separation, we calculated L-ratio as well as isolation distance between clusters.

Two small light-emitting diodes (LEDs) were mounted to the head-stage to track the rats' speed, position and orientation via an overhead video camera. Only spikes with instantaneous running speeds >2.5 cm/s were chosen for further analysis in order to exclude confounding behaviors such as immobility, grooming and rearing.

To classify firing fields and firing rate distributions, the position data were divided into 2.5 cm × 2.5 cm bins, and the path was smoothed with a 21-sample boxcar window filter (400 ms; 10 samples on each side).[7,17,44] Cells with > 100 spikes per session and with a coverage of > 80% were included for further analyses. Maps for spike numbers and spike times were smoothed with a quasi-Gaussian kernel over the neighboring 5 × 5 bins.[7,17,44] Spatial firing rates were calculated by dividing the smoothed map of spike numbers with spike times. The peak firing rate was defined as the highest rate in the corresponding bin in the spatial firing rate map. Mean firing rates were calculated from the whole session data.

Analysis of place cells

Spatial information is a quantification of the extent to which a neuron's firing pattern can predict the position of freely moving animals and is expressed in the unit of bits per spike. The spatial information was calculated as:

$$\text{spatial information} = \sum_i p_i \frac{\lambda_i}{\lambda} \log_2 \frac{\lambda_i}{\lambda},$$

where λ_i is the mean firing rate of the cell in the i-th bin, λ is the overall mean firing rate of the cell in the trial, and p_i is the probability for the animal being at the location of the i-th bin.

Adaptive smoothing[45] was applied to optimize the trade-off between spatial resolution and sampling error before the calculation of spatial information. The data were first divided into 2.5 cm × 2.5 cm bins, and then the firing rate within each bin was calculated by expanding a circle centered on the bin until

$$\gamma \geq \frac{\alpha}{n\sqrt{s}},$$

where γ is the circle's radius in bins, n is the number of occupancy of samples within the circle, s is the total number of spikes fired within the circle, and α is a constant parameter set to 10,000. With a position sampling frequency at 50 Hz, the firing rate assigned to that bin was then set to $50 \cdot s/n$.

A place cell was classified as a cell with the spatial information above chance level, which was computed by a random permutation process using all recorded cells. For each round of the shuffling process, the entire sequence of spike trains from each cell was time-shifted along the animal's trajectory by a random period between 20 s and the trial duration minus 20 s, with the end wrapped to the beginning of the trial. A spatial firing rate map was then constructed, and spatial information was calculated. This shuffling process was repeated 100 times for each cell, generating a total of 202,500 permutations for the 2025 somatosensory neurons. This shuffling procedure preserved the temporal firing characteristics in the unshuffled data while disrupting the spatial structure at the same time.

Spatial information score was then measured for each shuffled rate map. The distribution of spatial information values across all 100 permutations of all cells was computed and finally, the 99th percentile of the significant level was determined. The threshold values for categorizing cells into place cells were defined as the spatial information scores above the 99th percentile of the distribution from shuffled populations. In addition to the population shuffling, the within-cell shuffling was also performed within the entire sequence of spike trains from each individual neuron, and the shuffling process was also repeated 100 times for each single cell.

Spatial sparsity was used to measure how compact and selective the place field of each place cell is relative to the recording enclosure. The spatial sparsity was calculated using the formula as follows:[45]

$$\text{Sparsity} = \frac{\left(\sum p_i \lambda_i\right)^2}{\sum p_i \lambda_i^2}$$

Where p_i is the occupancy probability for the animal being at the location of the i-th bin in the map and λ_i is the mean firing rate of the cell in bin i.

Spatial coherence was estimated by calculating the mean correlation between the firing rate of each bin in the map and the aggregate firing rate of the eight nearest bins.[34] We used unsmoothed firing rate maps for computing the spatial coherence.

The spatial correlation across trials from the same recording arena was computed for each cell by correlating the firing rates in corresponding paired bins of two smoothed rate maps. The spatial stability within trials was estimated by calculating spatial (2D) correlations between firing rate maps generated from the first and second halves of the same trial. Place cells with spatial stability lower than 0.3 were excluded for further analysis.

Analysis of grid cells

Spatial autocorrelation was calculated with smoothed rate maps.[7,17,44] Autocorrelograms were derived from Pearson's product-moment correlation coefficient correcting for edge effects and unvisited locations.

With $\lambda(x, y)$ representing the average firing rate of a cell at coordinate (x, y), the autocorrelation between the spatial firing field itself and the spatial firing field with lags of τ_x and τ_y was calculated as:

$$r(\tau_x, \tau_y) = \frac{n\sum \lambda(x,y)\lambda(x - \tau_x, y - \tau_y) - \sum \lambda(x,y)\sum \lambda(x - \tau_x, y - \tau_y)}{\sqrt{n\sum \lambda(x,y)^2 - \left[\sum \lambda(x,y)\right]^2}\sqrt{n\sum \lambda(x - \tau_x, y - \tau_y)^2 - \left[\sum \lambda(x - \tau_x, y - \tau_y)\right]^2}},$$

where the summation is over whole n pixels in $\lambda(x, y)$ for which firing rate was calculated for both $\lambda(x, y)$ and $\lambda(x-\tau_x, y-\tau_y)$. Autocorrelations were not calculated for spatial lags of τ_x, τ_y where $n < 20$.

The degree of spatial regularity ("gridness" or "grid score") was calculated for each unit by using a circular sample centered on the central peak of the auto-correlogram but excluding the central

peak itself, and by comparing rotated versions of this circular sample.[7,17] The Pearson's correlations between this circular sample and its rotated versions were calculated, with the angles of rotation of 60° and 120° in the first group, and 30°, 90° and 150° in the second group. Gridness or the neuron's grid score was defined as the minimal difference between any of the coefficients in the first group and any of the coefficients in the second group. Shuffling was performed in the same procedure used for defining place cells. Grid cells were categorized as cells with the rotational symmetry-based grid scores exceeding the 99th percentile of the distribution of grid scores for shuffled data from the entire population of identified somatosensory cells.

Grid spacing was computed as the median distance from the grid center to the closest peak among six neighboring firing fields in the auto-correlogram of the spatial firing map. Since such analysis is sensitive to noise in the grid auto-correlogram, grid spacing was computed only when the median distance to the six neighboring peaks for the analyzed cell was comparable to the radius of the circle centered on the grid auto-correlogram with the highest grid score. The radius of this circle around the center of the auto-correlogram was also referred to as the grid field size.

Grid orientation was calculated by first computing vectors from the center of the auto-correlogram to each of the three adjacent peaks among six neighboring firing fields in the auto-correlogram of the spatial firing map in the counterclockwise direction, beginning with a camera-based reference line of zero degree. The angle between the minimal orientation of those three vectors and the camera-based reference line was defined as the grid orientation.

Analysis of head direction cells

The rat's head direction was estimated by the relative position of the LEDs differentiated through their sizes.[7,17,44] The directional tuning curve for each recorded cell was drawn by plotting the firing rate as a function of the rat's head angle, which is divided into bins of 3 degrees and then smoothed with a 14.5° mean window filter (2 bins on each side). To avoid bias, data were only used if all head angle bins contain data.

The strength of directionality was calculated by computing the mean vector length from circular distributed firing rates. The chance values were determined by a shuffling process simulated in the same way as for place cells, with the entire sequence of spike trains time-shifted between 20 s and the whole trail length minus 20 s along the animal's trajectory. Cells were defined as head direction cells if the mean vector lengths of the recorded cells were larger than the 99th percentile of the mean vector lengths in the shuffled distribution. Angular stability was computed by calculating the correlation of firing rates across directional bins generated from the first and second halves of the same trial. Head direction cells with angular stability lower than 0.3 were excluded for further analysis.

Analysis of border cells

Border or boundary vector cells were identified by calculating, for each recorded cell, the difference between the maximal length of any of the four walls touching on any single spatial firing field of the cell and the average distance of the firing field to the nearest wall, divided by the sum of those two values.[5,17,44] Border scores ranged from −1 for cells with perfect central firing fields to +1 for cells with firing fields that exactly line up with at least one entire wall. Firing fields were defined as summation of neighboring pixels with total firing rates higher than 0.3 times the cell's maximum firing rate that covered a total area of at least 200 cm^2.

Border cell classification was verified in the same way as for place cells, head direction cells and grid cells. For each permutation trial, the whole sequence of spike trains was time-shifted along the animal's trajectory by a random period between 20 s and 20 s less than the length of the entire trial, with the end wrapped to the start of the trial. A spatial firing rate map was then obtained, and a border score was estimated.

The distribution of border scores was calculated for the entire set of permutation trials from all recorded cells, and the 99th percentile was then determined. Cells were defined as border cells if the border score from the observed data was higher than the 99th percentile for border scores in the entire distribution generated from the permutated data.

Maximum-likelihood estimation of place cells and head direction cells

To evaluate inhomogeneous sampling biases on the spatial firing properties of place cells and head direction cells, we applied the maximum-likelihood correction algorithm previously published by Burgess and colleagues[47] and used the same analytical codes generously provided by original authors. Specifically, the position data were sorted into 2.5 cm × 2.5 cm bins and directional date into 120 bins of 3 degrees each before applying MLM correction technique. The field plot and polar plot are shown after smoothing with 5 boxcar × 5 boxcar smoothing and 5 boxcar smoothing, respectively.

Whisker-trimming

To test how the removal of another salient somatosensation mediate by vibrissae input impacts on FL/HL S1 spatial cells, we carefully trimmed all whiskers of each recoded rat bilaterally using blunt surgical scissors to within 1–2 mm above the skin surface just right before each recording session every day. Since bilateral whisker-trimmed rats did not exhibit abnormal behaviors, we continuously recorded somatosensory units from two whiskers-trimmed rats.

Environmental manipulations

For visual landmark rotation, we first recorded neuronal activity in the standard session followed by a 90° cue-card rotation in the clockwise or counterclockwise direction. Then another standard session was performed with the cue-card rotated back to the original position. For recording in the elevated platform without walls, we first recorded the somatosensory spatial cells in the square box, followed by the recording in the elevated platform without walls. Finally, the animals were returned to the original square box for another recording session. For the recording of border cells in the presence of inserted wall, we first identified the somatosensory border cells in the square box. Then the recording session was followed by the insertion of a wall along the center of the external wall. Another recording session was performed after removing the inserted wall. For food/no food comparisons, we recorded the somatosensory spatial cells in two consecutive recording sessions without and with throwing food pellets into the running enclosure.

Histology and reconstruction of recording positions

At the end of the experiment, rats were euthanized with an overdose of sodium pentobarbital and transcardially perfused with phosphate-buffered saline (PBS) followed by 4% paraformaldehyde (PFA). Brains were removed and stored in 4% PFA overnight. The brain was then placed in 10%, 20% and 30% sucrose/PFA solution sequentially across 72 h before sectioning using a cyrotome. Thirty-micron sections were obtained through the implant region. Sections were mounted on glass slides and stained with cresyl violet (Sigma-Aldrich). The final recording positions were determined from digitized images of the Nissl-stained sections scanned on the Olympus Slideview VS200 Digital Slide Scanner. Positions of each individual recordings were estimated from the deepest tetrode track, notes on tetrode advancement with tissue shrinkage correction by dividing the distance between the brain surface and electrode tips by the last advanced depth of the recording electrodes. Electrode traces were

confirmed to be located within the hindlimb region (S1HL) from eight implanted rats, the forelimb region (S1FL) from one rat and the shoulder region (S1Sh) from another rat but all electrode tracking paths from 10 implanted rats were verified to be away from the barrel field (S1BF) of the primary somatosensory cortex according to the Rat Brain Atlas.[88]

DATA AVAILABILITY
Recording dataset will be prepared available in a forthcoming public domain, and inquiries into acquiring the recording dataset beforehand should be directed to the corresponding author.

AUTHOR CONTRIBUTIONS
S.J.Z. conceived and designed the study. X.L. and S.J.Z. performed the experiments, collected the data and performed the analyses. X.L. made the figures. S.J.Z. wrote the manuscript.

ADDITIONAL INFORMATION
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