

A cellular resolution map of barrel cortex activity during tactile behavior

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Abstract

Comprehensive measurement of neural activity remains challenging due to the large numbers of neurons in each brain area. We used volumetric calcium imaging in mice expressing GCaMP6s and nuclear red fluorescent proteins to sample activity in 75 % of superficial barrel cortex neurons across the relevant cortical columns, approximately 12,000 neurons per animal, during performance of a single whisker object localization task. Task-related activity peaked during object palpation. An encoding model related activity to behavioral variables. In the column corresponding to the spared whisker, 300 layer (L) 2/3 pyramidal neurons (17 %) each encoded touch and whisker movements. Touch representation declined by half in surrounding columns; whisker movement representation was unchanged. Following the emergence of stereotyped task-related movement, sensory representations showed no measurable plasticity. Touch direction was topographically organized, with distinct organization for passive and active touch. Our work reveals sparse and spatially intermingled representations of multiple tactile features.

Introduction

Neural coding is a population-level phenomenon, with even simple choice behaviors involving large numbers of neurons in multiple brain areas (Guo et al., 2014b; Romo, 2013). Due to technical limitations, neurophysiological recordings typically sample only a small subset of neurons in a cortical region (Stevenson and Kording, 2011). Incomplete sampling limits our understanding of neural representations and their relationship to neural circuit structure. Two-photon microscopy (Denk et al., 1994; Svoboda and Yasuda, 2006) has been used to image the activity of populations of individual neurons in anesthetized (Kerr et al., 2007; Kerr et al., 2005; Ohki et al., 2006; Sato et al., 2007; Stosiek et al., 2003) and behaving (Andermann et al., 2010; Dombeck et al., 2007; Huber et al., 2012; Komiyama et al., 2010) animals. Recent advances in genetically encoded calcium indicators (GECIs; (Chen et al., 2013b; Nagai et al., 2004; Tian et al., 2009)) permit sensitive detection of neural activity in individual neurons and tracking of activity in neural populations across days or weeks (Huber et al., 2012; Margolis et al., 2012; Peters et al., 2014). Two-photon microscopy can sample neurons densely, localize them, and measure activity in genetic and projection classes (Chen et al., 2013a; Chen et al., 2013b; Sato and Svoboda, 2010). To date, cellular imaging in the mammalian brain has been limited to hundreds of neurons in small tissue volumes.

Mice move their whiskers over objects to localize and recognize them (Diamond et al., 2008). The vibrissal primary somatosensory cortex (vS1, or ‘barrel cortex’) contains a somatotopic map of the large facial whiskers (Woolsey and Van der Loos, 1970). Tactile information from individual whiskers is processed in single barrel columns in vS1 (Feldmeyer et al., 2013; Simons, 1978). Each column (diameter, 300 μm) contains approximately 10,000 neurons (Hooks et al., 2011; Lefort et al., 2009). Rodents can localize objects and walls under head-fixed (O’Connor et al.; Sofroniew et al.) and freely moving (Hutson and Masterton, 1986; Knutsen et al., 2006) conditions using a single whisker. Single barrel columns thus provide a defined target for comprehensive and dense, cellular-resolution imaging.

Vibrissal S1 comprises distinct layers, with each layer harboring neuron types with distinct inputs and outputs. Input from VPM primarily targets L4, with minor projections to L3, L5B and L6. Input from POM terminates in L5A and L1 (Lu and Lin, 1993; Petreanu et al., 2009; Wimmer et al., 2010). Other cortical areas send projections to specific laminae in vS1. Vibrissal motor cortex axons terminate in L6 and L1, synapsing onto L2/3, L5, and L6 neurons (Kinnischtzke et al., 2014; Petreanu et al., 2009). Within the cortex, L4 neurons project to L3, and L3 and L5A neurons project to L2 (Feldmeyer, 2012; Staiger et al., 2014). L2 and L3 neurons project to L5 and other parts of the neocortex. L2/3 is therefore a site of integration of ascending sensory input and top-down modulation from higher cortical areas.

Vibrissal S1 neurons are sensitive to whisker deflections (Ahissar et al., 2001; Armstrong-James et al., 1992; Kerr et al., 2007; Sato et al., 2007; Simons, 1978), with deflection direction mapped topographically in superficial cortical layers (Andermann and Moore, 2006; Kremer et al., 2011). During active behavior, vS1 is modulated by

object touch and whisker movement (Crochet et al., 2011; Curtis and Kleinfeld, 2009; de Kock and Sakmann, 2009; Krupa et al., 2004; O'Connor et al., 2010b; Petersen and Crochet, 2013). However, little is known about the prevalence of different sensory representations and their spatial distribution within and across barrel columns.

We used volumetric two-photon laser scanning microscopy, combined with expression of GCaMP6s (Chen et al., 2013b), to measure behavior-related activity in more than 10,000 neurons per mouse in and around the barrel column corresponding to the whisker used by mice to solve a tactile task. This includes the majority of neurons in superficial vS1.

Using an encoding model, we quantified the contribution of behavioral variables to the activity of individual vS1 neurons. We thereby constructed a cellular resolution map of behavior-related activity for vS1.

Results

Imaging large neuronal populations in behaving mice

Head-fixed mice were trained to perform an object localization task with a single whisker. A pole was either presented in a range of proximal positions that predicted a reward in the right lickport (of two) or in a distal position, which predicted reward in the left lickport (Guo et al., 2014a; Guo et al., 2014b) (**Figure 1A**). Mice had to make a decision about object location and hold this decision in memory during a delay epoch before signaling their decision by licking. We used videography and automated whisker tracking (**Figure 1B**) to measure whisker movements and tactile input (**Figure 1C**) (Clack et al., 2012; Pammer et al., 2013). Mice searched for the pole with their whisker, mainly towards the end of the sample epoch (**Figure 1C**). Whisker position was measured as the azimuthal angle of the spared vibrissa (θ). As a measure of tactile input we extracted the touch-induced changes in whisker curvature ($\Delta\kappa$), which are proportional to the forces acting on mechanoreceptors in the follicle (Birdwell et al., 2007; Pammer et al., 2013).

In most cases, imaging began after mice attained stable behavior ('Volume imaging', **Figure 1D**, **Table S1**). A subset of mice was also imaged during training ('Imaging during learning'). Mice reached criterion level performance (d -prime > 1.5, or approximately 70% trials correct; **Figure 1E**) after 11 ± 4 days (mean \pm S.D.; $n = 8$ mice) of training. Whisker movements became stereotyped early during training (**Figure 1G**; Methods; (Huber et al., 2012)), whereas behavioral performance increased more gradually (**Figure 1E**). Mice performed 245 ± 71 trials per session during imaging. Compared to tasks with two pole locations (Guo et al., 2014b), our task elicited larger amplitude whisker movements (approximately 20° vs. $55.5 \pm 13.3^\circ$; mean \pm S.D.; $n = 8$ mice). A large range of whisking angles and whisker curvatures were sampled during behavior, allowing us to relate neural activity to the dynamics of whiskers.

We performed large-scale calcium imaging in vS1 within the column of the spared whisker (principal column) and its neighbors (**Figure 2A**; **Figure S1**). In most experiments (**Table S1**), mice expressed a red fluorescent protein (mCherry, (Shaner et al., 2004)) in the nuclei of cortical glutamatergic neurons (emx1-cre x Rosa26-LSL-CAG-H2B-mCherry; **Figure S2**). One mouse expressed mCherry in the nuclei of GABAergic neurons (Gad2-T2A-NLS-mCherry). The labeling allowed us to distinguish glutamatergic ('excitatory') and GABAergic neurons. In addition, the red nuclear fluorescence was used to detect neurons in images independent of functional signals. We infected all neurons in and around the principal column with AAV2/1 syn-GCaMP6s, yielding labeled neurons in layers 1, 2, 3, 5, but not layer 4 (**Figure 2B**). A cranial window was placed over the infected area (Huber et al., 2012; Trachtenberg et al.).

Following viral injection, mice were water restricted and trained on the pole localization task (Guo et al., 2014a). Two-photon excitation with 1000 nm light produced emission from both GCaMP6s (green) and the nuclear mCherry (red) (**Figure 2A,C**). Laser

scanning in the plane was performed with a custom resonant scanning system (line frequency 16 kHz) (Fan et al., 1999), with axial scanning controlled by a piezo collar. For each trial (approximately 10 s), we imaged subvolumes comprising three planes ($600 \times 600 \mu\text{m}^2$; 512x512 pixels) 15 μm apart in depth. Each subvolume was imaged at 7 Hz for 32 ± 7 trials, followed by another subvolume, and so on. Over approximately two behavioral sessions, we visited 16 subvolumes, spanning 6-8 barrel columns. Each subvolume was imaged during 4.0 ± 1.2 behavioral sessions.

Regions of interest (ROIs) were drawn in a semi-automated manner around individual somata, aided by the nuclear mCherry fluorescence (**Figure 2D**; see Methods). Our core dataset comprised 82,732 excitatory L2/3 neurons, 777 L1 neurons, and 3,806 L2/3 GABAergic neurons (n=8 mice; **Table S1**). The mCherry labeling allowed us to quantify the fraction of recorded excitatory L2/3 neurons. Red nuclei were counted in high-resolution stacks of the imaged volumes (**Figure S2**; see Methods). After aligning the imaging planes to these stacks, we estimated the fraction of excitatory neurons for which ROIs had been defined. In L2/3, 76 ± 6 % of red nuclei had corresponding ROIs, implying that we had recorded from $\sim 3/4$ of the excitatory neurons in L2/3 (**Figure 2E**). Our count for L2/3 pyramidal neurons per barrel column (1796 ± 299 ; n = 5 volumes, see Methods) is in agreement with reported neuronal counts for mouse vS1 (Hooks et al., 2011; Lefort et al., 2009).

In most mice, GABAergic neurons were analyzed based on GCaMP6s fluorescence alone. Since fluorescence increases with neural activity our sampling was biased toward active GABAergic neurons. We found 3,806 GABAergic neurons among 86,538 neurons in L2/3 (4.3 %). Given that approximately 15% of neurons in L2/3 are GABAergic (Lefort et al., 2009) this implies that we are recording from 1/3 of the GABAergic neurons present in our imaging planes.

Viral transfections produce densely labeled neuropil, consisting of GCaMP6s-expressing axons and dendrites, showing behavior-related activity (**Figure S3**). Because of the limited resolution of 2-photon microscopy *in vivo*, especially axially (Ji et al., 2012), the neuropil signal bleeds into the somatic signal (Chen et al., 2013b; Ji et al., 2012). We corrected for neuropil contamination by subtracting the local, peri-somatic neuropil signal (**Figure 2F**; **Figure S3**; see Methods) (Kerlin et al., 2010) and computed $\Delta F/F$ for each neuron (**Figure 2G**). Fluorescence events corresponding to neural activity were extracted using a greedy template-fitting algorithm, similar to the peeling method described in (Lutcke et al., 2013), but incorporating the variable decay time constants of GCaMP6s (see Methods; **Figure 2H**). Events were then convolved with their respective rise and decay times, to generate a de-noised, event-based $\Delta F/F$ trace. All subsequent analyses employed this event-based $\Delta F/F$ trace, unless noted.

In deep L3, the apical dendrites of L5 neurons appeared as small, high contrast circles, with large fluorescence transients (**Figure S4**). We verified that $\Delta F/F$ in apical dendrites reflected somatic activity by simultaneously imaging apical dendrites in deep L3 and the corresponding somata in L5A. First, manual inspection of 845 events (n=42 dendrite-soma pairs, n=3 mice) revealed that 91.0 ± 15.1 % of somatic events (mean \pm S.D) had

corresponding dendritic events, and $90.9 \pm 12.7\%$ of dendritic events had corresponding somatic events. Second, the correlation between the somatic and dendritic signals was 0.63 ± 0.11 (Pearson's R; $p < 0.001$, $n = 42$ pairs). This correlation has to be interpreted in the context of the different fluorescence dynamics in the two compartments.

Specifically, dendritic responses were larger and briefer compared to somatic transients (**Figure S4**; (Helmchen et al., 1999; Hill et al., 2013; Svoboda et al., 1997; Svoboda et al., 1999)). We computed the correlation in a model with perfect correspondence between somatic and dendritic events, where the events were convolved with compartment-specific kernels (to reflect different amplitude and kinetics) and compartment-specific noise. This estimate, $R = 0.69 \pm 0.28$ (mean \pm S.D., correlation $p < 0.001$; $n = 42$ pairs), was similar to the measured correlation (Wilcoxon signed rank test, $p = 0.063$). Thus activity measured in L5 dendrites measured in L3 reports activity of L5 somata. We imaged a total of 2,469 apical dendrites as a proxy for L5A somatic activity. Based on estimates of neuronal density in L5A (Lefort et al., 2009), we recorded from less than 10% of these neurons.

Task-related activity

Our goal was to understand how the coding of behavioral variables is distributed across neurons in the barrel cortex. The majority of neurons were not silent (event rate > 0.0083 Hz; **Figure 3A**). The largest proportion of active neurons was seen in L2/3 excitatory neurons ($67 \pm 5.4\%$; $n=8$ mice), in quantitative agreement with electrophysiological recordings (O'Connor et al., 2010b).

To determine if neural activity was task-related we examined whether activity was temporally locked to the trial (**Figure 4A, B**; see Methods). Overall, 40% of neurons showed task-related activity (**Figure 3B**). L2/3 excitatory neurons showed the lowest proportion of task-related neurons ($39 \pm 2.9\%$), L5 excitatory neurons were intermediate ($43 \pm 4.1\%$), and GABAergic neurons had the highest proportion of task-related neurons, both in L1 and L2/3 (57 ± 12 and $63 \pm 7\%$, respectively).

The temporal dynamics of L2/3 neurons were diverse. Across the population, activity spanned the behavioral trial (**Figure 4C**), with a pronounced bias toward the sample epoch during which the animal interacted with the stimulus. In the principal barrel column, activity in the sample epoch was substantially elevated relative to surround columns for all excitatory neuron classes examined (**Figure 4D**), consistent with somatotopically organized touch input. GABAergic neurons showed distinct dynamics. In addition to a response peak during the sample epoch, GABAergic neurons in L1 and L2/3 showed a pronounced peak in activity around the reward cue, which was especially prominent in surround columns due to lack of touch input (**Figure 4D**).

Activity in vS1 during the sample epoch is necessary for pole localization (Guo et al., 2014b). The task-related activity observed during the sample epoch may therefore be used by mice to judge object location. To assess the discriminative capacity of individual neurons, we measured how neurons differentiate trial types using receiver operating characteristic analysis (see Methods). Only activity up to the reward cue was used.

Among L2/3 excitatory neurons in the principal column, $9.7 \pm 8.5\%$ discriminated trial type above chance level ($n=8$ mice; **Figure 3C**). L5 excitatory ($17.3 \pm 7.1\%$) and L2/3 GABAergic neurons ($20.7 \pm 11.5\%$) performed better. A majority of neuropil ROIs were discriminative ($56.1 \pm 31.4\%$). Thus, representation of task-related parameters in L2/3 excitatory neurons of vS1 is sparse (Barth and Poulet, 2012).

Encoding model of neural coding

Vibrissal S1 neurons encode aspects of forces acting on the whisker and whisker movements (Crochet et al., 2011; de Kock and Sakmann, 2009; Fee et al., 1997; O'Connor et al., 2013; O'Connor et al., 2010b). To investigate which behavioral parameters are represented by individual vS1 neurons during active somatosensation, we quantified how whisker movements and touches contribute to neural activity.

The data were fit using a cascaded generalized linear model with input nonlinearities (Ahrens et al., 2008; Paninski et al., 2004). A receptive field, modeled as a piecewise static non-linearity applied to each input variable, describes the relationship of each variable to neural activity. A linear temporal kernel models the temporal dynamics of the neuron, including the time-course of calcium transients (**Figure 5A**). The model was fit to each neuron by maximizing the probability of the response as a function of the behavioral input variables (see Methods). The behavioral data consisted of whisker movements or curvature changes collated across multiple sessions (128 ± 30 behavioral trials per neuron; **Figure 5B**). To quantify model performance, the response predicted by the model for each neuron was compared to the actual response using Pearson's correlation (R_{fit}). If a behavioral variable predicted neuronal $\Delta F/F$ with an R_{fit} that exceeded the 95th percentile of shuffled R_{fit} s, the cell was classified as encoding that behavioral variable (**Figure 5C-E**; see Methods).

Touch and whisking neuron responses were aligned to the first touch and whisking bout (see Methods) of a trial, respectively (**Figure 5F**). Layer 2/3 excitatory neurons were equally likely to show representation of whisking and touch within the principal column ($17.0 \pm 5.2\%$ and $16.8 \pm 9.0\%$, respectively; $p=0.938$, Wilcoxon signed rank test, $n=8$ mice) (**Figure 3D**). In surround columns, whisking representation dominated ($15.0 \pm 3.9\%$ whisking vs. $8.9 \pm 2.0\%$ touch; $p=0.008$). Overall, about a quarter of L2/3 excitatory neurons encoded whisker movement or touch, the lowest proportion observed for any group. Among L2/3 GABAergic neurons as well as L5 excitatory neurons, whisker movements were represented by a larger proportion of neurons compared to touch. The neuropil was dominated by touch, but also showed detectable encoding of whisker movements in the majority of ROIs.

Imaging during learning

We examined the dynamics of neural representations during learning of the task (**Figure 6A**). In four mice (**Table S1**), we imaged a subset of L2/3 excitatory neurons spanning the principal column and portions of adjacent columns from the onset of training until expert performance was achieved (7, 10, 13, and 17 sessions) (**Figure 1D, E**).

We first asked whether the fraction of touch and whisking neurons changed during learning (**Figure 6B**). The fraction of touch neurons was constant from the start or middle of training relative to the end of training (days 1, 2 touch fraction: 0.15 ± 0.05 , mean \pm SD, n=4 mice, 2 days per mouse; 5 and 4 days before end: 0.14 ± 0.06 ; final two days: 0.15 ± 0.07 ; first two vs. final two p = 1, Wilcoxon rank sum test; middle vs. final p = 0.573). The fraction of whisking neurons increased from the start of training to the final training day. This change occurred early during training (days 1, 2 whisking fraction: 0.13 ± 0.04 ; 5 and 4 days before end: 0.20 ± 0.09 ; final two days: 0.19 ± 0.05 ; first vs. final p-value = 0.010; middle vs. final p-value = 0.798). This early increase in the number of whisking neurons parallels early changes in task-related movement (**Figure 1F, G**), suggesting that changing behavior rather than neural plasticity underlies the observed dynamics.

We next looked for functional expansion of the spared whisker representation following trimming, a classic measure of receptive field plasticity. We compared the fraction of neurons representing touch by the eventual spared whisker inside and outside the principal column after trimming to a single whisker (n = 3 mice, all imaged for at least 10 days after trimming; **Figure 6C**). We found no change in the ratio of the number of touch neurons (inside-to-outside) following trimming (days 0 and 1: 1.29 ± 0.31 ; days 4 and 5: 1.55 ± 0.71 ; days 9 and 10: 1.31 ± 0.39 ; early vs. middle: p = 1; early vs. late p = 0.937). Thus, we did not detect enhanced representation of the spared whisker.

Improved task performance during training (**Figure 1E**) could be a consequence of enhanced representation of object location in vS1, or improved readout by downstream brain areas. We measured discrimination by vS1 neurons during learning. Discrimination by small L2/3 ensembles (10 individually discriminative neurons; Methods) exceeded behavioral performance (O'Connor et al., 2010b), even during early training (**Figure 6D**). Neurometric performance remained flat during learning (first two days: $75 \pm 4\%$, middle two days: $75 \pm 5\%$, final two days: $78 \pm 5\%$; first two vs. middle two: p = 0.959, first two vs. final two: p = 0.721 ; middle two vs. final two: 0.505). In contrast, psychometric performance improved gradually from chance level to criterion levels over the course of training (first two days: $52 \pm 2\%$; final two days: $66 \pm 7\%$). This indicates that at the outset, suitable tactile information is present in vS1 to perform the task, but is not used by the animal; following learning, mice interpret this information more effectively to gather reward.

We next asked how the representation of touch and whisking evolved at the single neuron level. Individual L2/3 neuron representations stabilized after the first few days of training (**Figure 6E**): population R_{fit} vectors from early training days were poor predictors of R_{fit} vectors on subsequent days (quantified using Pearson's R), whereas the R_{fit} vectors in later training were better predictors of R_{fit} vectors of subsequent days (**Figure 6F, G**).

Prediction was imperfect across sessions even late in training (i.e., correlation between R_{fit} vectors < 1). This could reflect genuine plasticity in representations, or lack of

sensitivity when comparing encoding model fits across days. To disambiguate these possibilities, we computed a distribution of correlation values from data without plasticity. Each session was partitioned trial-wise into two equal-length, temporally interdigitated ‘pseudo sessions’. The encoding model was fit to each ‘pseudo session’ independently, and the correlation between the two resulting R_{fit} vectors was measured. By repeating this 25 times per session, a distribution of intra-day correlation values was obtained. For a given imaging day, subsequent days were considered significantly distinct if the correlation of the R_{fit} vectors fell below the 5th percentile of intra-day correlations for that day.

By this measure, touch representations stabilized during training (**Figure 6H**). Relative to the first day of imaging $77 \pm 26\%$ (n=4 mice) of subsequent days were different, whereas 5 days prior to the final training day, only $17 \pm 33\%$ of subsequent days were different. Whisking representations also stabilized. Relative to the first day, $60 \pm 43\%$ of subsequent days were different, falling to $33 \pm 36\%$ 5 days before the end of training.

Our data show that encoding as well as discriminative performance stabilizes after the first few days of training, both at the level of single neurons and populations. The time course of stabilization mirrors the time course of the emergence of motor stereotypy (**Figure 1F, G**), rather than improvement of behavioral performance (**Figure 1E**). The stability of representations after the first few days of training justifies sampling across behavioral sessions to characterize large numbers of vS1 neurons in well-trained mice.

Spatial organization of sensory coding

We next investigated how the encoding of behavioral variables is organized spatially. Among L2/3 pyramidal neurons, touch and whisking cells were intermingled in a salt and pepper manner within the principal barrel column and also in the surrounding columns (**Figure 7A, B**). The proportion of L2/3 excitatory neurons encoding touch declined by half when moving from the principal column to the surround columns (**Figure 7B, C**; $p = 0.008$, Wilcoxon signed rank test, n=8 mice). A similar spatial profile was observed in L1 ($p = 0.031$). In contrast, touch-encoding L5 neurons ($p = 0.383$) and L2/3 GABAergic neurons ($p = 0.578$) were equally likely in the center and surround columns (**Figure 3D**). Touch thus activates L2/3 GABAergic neurons over multiple columns, producing an inhibitory surround in L2/3 (Derdikman et al., 2003). Representation of whisking was uniform across neurons inside and outside the principal column. Thus, touch neurons were primarily confined to the principal column, whereas whisking neurons were distributed uniformly.

The L2/3 neuropil signal had different dynamics and structure from L2/3 somata (**Figure 4C, D; Figure S3**). The neuropil signal was temporally more concentrated in the sample epoch and spatially more concentrated in the principal column. This suggests that it mostly reflects active touch. The neuropil signal did not directly reflect activity in L4 axons as suggested previously (Kerr et al., 2005) because L4 neurons were not expressing GCaMP6 infection in our preparation. We measured L4 axon activity in a separate set of mice (**Figure S5**). The spatial distribution of neuropil activity most closely resembled L4

activity. Specifically, the correlation between L4 axon and L2/3 neuropil touch maps was highest of the three possible pairings (L4-neuropil Pearson's R: 0.60 ; L4-L2/3 somata: 0.41; L2/3 somata-neuropil: 0.36). Temporally, L4 axons and L2/3 neuropil both had a large fraction of activity confined to the sample epoch (**Figure 4C, D**). This suggests that the neuropil corresponds to postsynaptic calcium dynamics in L2/3 neuron dendrites, independent of somatic spikes. Consistent with this interpretation, all L2/3 neurons show strong subthreshold responses to touch (Crochet et al., 2011).

We next examined the spatial organization of encoded variables at the fine scale. We found no difference when comparing the distribution of pairwise distances among neurons of a category (touch, whisking, mixed) to the pairwise distance distribution among all neurons (**Figure S6**). Thus, different representations were randomly intermixed in L2/3 within a column, but the touch representation showed the expected somatotopic organization on the scale of multiple columns (Kerr et al., 2007; Sato et al., 2007) (**Figure 7B**).

Directional tuning in vS1

Directional tuning maps have been observed in rat vS1 (Kremer et al., 2011) (Andermann and Moore, 2006). Our encoding model yields a description of how response amplitude varies as a function of whisker curvature (**Figure 5A, C, D**). Because the sign of curvature is different for protraction and retraction touches (negative and positive, respectively), the receptive fields reflect direction preference, allowing us to examine the topography of directional tuning in vS1.

The majority of L2/3 touch neurons were directionally tuned (**Figure 8A, B**), with an approximately equal number of neurons preferring protraction and retraction. Directional tuning was typically strong, with neurons either responding almost exclusively to protraction (directionality index, DI = 1) or retraction (DI = -1) touches. Relatively few neurons showed an intermediate level of tuning. To assess stability of directional tuning over time, we measured the average pairwise root mean square (RMS) difference among touch receptive fields in the animals imaged during training (n=4 mice). The difference between touch receptive fields among days (grand mean of RMS difference: 0.273; n = 118 neurons) was below the 5th percentile of the shuffled distribution (0.425; shuffling by neuron identity) for 97.8 % of the neurons examined (**Figure S4**). Thus, L2/3 directional tuning was stable over the course of learning.

Consistent with observations in anesthetized rats (Andermann and Moore, 2006; Kremer et al., 2011), neurons sensitive to protraction touch were more abundant in and near the barrel column of the whisker anterior to the spared one, with retraction sensitive neurons more abundant in and near the column of the whisker posterior to the spared one (**Figure 8C, D, E**). This trend in direction selectivity within a barrel row (**Figure 8E**) could be a consequence of neurons becoming more biased (increasing DI magnitude) with distance from the principal column, or it could reflect changes in relative fractions of comparably biased neurons (constant DI magnitude). DI magnitudes inside and outside the principal

column were similar ($p = 0.333$, Wilcoxon rank sum test), implying that the spatial trend reflects a change in overall directional preference of neurons with similar DI magnitude.

Prior examination of directional tuning using extracellular recordings failed to reveal topographic organization in L4 (Andermann and Moore, 2006). In contrast, we observed a directional map, similar to that observed in L2/3 excitatory neurons, in the activity of both L4 axons and the L2/3 neuropil (**Figure 8E**). Thus, at least in mice, the directional topography of L2/3 is likely inherited from L4.

Previous work in anesthetized rats showed that forward whisker deflection produced stronger responses in the column of the whisker immediately anterior to that of the deflected one (Andermann and Moore, 2006; Kremer et al., 2011). Active touches during protraction (whisker movement in the anterior direction) produce negative curvature changes, whereas passive whisker deflection in the anterior direction produces positive curvature changes. Despite this difference in curvature sign, both cases evoke the most response among neurons in and near the column of the whisker anterior to the spared one (**Figure 8F**). Since the change in curvature is proportional to the lateral force exerted on the whisker, this implies that the somatotopy in active and passive touch is reversed in terms of lateral forces.

To explore this discrepancy we compared neural responses to passive touch in anesthetized mice and during active tactile behavior ($n=3$ additional mice). In anesthetized mice, whiskers were stimulated with a pole mounted on a piezo (see Methods). We found that most neurons responded to touch in only one or the other condition, and rarely both. Surprisingly, only 8% of neurons showing touch responses under anesthesia showed touch responses during behavior, with 6.5% expected by chance (**Figure 8G, H**). Therefore different neurons encode passive whisker deflection and touch during active behavior. Furthermore, in contrast to the topography observed in anesthetized rats, we did not detect topography in directional tuning for neurons in anesthetized mice (**Figure 8I**).

The neuropil signal was still direction selective under anesthesia (**Figure 8J**). Moreover, the topography of neuropil directionality under anesthesia was consistent with that observed in anesthetized rats (Kremer et al., 2011): forward deflection elicited responses in and near the column of the whisker immediately anterior to the one deflected (**Figure 8K**). Thus, both individual neuron touch sensitivity and the overall organization of the directional map differed between actively sensing and anesthetized mice.

Discussion

Sensory cortex builds representations of the world to guide behavior. Even a single brain area such as vS1 contains approximately 400,000 neurons spanning three cubic millimeters of brain tissue (Hooks et al., 2011). These numbers present a sampling problem for comprehensive measurement of neural activity. To begin to overcome this challenge, we employed single whisker behaviors that depend on activity in the spared barrel column (Guo et al., 2014b; Hutson and Masterton, 1986; O'Connor et al., 2013; Shih et al., 2013). The superficial layers of a single barrel column contain approximately 2,000 neurons (Lefort et al., 2009), providing a tractable target for comprehensive imaging.

We sampled activity in 10-20,000 neurons per mouse, including the majority of L1-3 neurons in the principal and neighboring columns, while mice performed an active tactile task. The majority of neurons were active during behavior, with nearly half showing task-related dynamics. Here we only analyzed increases in fluorescence. Task-related decreases in fluorescence, consistent with reduced activity, were also observed, but were slow and weak, and not analyzed further. Our estimate of the fraction of silent neurons was in agreement with measurements based on unbiased electrophysiology (O'Connor et al., 2010b).

In the principal column, 12% of L2/3 neurons reported touch, 12 % reported whisker movements, and 5% reported both. Representations of whisker movements were distributed uniformly across multiple barrel columns, whereas representation of touch was concentrated in the spared column. We did not observe functional clustering on local scales: neighboring neurons were no more likely to be part of the same representation than expected by chance. Representations of different variables are therefore intermingled in L2/3.

We sampled about 1,200 neurons at 7 Hz. This rate is constrained by the need to sample serially, limited dynamic range of the protein calcium indicator, attenuation of light in tissue, and raster scanning, which makes non-optimal use of imaging time. Our finding of strong and correlated neuropil signals suggest that current methods for parallel sampling, such as light-field microscopy (Prevedel et al., 2014) and light-sheet microscopy (Holekamp et al., 2008) may be inadequate for isolating signals from single neurons in scattering tissue.

Extracting neural activity from population imaging data requires multiple steps. Images must first be registered to correct for motion (Dombeck et al., 2007; Greenberg and Kerr, 2009). Neurons or other structures must then be identified in the images. Finally, fluorescence dynamics must be converted to an approximation of neural activity (Dombeck et al., 2007; Greenberg et al., 2008; Vogelstein et al., 2010). Extensive automation is essential for image processing with the large datasets presented here (100,000 neurons or more). We made extensive use of cluster-computing and selected algorithms based on their robustness. Future progress will depend on the development of robust and standard image processing algorithms.

Relating neural activity to behavioral variables poses conceptual and computational challenges. Direct comparisons between sensory variables and calcium activity using methods such as correlation fail to incorporate the nonlinear aspects of neuronal tuning and the kinetics of calcium. Moreover, correlation-based methods are difficult to interpret when comparing coding of variables with distinct temporal structure. In our case, whisker angle is temporally dense, whereas touch is temporally sparse. Pearson's correlation provides a reasonable estimate of the linear relationship between activity and whisker angle, but will tend to overestimate the strength of the relationship between activity and object touch, and will underestimate the relationship strength with whisking if it is nonlinear. In contrast, encoding models that predict calcium activity explicitly enable principled comparisons between the strengths of coding for touch, whisker movements, and other behavioral variables (Ahrens et al., 2008; Huber et al., 2012; Miri et al., 2011). Applying a correlation-based approach to our dataset classifies a larger proportion of neurons as touch, likely explaining the discrepancy with other imaging studies in the barrel cortex (Chen et al., 2013a).

We employed a probabilistic encoding model to relate neural activity and behavior (Ahrens et al., 2008; Pillow et al., 2008) by explicitly modeling calcium kinetics and nonlinear receptive field shape. In contrast to other approaches, such as random forests (Huber et al., 2012), this approach is computationally efficient and yields transfer functions that are interpretable as receptive fields (Ahrens et al., 2008) (**Figure 5**). The computational efficiency of our approach allowed us to run numerous iterations of the model and perform rigorous statistical testing. In addition to more accurate classification, the model produced interpretable receptive fields for whisker curvature, allowing us to examine the topography of directional selectivity.

We related neural activity to the angle and curvature of the spared whisker. These parameters describe the motion of the whisker and the stresses at the follicle where mechanosensation takes place (Birdwell et al., 2007; Pammer et al., 2013; Quist and Hartmann, 2012). Though our videography ignored the three dimensional structure of the whisker, this should have little impact on neuronal classification: torsion about the whisker axis is tightly coupled to whisker angle (Knutsen et al., 2008). Our encoding model produced meaningful $\Delta\kappa$ kernels, but not whisker angle kernels (**Figure 5C**). Touch neurons with an angular preference should have produced a discernible peak in the whisker angle kernel. This implies that we did not detect tuning to position-at-touch. The encoding model captures nonlinear activity-variable relationships, as demonstrated by the curvature kernels (**Figure 5C**, **Figure S7**). It does not capture non-stationary activity relationships, such as adaptation during repeated touches. Moreover, our choice of behavior limits the stimulus space that was probed. For example, we did not explore vS1 responses to slip events typically encountered during exploration of textures (Chen et al., 2013a; Hires et al., 2013; Jadhav et al., 2009; von Heimendahl et al., 2007). It is possible that additional representations of tactile information will be discovered in other behavioral conditions and with the aid of more sensitive encoding models.

Topographic organization of direction selectivity has been reported in both cortex (Andermann and Moore, 2006; Kremer et al., 2011) and thalamus (Timofeeva et al., 2003) of anesthetized rats. We detected directional topography in actively sensing mice. In passively stimulated anesthetized mice we observed directional topography in the neuropil of anesthetized mice, but not at the level of neurons, suggesting that topography is present but very weak. Passive forward deflection of the whisker produces lateral forces opposite in sign to those experienced upon object contact during forward whisker movement (Birdwell et al., 2007; Hires et al., 2013). Thus, the anesthetized passive deflection tuning maps exhibit opposite organization in terms of lateral force from the map observed in actively sensing mice. Further investigation will be necessary to elucidate the origin of this discrepancy.

A major advantage of imaging is the ability to track activity over multiple days. We find the L2/3 population to be stable in terms of the fraction of neurons participating in the touch representation (**Figure 6**). Whisking neuron fraction and psychometric performance stabilized after the first few days of training, as did the distribution of touch and whisking encoding strength. This time course was more consistent with the time course of motor stereotypy emergence rather than performance improvement (**Figure 1E, G**). Because encoding classification as well as neurometric performance can change in response to the changes in motor strategy, it is possible that these early dynamics are merely a reflection of changing behavior and do not reflect genuine plasticity. Although our data are thus ambiguous regarding the nature of early learning, we do find stability during later learning. Interestingly, force tuning curves (and by extension, directional preference) among individual touch neurons as well as the fraction of touch neurons were stable from the outset of training. Such stability contrasts with motor cortex, where representations seem more dynamic (Huber et al., 2012; Masamizu et al., 2014; Peters et al., 2014). We did not observe the expansion of the representation of the spared whisker observed in some trimming studies (Margolis et al., 2012), nor did we observe the spared whisker contraction reported in animals exposed to enriched environments (Polley et al., 2004). The failure to observe map expansion may reflect lack of sensitivity in our imaging, or it could be a consequence of our measuring expansion in behaving rather than anesthetized animals.

Methods

Transgenic mice

The majority of mice in this study expressed mCherry in the nuclei of excitatory neurons. A custom reporter mouse (Rosa26 LSL-H2B-mCherry; **Figure S2**) was crossed with emx1-IRES-Cre (Jax 005628) (Gorski et al., 2002). The Rosa26 LSL-H2B-mCherry mouse was generated by targeted insertion of a construct containing the CAG promoter followed by a floxed-Stop cassette-controlled nuclear red fluorescent protein (Madisen et al., 2012) at the Rosa26 locus. The woodchuck hepatitis virus posttranscriptional regulatory element (WPRE) was used to enhance mRNA transcript stability. We chose mCherry as the fluorescent protein because its fluorescence emission spectrum does not overlap with GCaMP6 (Shaner et al., 2004). The nucleus was targeted by fusing mCherry to the histone protein H2B. Using the pair of PhiC31 recognition sites, AttB/AttP, the PGK-Neo marker can be deleted from the reporter lines in mice. Removal of the Stop cassette by Cre expression produces strong, red nuclear fluorescence in Cre-expressing cells.

We also generated a Gad2-T2A-NLS-mCherry mouse (**Figure S2**), which expresses mCherry in the nuclei of GABAergic interneurons. T2A-NLS-mCherry was inserted in-frame after the last codon of the GAD2 gene. Insertion was verified using a frt-Neo-frt cassette. This was removed by crossing the chimeras with R26FLP females (Jax 003946). Nuclear targeting was achieved using by fusing mCherry to a nuclear localization signal (NLS) sequence.

Surgeries

Mice (6-8 weeks old) were anesthetized (2% isoflurane, by volume) and infected with AAV2/1 syn-GCaMP6s (University of Pennsylvania viral vector core #AV-5-PV2824; **Table S1**). A craniotomy was made over vS1 and 9-12 viral injections (20 nL each) were delivered 450 μ m below the dura. Injections were laid out on a 300 μ m grid centered 3.7 mm lateral and 1.7 mm posterior to Bregma. Viral suspension was delivered at 15 nL/min using beveled glass pipettes, followed by a pause (1 minute) before withdrawal. A double-layer (4.5 mm external diameter, 3.5 mm inner diameter; No. 2 cover glass, Corning) window was placed over the craniotomy (Huber et al., 2012). The window and a titanium headpost were affixed to the skull with dental acrylic (OrthoJet). For L4 axonal imaging, a 2-by-2 400 μ m spaced grid was employed, with two 50 nL injections of AAV2/1 FLEX-CAG-GCaMP6s per site (400 and 700 μ m below dura). All animal procedures were in compliance with protocols approved by the Janelia Farm Research Campus Institutional Animal Care and Use Committee.

Behavioral training

The behavioral protocol, apparatus, and training procedures have been described (Guo et al., 2014b). Briefly, animals were placed on water restriction (1 ml daily) 3-5 days following viral injection. Training started one week later, in some cases under the

microscope (**Figure 6; Table S1**). On the first training day, head-fixed animals were trained to lick for water, after which they were introduced to the temporal structure of the task. On subsequent days, animals were first trained with two fixed pole positions and later with a range of positions for right lick trials (**Figure 1A**). The delay epoch was increased gradually to one second.

Whiskers were trimmed to the base with scissors every 2-3 days. On the day of surgery, animals were trimmed to 2 rows (C, D). One week later, intrinsic optical imaging was used to map somatotopy in vS1. Images of vasculature referenced to injection site images taken during surgery were used to determine the whisker row least suited for imaging, which was trimmed. Training started with one row. Once criteria performance was reached with a single row, animals were trimmed to one whisker. The chosen whisker corresponded to a barrel column with good viral expression and minimal surface vasculature occluding imaging (**Table S1**). After trimming all whiskers, performance fell to chance level (d-prime: -0.03 ± 0.11 , mean \pm S.D., n=4 mice), confirming that mice used tactile cues to solve the task.

The task was implemented using BControl (brodylab.princeton.edu/bcontrol). A behavioral trial began with a pre-trial period during which the pole (0.3 mm diameter; Wiretrol) was moved into position along the anterior-posterior axis using linear actuators (Zaber). The pole ascended into the whisker field (Festo), and remained in place for the 1 s sample epoch. The pole was withdrawn and, 1 s later, a linear actuator (Zaber) moved the two lick ports within reach of the tongue. A 50 ms auditory reward cue signaled the animal to respond. On correct trials, reward was delivered. Following reward collection, or after an incorrect lick, the lick ports were moved to an out-of-reach position. During periods where the lick ports were out-of-reach, licking was monitored using a laser (Thorlabs) and photodiode.

Whisker video was collected at 500 Hz using a high speed imaging system (Norpix) and high-intensity infrared illumination (LED 940 nm; Roithner Lasertechnik). Identification of whisker segments within frames was performed using the Janelia Whisker Tracker (openwiki.janelia.org/wiki/display/MyersLab/Whisker+Tracking) (Clack et al., 2012).

The identity of whiskers across frames was tracked using a custom, automated MATLAB package. Whisker angle was measured by fitting a polynomial to the traced whisker and determining the angle at the point the polynomial intersected a parabola that was a fixed distance away from the face. By convention, an angle of 0° was perpendicular to the animal's midline (leftward in **Figure 1B**), with positive angles corresponding to protractions (**Figure 1B**). Whisker curvature was measured a fixed distance range along the fitted polynomial (Pammer et al., 2013). Whisker touches were first detected using an automated algorithm, then verified manually in a custom MATLAB interface.

Whisker angle stereotypy (**Figure 1F**) was measured by taking the mean of whisker angle vectors across the trials of a given type (left, right; correct, error) and, for each trial, measuring its whisker angle vector's correlation with the mean vector (R^2). The mean of these, weighed by number of trials per trial type, was used as a single day's stereotypy measure. The start of whisking bouts (**Figure 5F, 6A**) was defined as the moment when

the amplitude of whisking, obtained using the Hilbert transform (as described previously; see (Huber et al., 2012)), exceeded 10 degrees.

For passive whisker stimulation (**Figure 8**) mice (n=3, emx1-Cre X LSL-H2B-mCherry) were lightly anesthetized using isoflurane (0.5%) and head-fixed under the microscope on a heating pad. Stimulation was provided using a “Y” shaped pole (0.3 mm diameter; Wiretrol) connected to a piezo. The whisker sat inside the Y, allowing for stimulation in both directions. Stimulation consisted of 5-10° ramp-and-hold deflections applied at 1,000-2,000°/s, 0.5-1 mm from the follicle base. Deflection parameters were verified using high-speed (500 Hz) videography. The stimulator was positioned so that the whisker was adjacent to but not touching the pole. Forward and rearward trials were interleaved every 10 or 20 trials.

Two photon imaging

Calcium imaging was performed using a custom two-photon microscope (openwiki.janelia.org/wiki/display/shareddesigns/MIMMS). Images were acquired using a 16X 0.8 NA objective (Nikon) and GaAsP PMTs (Hamamatsu). Green (GCaMP; BG22, Chroma) and red (mCherry; 675/70 emission filter, Chroma) fluorescence channels were collected simultaneously. Horizontal scanning was accomplished using a resonant galvanometer (Thorlabs; 16 kHz line rate, bidirectional). Axial motion was controlled by a piezo collar (Physik Instrumente). Powers at the sample ranged from 30 mW in L1 to 300 mW in L5. Exponential power modulation ($\lambda = 250 \mu\text{m}$) was employed during piezo scanning. Four 600 by 600 μm (512 by 512 pixels) imaging planes were acquired at 7 Hz each across depth. The system was controlled using ScanImage (scanimage.org; (Pologruto et al., 2003)).

The three imaging planes were either 15 μm (volumetric and L4 axon data) or 120 μm (imaging during learning and L5-L3 simultaneous imaging data) apart: the fourth fly-back frame was discarded. Each set of three planes constituted a subvolume (**Figure 2A**). Learning data tracked a single subvolume over multiple sessions. For volume imaging, each subvolume was imaged for 4.0 sessions (mean; range, 1-8), with data pooled across sessions. Eight subvolumes comprised a single volume, and two volumes were imaged per animal. These spanned 360 μm in depth (L1-L3). For L4 axon imaging, six subvolumes in a single volume over the spared whisker barrel were imaged (L2-L3; spanning 270 μm in depth). For simultaneous imaging of L5 somata and deep L3 apical dendrites the top and bottom planes of the subvolume were employed, for an inter-plane distance of approximately 240 μm .

After the first imaging session average images of each imaged plane were generated. At the beginning of all subsequent imaging sessions, these four images (three imaged planes, one fly-back plane) were employed as references while imaging. Positional adjustments were made to ensure that the same neurons were imaged over the course of the session (Huber et al.). On the middle imaging day, a high-resolution stack (1 μm spacing) was acquired, and all imaging planes were aligned to this stack to establish the relative positions of neurons. In emx1-Cre X LSL-H2B-mCherry animals, the locations of

individual neuronal nuclei were determined using the red channel of this reference stack (**Figure S2**) and a three-dimensional Gaussian mixture model. Briefly, the model looked for spheres whose diameter was 10 μm in a luminance-normalized stack. The expected fraction of image pixels belonging to nuclei was adjusted manually for each stack until the number of detected nuclei was reasonable, as judged by visual inspection.

Laminar boundaries were defined as follows. The L1-L2 border was the depth at which pyramidal neurons first appeared. The L3-L4 border was defined as the point where a drop off in labeled somata occurred. The L4-L5A border was defined as the point at which labeled somata appeared. These boundaries were linearly adjusted to fit into the relative positioning defined by (Hooks et al., 2011); the L2-L3 border was then extrapolated to occur at a defined relative position (13% of cortical depth; (Hooks et al., 2011)).

Calcium imaging analysis

Imaging data were processed in five steps: intra-session registration, inter-session registration, ROI selection, $\Delta\text{F/F}$ calculation, and Ca^{2+} event extraction.

1.) Images were motion-corrected using an automated pipeline (Huber et al., 2012). First, registration was performed on individual behavioral trials: rigid registration was performed using a down sampled fast Fourier transform (dFFT; reference image, 5 consecutive frames of trial with minimal luminance change), followed by registration using a custom line-by-line algorithm (similar to (Greenberg and Kerr, 2009)). Second, registration was performed across trials within a session. The mean of a luminance-stable trial toward the middle of the session was employed as the reference image. All other trials were aligned to this trial using the dFFT, sometimes followed by a non-rigid interpolated warp field transform.

2.) Images were aligned across imaging sessions using the same interpolated warp field algorithm employed in the last step of inter-trial registration (Huber et al., 2012). The reference image for each session of a given imaging plane was registered to the reference images of other sessions. For all session pairs of a particular imaging plane, normalized cross correlations were computed among the warp field-registered reference images. The session with the highest median correlation to all other sessions was selected as the master reference image for that plane (typically, one of the middle imaging sessions). Manual inspection was used to exclude sessions whose reference image differed excessively from the master reference.

3.) ROIs were drawn using the master reference image from step 2, then propagated to other imaging sessions using the warp-field transform. ROI drawing employed a custom user interface (MATLAB). A point near the cell center was selected. The algorithm constructed a matrix of intensities from the red channel (mCherry) where each column spanned a range of distances from the point and each row spanned all angles around the point. The sharpest intensity ridge along minimally varying distance was found, and this was treated as the border of the nucleus. The process was repeated, this time in the green

channel (GCaMP) and starting with the nuclear border. This next ridge in angle-distance space was considered the outer border of the cytoplasm (Chen et al., 2013b). Pixels between these two ridges were assigned to the ROI. For L5 apical dendrite ROIs, the semi-automated algorithm looked for a single ridge in angle-distance space in the standard deviation image of the green channel (**Figure S4**). Pixels between the clicked location and the edge of the ridge were assigned to the ROI. For L4 axon imaging, ROIs consisted of a series of 12-by-12 μm squares on a 25 μm -spaced grid. For each ROI, a peri-somatic neuropil ROI was generated, consisting of an annulus 3 to 13 μm away from the outer edge of the ROI (**Figure S3**). Pixels with a correlation above 0.2 to an adjacent pixel as well as pixels belonging to non-neuropil ROIs were excluded from neuropil annuli.

4.) Raw fluorescence was extracted for each ROI, and fluorescence transients were neuropil-corrected ($F_{\text{ROI-corrected}} = F_{\text{ROI}} - \alpha F_{\text{neuropil}}$). If the corrected trace fell below a baseline ROI fluorescence, the value was set to baseline, so as to prevent over-correction. Neuropil correction used the local, peri-somatic neuropil signal, as using a global neuropil signal resulted in under-correction (**Figure S3**). The scaling factor, α , was set to 1, as manual inspection revealed that scaling factors below this produced under-correction. L4 axon imaging did not employ neuropil subtraction. F_0 was calculated using a three minute sliding window. For cells with a highly skewed raw fluorescence distribution (generally, active neurons), the 5th percentile of raw fluorescence within the window was used as F_0 ; for cells with a symmetric distribution (generally, inactive neurons), the median was used. For cells with intermediate fluorescence distribution skewness, an intermediate percentile was used. $\Delta F/F = (F - F_0)/F_0$ was then calculated for each ROI.

5.) Event detection was performed using greedy template fitting of the $\Delta F/F$ trace (**Figure 2G,H**), which is similar to peeling approaches described previously (Lutcke et al., 2013). The template bank consisted of sums of exponentials with rise times of 3 to 5 frames (426 to 714 ms; see (Chen et al., 2013b)) and decay times of 7 to 35 frames (1 to 5 s). Candidate events were selected based on large second derivatives, and convolved with the template bank. For each ROI, a noise estimate, σ , was obtained by subtracting a Savitzky-Golay fit of the trace and taking the standard deviation of the remaining trace. If any of the candidate events had a root mean square difference of less than σ from the actual trace but were, on average, larger than σ in amplitude, the event was accepted. That is, events had to be within the noise envelope of the actual trace, but had to have an amplitude exceeding noise. Accepted events were subtracted from the fluorescence trace, and the process repeated until no further events met the criteria.

We evaluated the performance of our event detector using simultaneous imaging and cell attached recording data previously obtained in L2/3 of mouse visual cortex (Chen et al., 2013b). The data was degraded to match the per-neuron dwell time and sampling rate of the volume imaging employed in this study. For a false positive detection rate of 0.01 Hz, $54 \pm 10\%$ of single action potentials (mean \pm S.D; $n = 7$ neurons) were detected

Following event detection, a de-noised $\Delta F/F$ trace was produced by convolving events with appropriate amplitude, rise, and decay time constants. This trace was employed for all subsequent analyses, unless otherwise noted.

Model-free analysis of task-related activity

Neurons were classified as task-related if their activity was distributed in a non-random manner relative to the structure of the trial. For each cell, we counted the fraction of trials during which a particular time bin had calcium events (**Figure 4A, B**). A null distribution was constructed by shuffling the calcium events in time for each ROI 10,000 times and performing the same analysis. For each shuffled iteration, the peak was determined (in units of fraction of trials with events). If the actual peak exceeded the 95th percentile of the shuffled peaks, the cell was scored as task-related. This analysis was done separately for correct left and right trials; neurons satisfying the criteria on at least one of these two trial types were classified as task-related.

Neurometric performance was measured for each neuron using ROC analysis. Individual trial responses were reduced to a decision variable based on the peri-stimulus time histogram (PSTH), as described previously (O'Connor et al., 2010b). Briefly, for all trials where the appropriate response was left or right (**Figure 1A**), a PSTH was computed using the event trace (event amplitude corresponded to $\Delta F/F$ amplitude, but no exponential decay was included). The dot product similarity of a trial's PSTH to the mean PSTH across respond-left trials minus the trial's dot product similarity to the respond-right PSTH was used as the decision variable. ROC analysis was performed using this decision variable, allowing us to determine how well an ideal observer could discriminate left from right trials. Both correct and incorrect trials were included. Trials in which the animal did not respond were excluded from the analysis. ROC analysis was performed on the real event trace as well as 100 traces with shuffled trial labels. Neurons with area under the curve (AUC) values exceeding the 95th percentile of shuffled AUCs were considered discriminative. Data were pooled across all volumetric sessions.

Population neurometric performance (**Figure 6D**) was assessed using a maximum likelihood decoder (naive Bayesian classifier), in which responses from different conditions were modeled as multivariate Gaussian distributions assuming zero covariance. The decoder was tasked with differentiating trials where the correct response was a left or a right lick; both hit and error trials were included. Five-fold cross-validation was employed, with each training set using 80% of trials, and testing on the remaining 20%. Only neurons that were individually discriminative based on ROC analysis were included. One hundred groups of 10 randomly selected discriminative neurons were tested for each day in each animal.

Model-based encoding analysis

Neuronal encoding of behavioral variables was assessed by fitting a statistical model to every neuron independently. The model was a cascaded generalized linear model that predicted neuronal activity as a function of the whisker angle and curvature under a

Gaussian noise model with input nonlinearities (Ahrens et al., 2008). Let s_1 be the time-varying whisker angle and let s_2 be the time-varying curvature. The model predicted the time-varying $\Delta F/F$, response r , as

$$r \sim Norm(z, \sigma^2)$$

$$z = f_1(s_1) * k_1 + f_2(s_2) * k_2$$

Where f_1 and f_2 are static, point-wise nonlinearities, and k_1 and k_2 are temporal kernels. k_1 and k_2 each consisted of 14 time points (2 seconds). f_1 and f_2 were each parameterized as a weighted sum of sixteen triangular “tent” basis functions

$$f = \sum_{i=1}^{16} w_i b_i(x)$$

Where x is the input (either s_1 or s_2), with each b_i given by

$$b_i = \begin{cases} (x - x_{i-1}) / (x_i - x_{i-1}), & i > 1, \quad x_{i-1} < x < x_i \\ (x_{i+1} - x) / (x_{i+1} - x_i), & i < N, \quad x_i \leq x < x_{i+1} \\ 0, & \text{otherwise} \end{cases}$$

Fitting of the model parameters k_1 , k_2 , f_1 , f_2 was performed using maximum likelihood with block coordinate descent. The model is bilinear in the parameters, and thus not globally convex, but it is convex when either the temporal kernels, or the nonlinearities, are held fixed (Ahrens et al., 2008). In these two cases, the remaining parameters can be estimated through convex optimization in closed form because of the Gaussian noise model (for other noise models it would require gradient descent). Thus, alternating coordinate descent on the full model can be performed (akin to alternating least squares; (Young et al., 1976)). Although not guaranteed to converge to a global minimum, this procedure was found to reliably estimate model parameters within approximately three to five iterations, achieving unique solutions both in simulation and in practice.

To remove degeneracy in the model associated with arbitrary scaling factors on either the kernels or the nonlinearities, the nonlinearities were forced to have minimum of 0 and maximum of 1. This constraint was applied after each iteration. No constraints were placed on the temporal kernels. In interpreting the results, however, the shape of the recovered nonlinearity is uninformative if the amplitude of the corresponding temporal kernel is near 0.

Whisker movements were sampled at 500 Hz while calcium responses were measured at 7 Hz. Given the nonlinearity in the model, additional information in the higher resolution whisker variables could be incorporated into the prediction. Specifically, the nonlinearity was applied to the whisker variables at their native temporal resolution, followed by linear down sampling to 7 Hz.

To prevent over-fitting, a prior was used to ensure smoothness of both the temporal kernels and the nonlinearities. Formally, this was implemented through a Gaussian prior on each set of parameters, with an inverse covariance given by the second derivative matrix (a matrix with values 2 and -1 for the main and off diagonals, respectively). Employing such a prior corresponds to maximizing the log-posterior, with the prior adding a small penalty to the objective function. In order to fit several thousand cells efficiently, the scale factor associated with this penalty was determined from a cross-validated inspection of a random subset of ROIs. On subsets of data, changing this parameter did not qualitatively change the fractions of neurons identified as touch or whisking.

Model fitting was performed using five-fold cross-validation across trials (80% used for fitting, 20% used for model evaluation, with 5 distinct groupings per fit). Quality of fit was assessed using the Pearson correlation between the actual and model-predicted $\Delta F/F$ traces (R_{fit} ; calculated on data not used for fitting). To assess the statistical significance of either the whisking or touch component of the response, a permutation test was employed. Specifically, the model was fit using only one of the two variables. In each case, the significance of that variable's contribution was assessed by repeating the fit using 100 time-shuffled $\Delta F/F$ traces. These were generated by taking the individual calcium events and distributing them randomly, while maintaining the rise and decay time constants associated with each event. If the actual model fit produced an R_{fit} in excess of the 95th percentile of shuffled R_{fit} values, the neuron was said to belong to that category. If both predictions were significant, the neuron was considered as belonging to both categories. The 95th percentile criteria was employed throughout, unless otherwise noted.

For imaging during learning, the encoding model was fit to data from individual sessions using a more stringent 99th percentile criteria. On days where multiple whiskers were present, only the eventual spared whisker's curvature and angle information were employed for encoding model fitting. For trend analysis during learning, two sessions per animal were employed to increase statistical power (**Figure 6C-E**).

For volumetric imaging and L4 axon imaging, the model was fit to data pooled across all sessions for which a given ROI was imaged. Volume maps (**Figure 7**, **Figure S5**) were constructed by pooling data across all relevant animals and collapsing from four quadrants into one in arc-row space. Maps were constructed with a more stringent inclusion criterion: only cells with $R_{fit} > 99^{\text{th}}$ percentile of event-shuffled R_{fit} s were included. Voxels for arc-depth and row-depth were 15 μm by 15 μm by a half barrel width in size, and the fraction was estimated for a voxel using pooled data. Arc-row voxels were 15 by 15 μm , spanning all depths, but restricted to specified cell types.

The encoding model-derived static nonlinearity for curvature in touch neurons was equivalent to a receptive field, with whisker curvature-dependent amplitude ranging from 0 to 1 (**Figure 5C, D; 8B**). Because curvature is proportional to the force experienced by the whisker follicle (Birdwell et al., 2007; Pammer et al., 2013), and because most whisker motion is along a single plane, the sign of the curvature change provided

directional tuning. Negative curvature changes correspond to pole contacts during protraction, whereas positive curvature changes correspond to contacts during retraction. We derived a directionality index, DI, that was negative for neurons preferring retraction touch and positive for neurons preferring protraction touch (**Figure 8B**, inset). The index, DI, was simply the ratio of the difference between the amplitude of the force kernel at the maximal protraction and retraction $\Delta\kappa$ value divided by the sum.

Directionality maps were constructed for cells with R_{fit} values above the 99th percentile of event-shuffled R_{fit} s (**Figure 8D**). The map consisted of 50 μm by 50 μm voxels spanning depth (300 μm). The stability of directional preference was assessed using the average pairwise root mean square (RMS) of the curvature static nonlinearities for touch neurons (**Figure S7**). To restrict the dataset to neurons responding mainly touch-induced curvature changes the 99th percentile criteria was employed and the touch R_{fit} had to exceed 0.25.

In anesthetized mice direction tuning was measured for ROIs showing a significant response to piezo deflection, defined as a trial-locked mean response whose amplitude exceeded the 95th percentile of an event-shuffled null distribution. ROIs were further required to have event rates above 0.01 Hz. The directionality index for ROIs meeting this criteria was defined as the difference between the peak $\Delta F/F$ response to rearward (R) and forward (F) deflection divided by their sum: $DI_{anesthetized} = (R-F)/(R+F)$.

Figure Legends

Figure 1. Whisker-based object localization behavior.

- A.** Mice were trained to lick the right (blue) lickport if the pole appears in the blue range of positions, or the left (red) lickport if it appears at the red position. Light blue fan: range of whisking.
- B.** Single frame from whisker video (500 Hz) with whisker position (θ) and curvature (κ).
- C.** Behavioral data for a series of trials. Individual trials consist of a sample epoch, during which the pole is within reach of the whisker (1-2 s); a delay epoch (1 s), during which mice have to withhold licking; and a response epoch, triggered by an auditory “reward cue”, when mice should signal their behavioral choice. Green, whisker angle (θ , left panel); blue, curvature change ($\Delta\kappa$, right panel); gray ticks, touches
- D.** Experimental timeline.
- E.** Performance (d-prime) as a function of training day for individual animals. Dotted line, criterion, d-prime > 1.5. White circle, no imaging; grey circle, imaging during learning; red circle, volume imaging.
- F.** Example whisker position trajectory for 10 randomly selected trials on first and final training days. Gray, sample epoch; magenta, reward cue.
- G.** Whisker angle stereotypy, as quantified by trial-to-mean correlation (see Methods) for the three mice with the longest period of videography during training. Black, mean; grey, imaging during learning; red, volume imaging.

Figure 2. Volume imaging in behaving animals.

- A.** The basic experiment. Subvolumes consist of three imaging planes (e.g., cyan lines) imaged simultaneously (inter-plane distance: 15 μm). Eight subvolumes (red shades, cyan) comprise a volume. Two volumes (gray boxes) were imaged per mouse. Inset, tangential view of the brain (gray, volume boundaries; red dot, center of the principal column). Green fluorescence indicates GCaMP6s expression. Right, three example imaging planes (green, GCaMP6s fluorescence; red, mCherry fluorescence).
- B.** Laminar distribution of GCaMP6s expression (green). Red, mCherry fluorescence (emx1-cre x Rosa26-LSL-CAG-H2B-mCherry mouse).
- C.** Closer view of plane 1 in **A** (dashed box). Gray arrows, putative GABAergic neurons.
- D.** Somatic region of interest (ROI) masks from plane 2 in **A**.
- E.** Average percentage of excitatory L2/3 neurons imaged ($n=5$ volumes).
- F.** Neuropil ROI generation. Left to right: raw image (green, GCaMP6s; red, mCherry), neuronal ROI, neuropil ROI (**Figure S3**).
- G.** Raw $\Delta F/F$ traces of neurons colored in **D**. Blue and red vertical regions indicate sample epochs for right and left trials, respectively. Gray, touches.
- H.** Event-based $\Delta F/F$ traces corresponding to **G**.

Figure 3. Neuronal activity and coding in the barrel cortex.

- A.** Fraction of active ROIs across layers and cell types (event rate > 0.0083 Hz).
- B.** Task-related activity fraction by population (**Figure 4**).
- C.** Object location discrimination fraction by population.

D. Encoding class fraction by population (**Figure 5**). Mixed neurons represented both touch and whisking.

Figure 4. Temporal distribution of neural activity.

- A.** Measurement of behavior-related activity. Neural activity for right and left trials (only correct trials are shown). Data are trial-aligned. White lines, sample epoch; magenta line, reward cue; red ticks, individual calcium events.
- B.** The fraction of trials on which a given trial-aligned time point has an event. Dotted horizontal line, 95% confidence interval of activity peaks based on shuffled data. This neuron shows trial-related activity for right, but not left, trials.
- C.** Activity distribution across all ROIs having significant task-related activity. Each line corresponds to a single ROI. The responses are averaged across all trials and z-scored. Neurons are sorted by activity peak time (pooled across animals: n=3 for L4, n=8 otherwise). Only ROIs from the principal column were employed, unless noted.
- D.** Temporal distribution of activity peaks for different neuronal types (red), aligned to trial time. Green, whisking amplitude; blue, touch fraction (n=8 mice).

Figure 5. Encoding model and neuronal classification.

- A.** The encoding model predicts neuronal $\Delta F/F$ (top-right, blue trace) from whisker dynamics (top-left; $\Delta \kappa$). The model consists of a nonlinearity, or receptive field (bottom-left), acting on the behavioral variable domain. This is multiplied with a temporal kernel (bottom-right), acting on the time domain. The predicted $\Delta F/F$ trace (blue) is compared to actual $\Delta F/F$ (black) using Pearson's correlation to compute R_{fit} (top-right).
- B.** Example behavioral variables used to fit the model for one subvolume. Curvature change ($\Delta \kappa$; right) was zero for non-touch periods. Vertical white dashed lines, sample epoch; magenta lines, reward cue. Blue vertical lines, right trials; red, left trials; stippled red and blue, error trials.
- C.** Fit for an example cell (corresponding to behavioral data shown in **B**). Top, temporal kernels; bottom, receptive field nonlinearities (green, whisker angle; blue, curvature change; mean \pm bootstrap 2.5 and 97.5 percentiles). Though the model produced a receptive field for θ , the temporal kernel was not significantly distinct from 0, implying that it is not informative. This is illustrated by scaling the amplitude of each nonlinearity by the norm of the corresponding temporal kernel.
- D.** Same as **C**, but for a whisking neuron.
- E.** Raw $\Delta F/F$ (left) and model-predicted $\Delta F/F$ (right) for the two cells in **C, D**.
- F.** Example neural responses aligned to behavior. Top traces, whisker curvature aligned to first touch (blue, left) and whisker angle aligned to first whisking bout (green, right; see Methods). Remaining traces, $\Delta F/F$ aligned to the first touch (blue) or first whisking bout (green) of a trial for example touch (left) and whisking (right) neurons. Light color, individual trials; dark color, mean.

Figure 6. Dynamics during learning.

- A.** Example neurons imaged during learning of the object localization task (before volume imaging). Left, touch cell; right, whisking cell.

- B.** Fraction of L2/3 excitatory neurons classified as touch or whisking during learning. Mean touch, blue; mean whisking, green; grey lines, individual animals (n=4).
- C.** Ratio of the fraction of neurons representing touch inside the principal column to fraction representing touch outside the principal column with respect to day following single whisker trimming (day 0: day of trimming). Mean is indicated in blue; grey lines, individual animals (n=3).
- D.** Neurometric and psychometric performance over the course of learning. Orange line, task performance of the best 10 neuron ensemble; grey lines, individual animals' (n=4) best ensemble performance; black, cross-animal psychometric performance (the first day of training consisted of a simplified form of the task where the performance metric did not apply and was thus excluded).
- E.** R_{fit} of top 50 touch (blue) and whisking (green) neurons over the course of learning, across animals (n=4). Grey, days where the neuron did not meet the $p < 0.01$ criteria to be counted as a neuron of that category (see Methods); white, missing data. Trace length identifies source animal (training length was unique).
- F.** Example relationship between the R_{fit} for touch neurons across days. Top panel: the relationship between the first and 5th day; bottom panel: 9th and 13th days.
- G.** Example animal's correlation of touch R_{fit} vectors from one day with another (i.e., the R values from F for all day-pairs). The diagonal elements show the median correlation of 25 pairs of intra-day R_{fit} vectors (see Results). Red dots, days for which the inter-day correlation fell below the 5th percentile of the intra-day correlation values for the first day of that row.
- H.** Fraction of days after the specified day for which the inter-day R_{fit} correlation fell below the 5th percentile of intra-day R_{fit} correlations (red dots in G). Blue, touch; green, whisking; grey, individual animals (n=4).

Figure 7. Spatial distribution of representations in L2/3.

- A.** Example imaging plane with somatic ROIs. Blue, touch neurons; green, whisking neurons; cyan, mixed; grey, unclassified. Intensity is proportional to R_{fit} . Thick dashed line, outline of the principal column; thin lines, surround columns.
- B.** Three dimensional distribution of response types in one mouse. Blue, touch neurons; green, whisking neurons; cyan, mixed; grey, unclassified; grey dashed line, outline of principal column. Radius indicates R_{fit} .
- C.** Map of touch and whisking excitatory neurons across L2/3 (pooled across 8 mice). Color represents the fraction of neurons in a given voxel classified as either touch or whisking. Location is in terms of distance from the center of the spared whisker column (dashed line). The maps on top represent the row-arc plane, averaging cell fractions through depth. The arc-depth (left) and row-depth (right) maps average cell fractions across a half-barrel width in the plane-orthogonal direction.

Figure 8. Directional selectivity maps in response to active and passive touch.

- A.** Two example direction selective touch neurons.
- B.** Distribution of directionality indices (DIs; inset) across touch neurons. Positive DI: protraction preferring; negative DI: retraction preferring ($\Delta\kappa < 0$ corresponds to protraction; see Methods).

C. Spatial distribution of directional selectivity in an example plane. Color indicates DI for touch neurons. Grey, unclassified. Thick dashed line, outline of the principal column; thin lines, surround columns.

D. Distribution of directional selectivity for excitatory L2/3 excitatory neurons across animals (n=7). Magenta, retraction preference ($DI < 0$); cyan, protraction preference ($DI > 0$). Here, it is assumed that the principal column was C2; in that case, the C1 and C3 columns would be where indicated (gray lines, barrel border). DI was averaged using 50 μm bins.

E. Directional selectivity along a whisker row for L2/3 excitatory neurons, L4 axons in L2/3, and L2/3 neuropil. DI was averaged using 45 μm bins along the row axis. Positive row axis values correspond to moving towards columns of whiskers anterior to the spared whisker (principal column: gray dotted line; columns labeled for case where PC was C2). Solid grey line, linear fit to points; Pearson correlation coefficient indicated along with p-value.

F. Directional preference map and its relationship to contact forces. In the active touch condition (top), protraction/retraction contacts elicit rearward/forward lateral forces along with a radial force (blue) at the follicle. Responding neurons (green) are predominantly confined to the segment of the barrel close to the whisker immediately anterior/posterior to the one contacted. Under anesthesia, forward/rearward passive deflection produces a spatial distribution of responding neurons similar to protraction/retraction contacts during active touch, but with opposite lateral forces.

G. Example responses among three neurons for the active, awake (top) and passive, anesthetized (bottom) conditions. Cyan, protraction touch; magenta, retraction touch; light cyan, forward anesthetized deflection; light magenta, rearward anesthetized deflection.

H. Fraction of neurons showing touch responses under active (blue) and passive conditions (light blue); dark blue, overlap (n=3 mice).

I. Directional selectivity within a whisker row for L2/3 excitatory neurons under anesthesia, as in **E**.

J. Fraction of neuropil ROIs showing touch responses, as in **H**.

K. Directional selectivity among neuropil ROIs in anesthetized mice, as in **E**.

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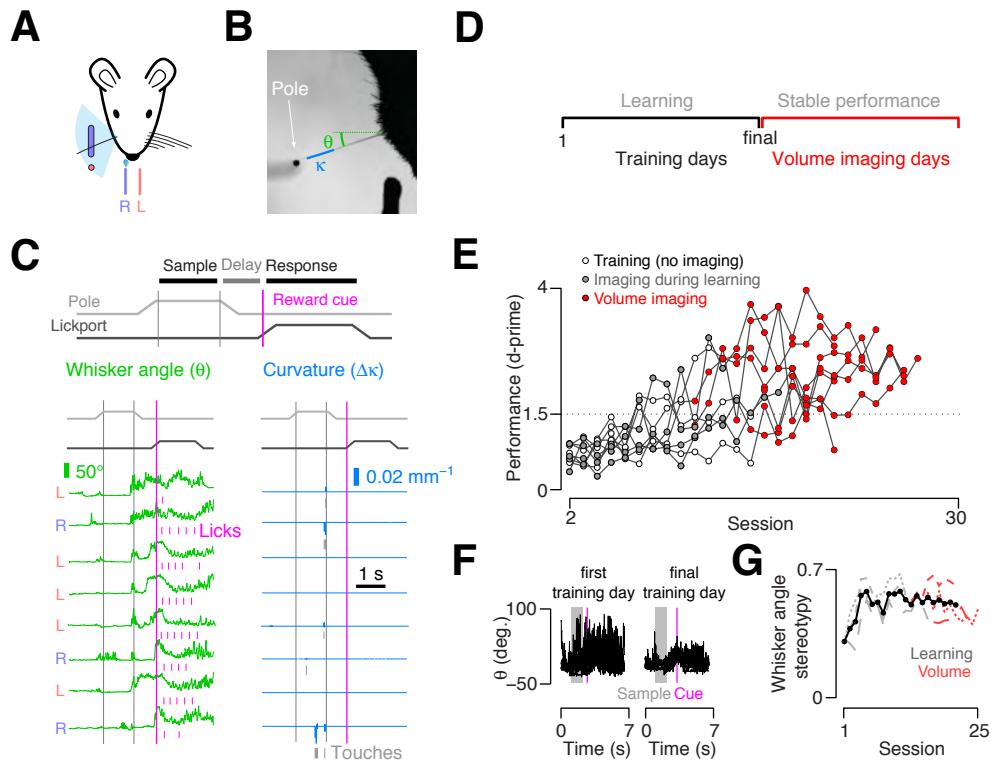
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Figure 1

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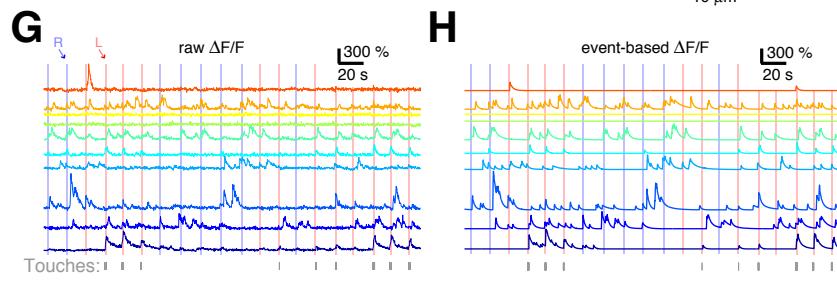
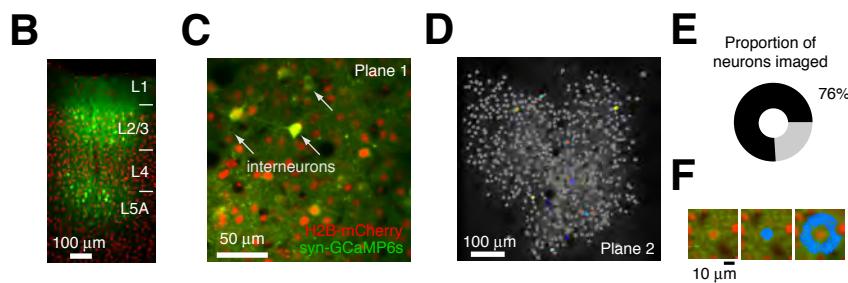
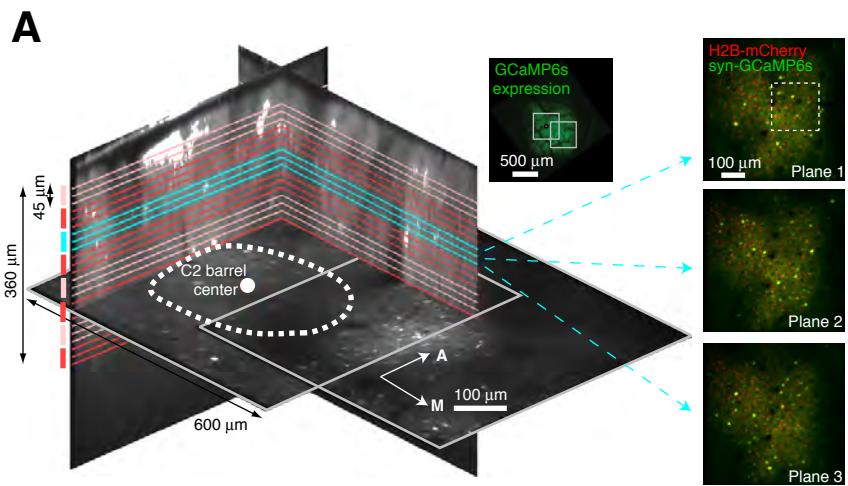


Figure 3

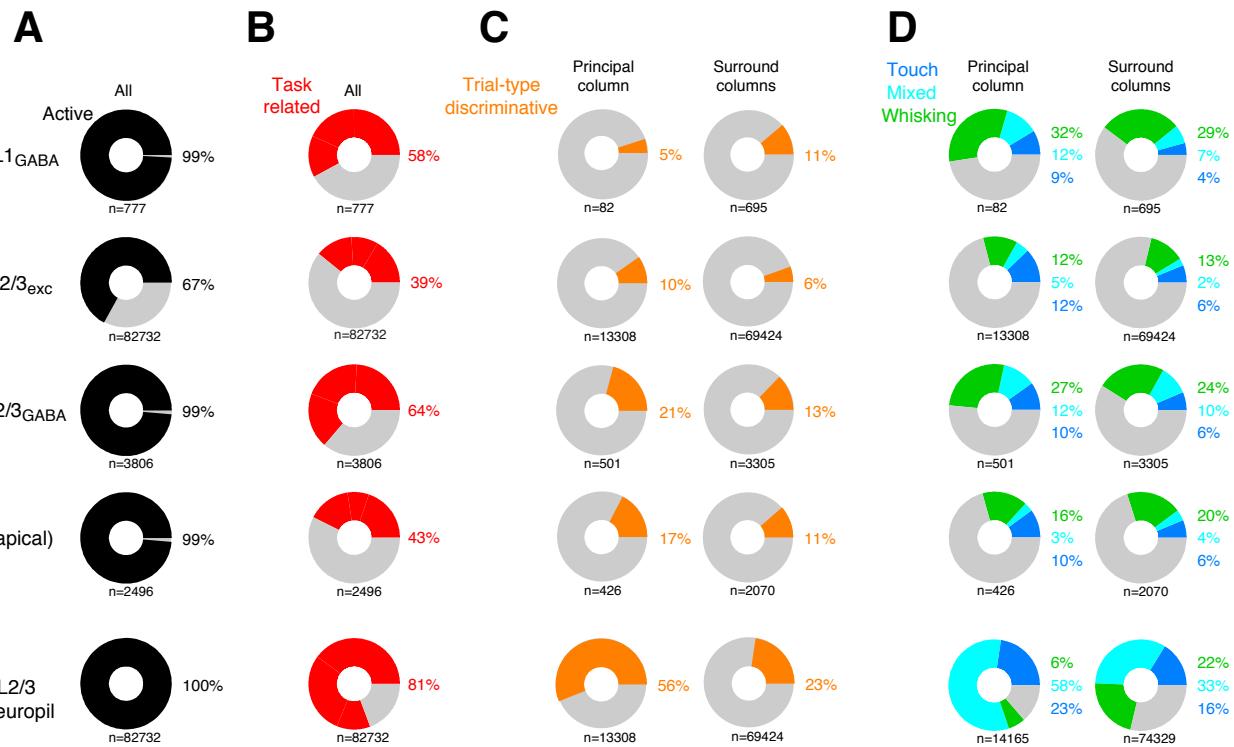
Figure 3, Peron et al.

Figure 4

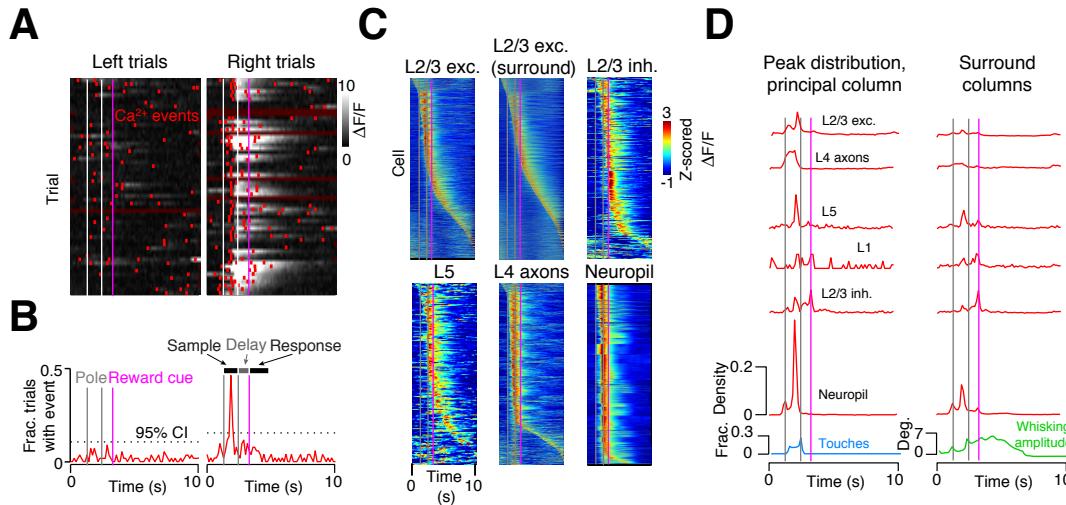
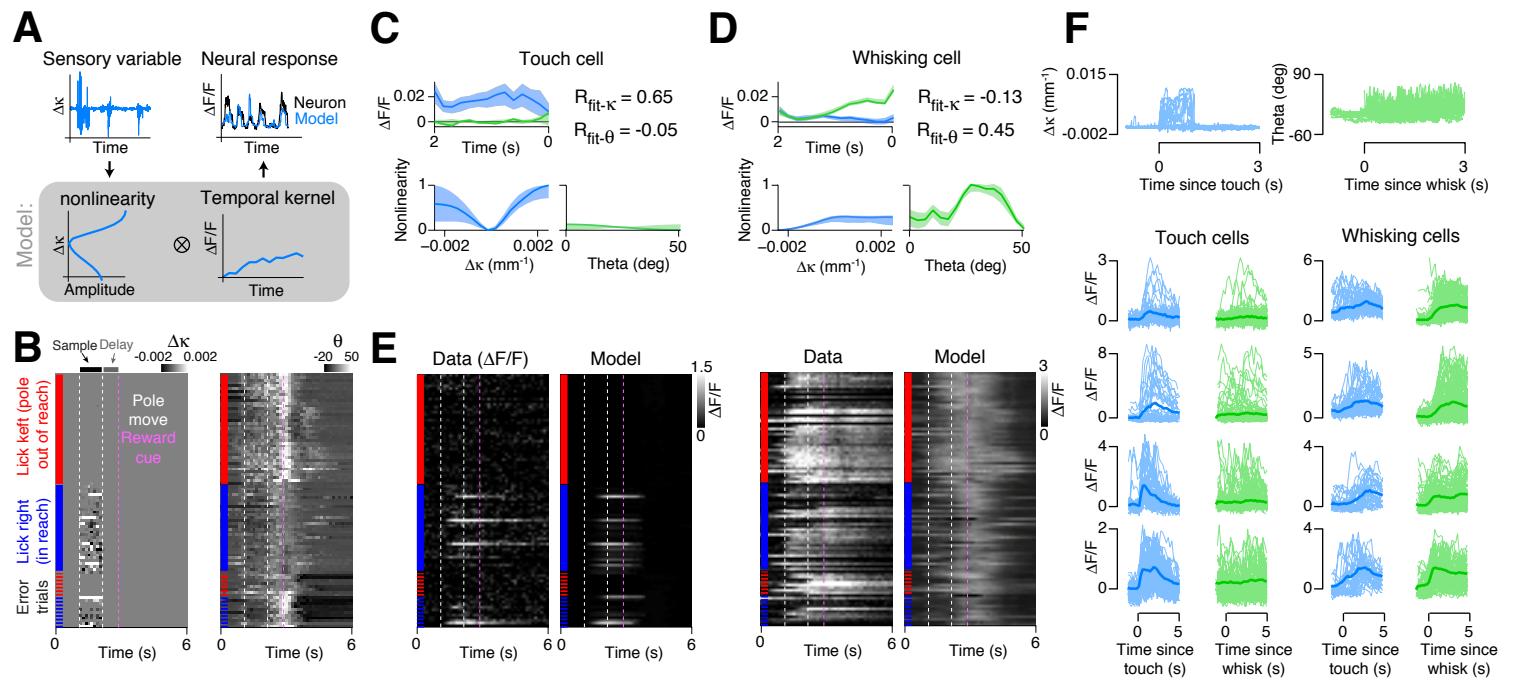
Figure 4, Peron et al.

Figure 5

Figure 5, Peron et al.



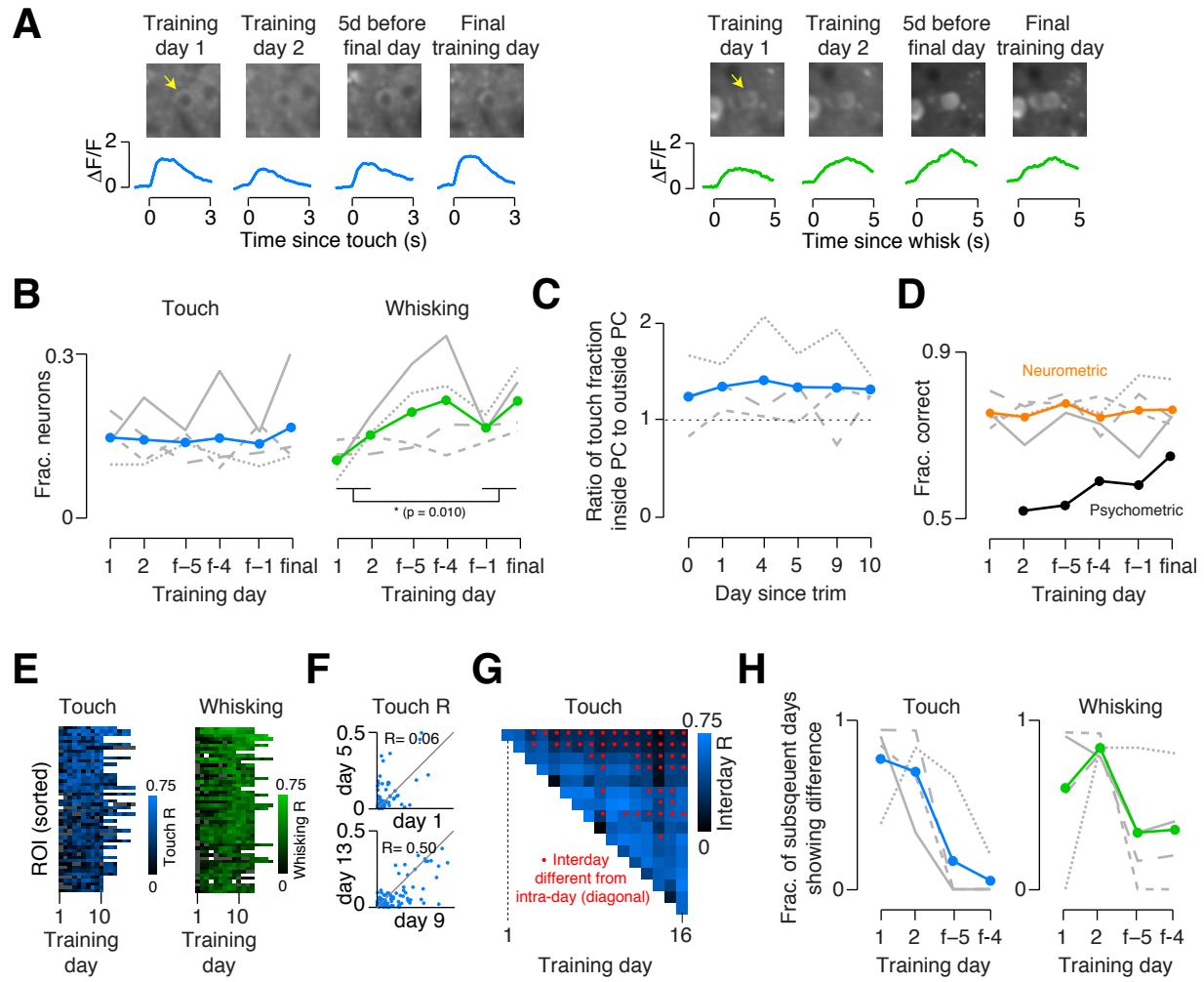


Figure 7

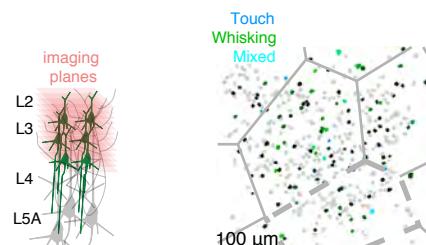
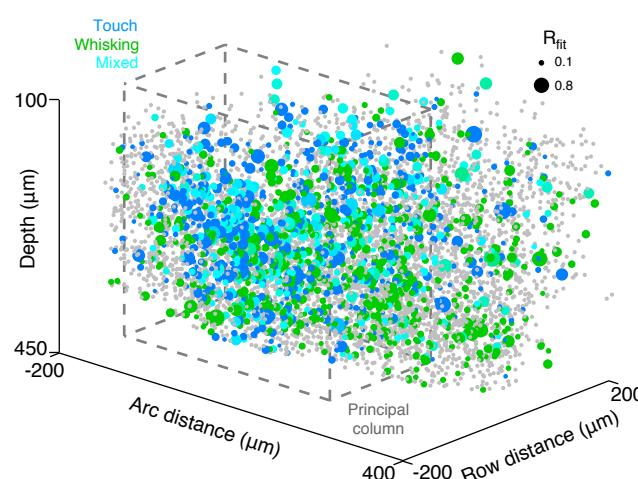
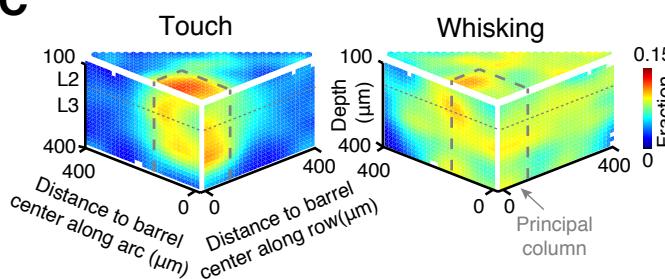
Figure 7, Peron et al.**A****B****C**

Figure 8

Figure 8, Peron et al.

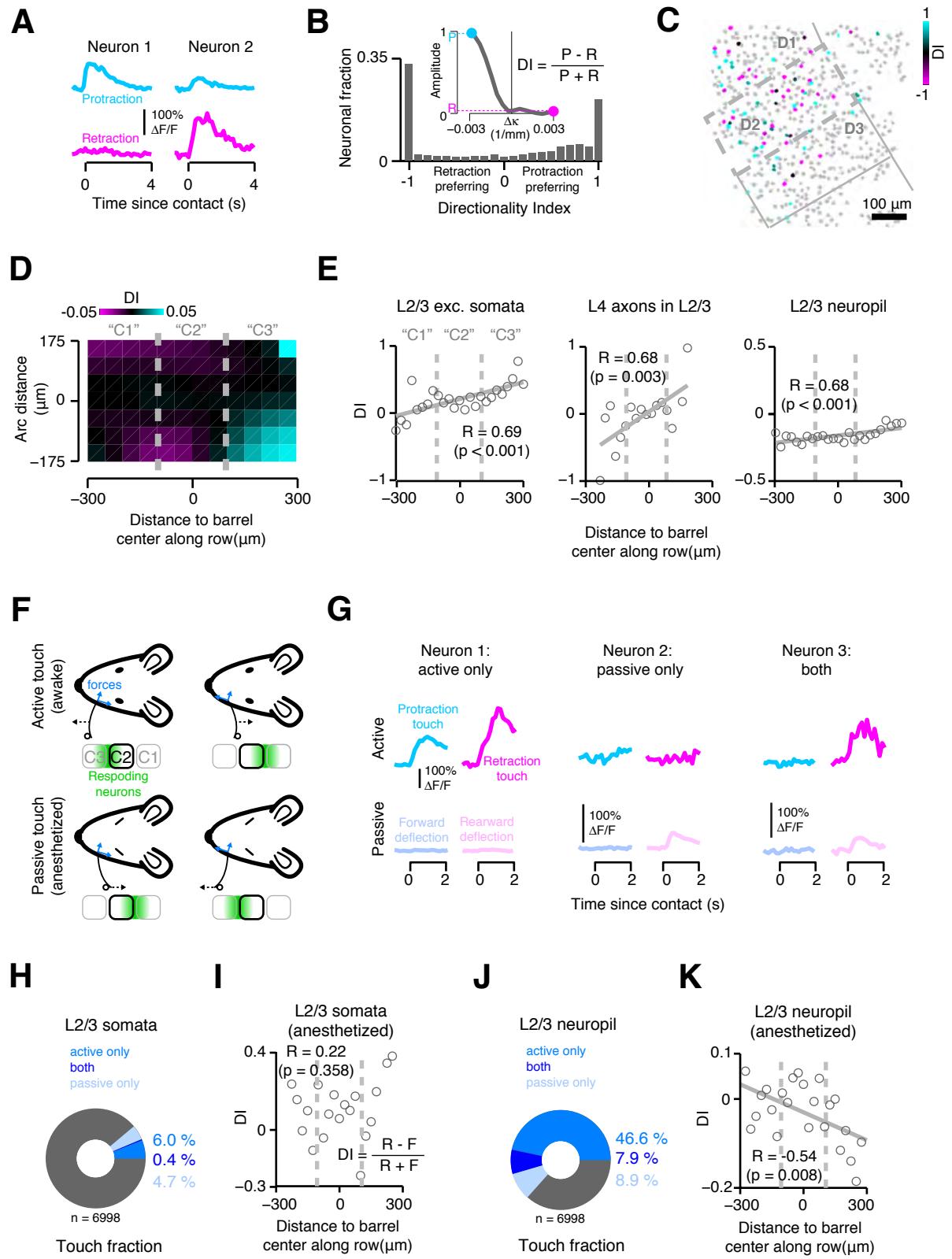


Figure S1. Identification of the principal barrel column.

Related to Figure 2A. Illustrates how the barrel column corresponding to the spared whisker was identified.

Figure S2. Red nuclear transgenics and segmentation.

Related to Figure 2C and D. Describes mice expressing the red nuclear marker. Also related to figure 2E. Illustrates how the fraction of L2/3 neurons imaged was estimated.

Figure S3. Neuropil correction.

Related to Figure 2F and 2G. Describes the neuropil subtraction.

Figure S4. Imaging Layer 5 apical dendrites.

Related to Figure 3.

Figure S5. Maps of touch and whisking for L4 axons and L2/3 neuropil.

Related to Figure 7. Describes spatial aspects of neuropil and L4 axon signals.

Figure S6. Spatial clustering among touch and whisking neurons.

Related to Figure 7. Additional analysis of spatial distribution of touch and whisking neurons.

Figure S7. Dynamics of single neuron directional preference during learning.

Related to Figures 8 and Figure 6. Shows stability of direction tuning during learning.

Table S1. Experimental animals.

Related to Figure 1. Lists animals that were used in various imaging protocols.

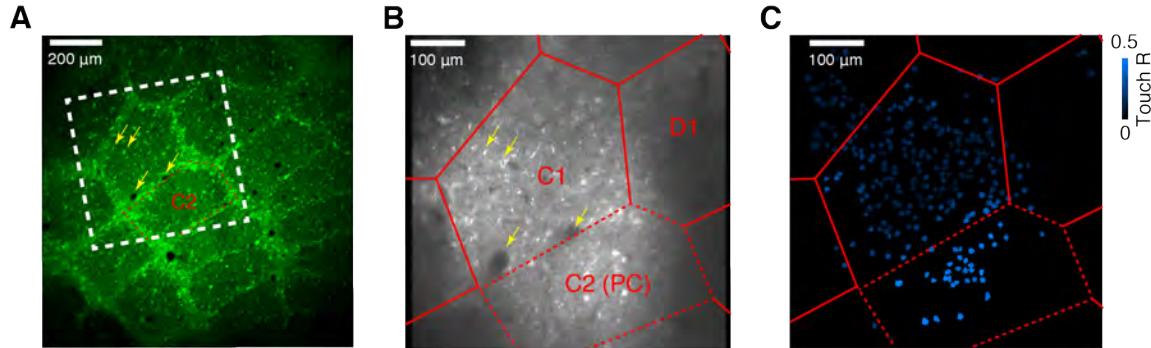


Figure S1. Identification of the principal barrel column.

A. Confocal image of a tangential section spanning the L4 barrel field in a GCaMP6s expressing animal. Red, principal column (C2).

B. Sample L4 plane from a two-photon image stack (white square in **A**). Red, barrel boundaries from **A**; dotted red, principal column. Confocal images were aligned with two-photon images using vasculature and dendrites (yellow arrows) in three mice. Because the barrel borders visible in **A** could also be identified in L4 two-photon images, the barrels in the remaining mice were identified using a combination of two-photon images and intrinsic signal imaging (which identified the barrel receiving input from the spared whisker). The neuropil touch signal (**C**) was used as confirmation.

C. Neuropil touch responses measured in a single whisker (C2) animal in a more superficial (L2/3) portion of the region shown in **B**. Strong neuropil touch response is confined to the principal column.

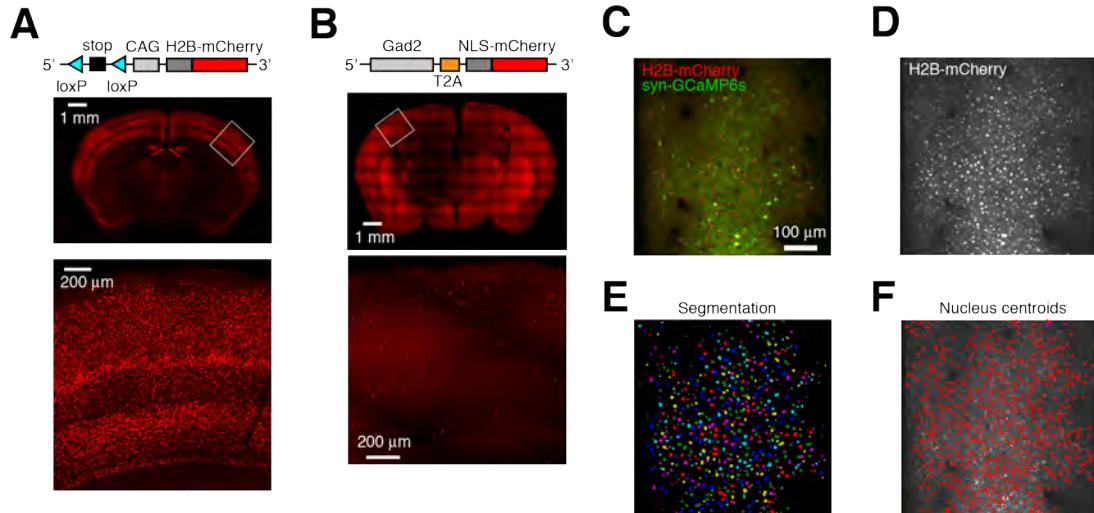


Figure S2. Transgenic mice expressing red fluorescent proteins and segmentation.

- A. Coronal section from a Rosa 26-LSL-H2B-mCherry X synapsin-Cre mouse. Inset, the LSL-H2B-mCherry gene inserted in the Rosa26 locus.
- B. Coronal section from a T2A-NLS-mCherry mouse. Inset, construct inserted at the Gad2 locus.
- C. Example plane from a 600-by-600-by-600 μm stack (depth resolution: 1 μm) in a Rosa26-LSL-H2B-mCherry X emx1-Cre mouse. Green, GCaMP6s fluorescence; red, mCherry fluorescence.
- D. Isolated mCherry fluorescence from C.
- E. Red nuclear segmentation of the plane using the Gaussian mixture model (Methods).
- F. Detected neuronal nuclei (red dots) overlaid on the mCherry image.

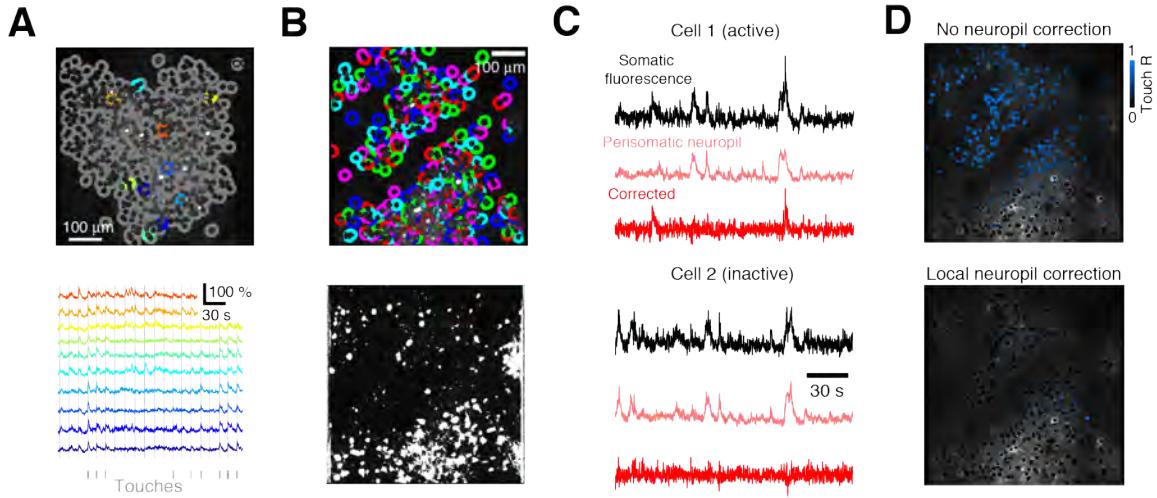


Figure S3. Neuropil correction.

- A.** Neuropil imaging. Top, image with neuropil masks. Bottom, fluorescence signals ($\Delta F/F$) in neuropil. Colors of the traces correspond to masks in image. The light blue and red vertical lines indicate the sample epoch for right and left trials; grey ticks, touches.
- B.** Example neuropil masks (top) and nearest-neighbor correlation mask (bottom). White pixels whose peak correlation with one of eight neighbors exceeds 0.2. These pixels, and neuronal ROI pixels, are excluded from the neuropil masks.
- C.** Example neuropil subtraction for two neurons. Black, somatic fluorescence; light red, peri-somatic neuropil signal; red, corrected fluorescence trace ($F_{\text{soma}} - F_{\text{neuropil}}$).
- D.** Touch fraction among neurons without peri-somatic neuropil correction (above) and with correction (below). The bottom plot is consistent with electrophysiological results.

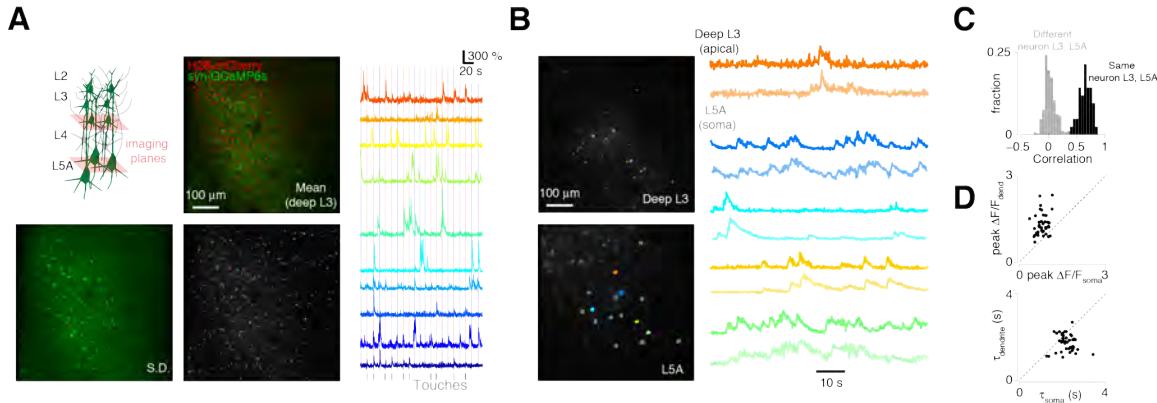


Figure S4. Imaging Layer 5 apical dendrites.

- A.** Imaging of L5A apical dendrites in L3. Top image, example plane deep in L3. Bottom-left, GCaMP6s fluorescence SD over a single imaging session (used for segmentation). Bottom-right, dendritic ROIS. Right, fluorescent transients from colored ROIs; grey ticks, touch.
- B.** Simultaneous imaging of deep L3 and L5A in behaving animals. Example responses are shown for both planes. Dark color, L3 apical dendritic response; light color, L5A somatic response for the same neuron.
- C.** Distribution of activity correlations between L3 and L5A for the same (black) and different neurons (grey).
- D.** Comparison of somatic (L5A) and dendritic (deep L3) peak $\Delta F/F$ and decay time constant, τ .

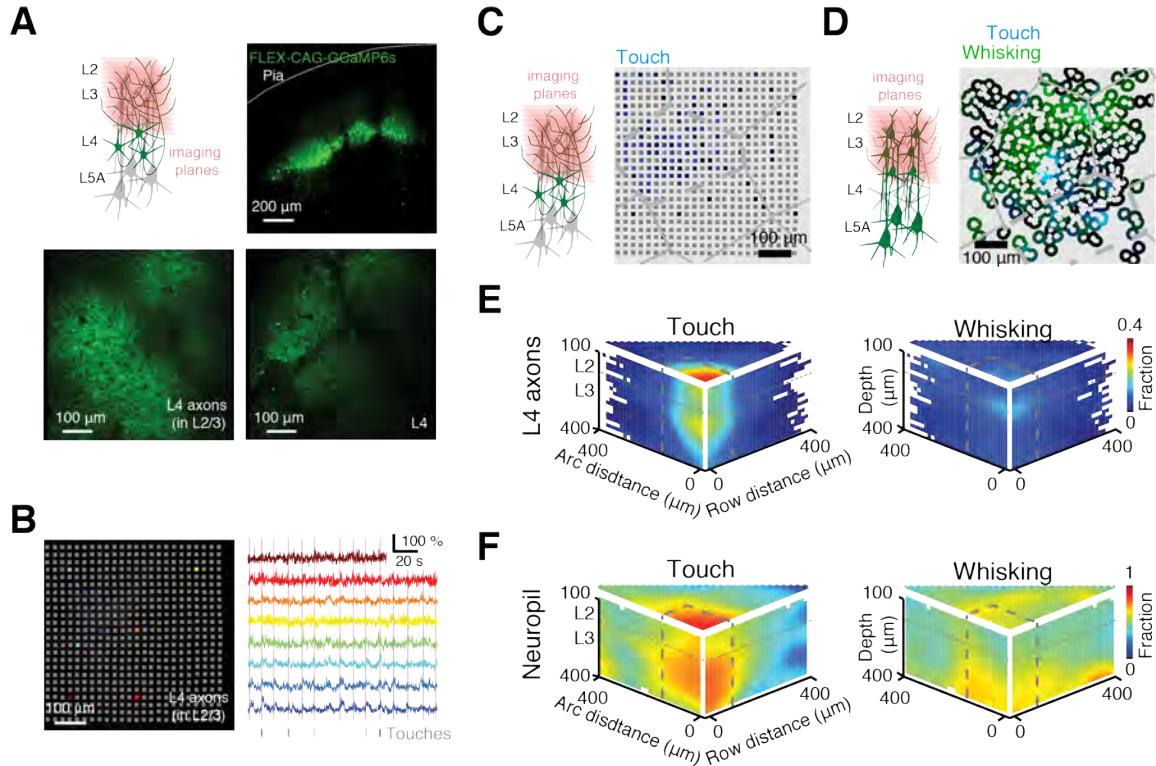


Figure S5. Maps of touch and whisking in L4 axons and L2/3 neuropil.

A. Imaging L4 axons in L2/3. Top, coronal section showing GCaMP6s expression obtained using FLEX-CAG-GCaMP6s injection into an *scnn1a-cre* mouse; bottom-left, tangential image of L2/3 showing L4 axonal labeling; bottom-right, tangential image of L4 showing somatic and process labeling, revealing barrels.

B. Example fluorescence ($\Delta F/F$) responses of L4 axons in L2/3. Fluorescent transients from ROIs colored in the top image are shown; grey ticks, touch.

C. Example imaging plane with L4 axon ROIs drawn in L2/3. Blue intensity, touch R_{fit} . Grey, unclassified. Thick dashed line, outline of the principal column; thin lines, surround columns.

D. Encoding in neuropil for same plane as in **Figure 7A**, with conventions as in **C**.

E. Touch and whisking fractional maps for L4 axons in L2/3 (n=3 mice). Color represents the fraction of ROIs in a given voxel classified as either touch or whisking. Location is given in terms of distance from the center of the spared whisker column (dashed grey line). The maps on top represent the row-arc plane, averaging cell fractions through depth. The arc-depth (left) and row-depth (right) maps average cell fractions across a half-barrel width in the plane-orthogonal direction. **E**.

F. Volumetric distribution of neuropil coding, as in **E** (n=8 mice).

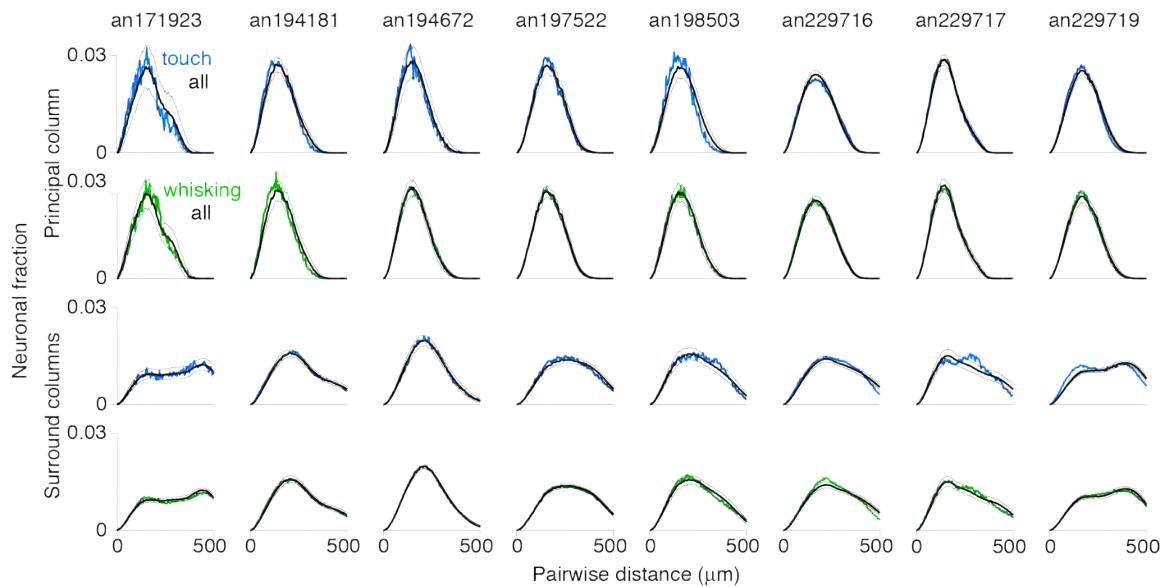


Figure S6. Spatial clustering among touch and whisking neurons.

Pairwise distance distributions for touch (blue) and whisking (green) excitatory L2/3 neurons, within and outside the spared whisker column, for each animal. Black lines, distance distribution for random drawings of the same number of neurons as the number belonging the relevant category from the entire population (thick lines, mean of 1,000 random drawings; thin lines, the 2.5 and 97.5 percentiles).

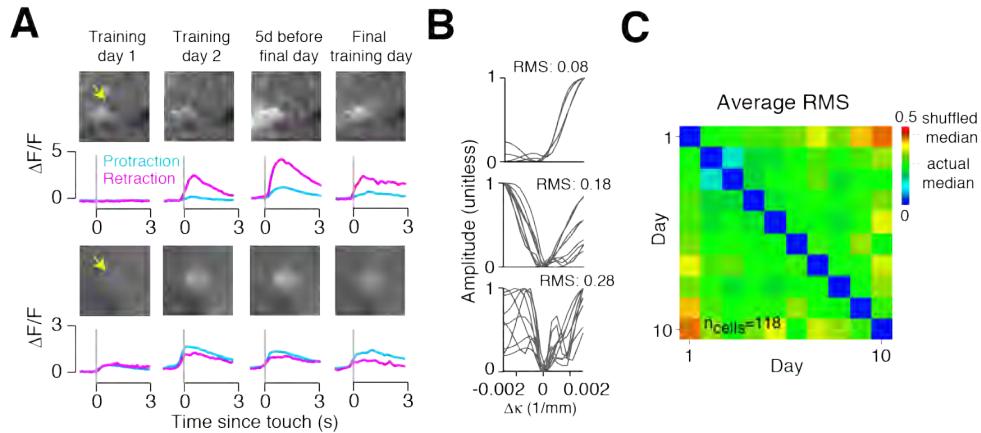


Figure S7. Dynamics of direction tuning during learning.

- A. Protraction and retraction aligned responses in two L2/3 touch neurons during learning.
- B. Touch receptive fields for 3 example L2/3 neurons across all days where the neuron was categorized as a touch cell. The mean pair-wise root mean square (RMS) was measured between all day pairs (i.e., all possible pairs of traces in each panel; Methods).
- C. Population RMS data for L2/3 pyramidal neurons strongly encoding touch (n=118 neurons; pooled across n=4 mice). Each square represents the mean pair-wise RMS across cells and animals when comparing those two days.

ID	Genotype	Red nuclear marker	Spared whisker	ROI count	Use
an171923	C57bl/6	None	C2	11,031	Volume imaging
an194181	Gad2-NLS-mCherry	GABAergic neurons	D2	9,637 (1,612)	Volume imaging, imaging during learning
an194672	emx1-Cre X LSL-H2B-mCherry	Excitatory neurons	C2	13,788 (1,913)	Volume imaging, imaging during learning
an197522	emx1-Cre X LSL-H2B-mCherry	Excitatory neurons	D2	16,200 (1,552)	Volume imaging, imaging during learning
an198503	emx1-Cre X LSL-H2B-mCherry	Excitatory neurons	C3	8,288 (1,301)	Volume imaging, imaging during learning
an229716	emx1-Cre X LSL-H2B-mCherry	Excitatory neurons	C2	12,293	Volume imaging
an229717	emx1-Cre X LSL-H2B-mCherry	Excitatory neurons	D3	19,012	Volume imaging
an229719	emx1-Cre X LSL-H2B-mCherry	Excitatory neurons	D2	18,125	Volume imaging
an224823	sccn1a-Tg3-Cre	None	D1	10,354	L4 axon imaging
an227254	sccn1a-Tg3-Cre	None	C1	13,681	L4 axon imaging
an238004	sccn1a-Tg3-Cre	None	C3	12,781	L4 axon imaging
an257218	emx1-Cre X LSL-H2B-mCherry	Excitatory neurons	C3	3,118	Anesthetized vs. awake directionality imaging
an257219	emx1-Cre X LSL-H2B-mCherry	Excitatory neurons	D2	1,804	Anesthetized vs. awake directionality imaging
an257221	emx1-Cre X LSL-H2B-mCherry	Excitatory neurons	C3	1,588	Anesthetized vs. awake directionality imaging

Table S1. Experimental animals. The first 8 mice were employed for volume imaging and imaging during learning (see **Figure 1D**). The 3 scnn1a mice were employed for bulk L4 axon imaging. Comparisons of direction maps in anesthetized and awake animals were carried out in the final 3 mice. Genotype dictated red nuclear expression, with Gad2-NLS-mCherry labeling GABAergic neurons and emx1-Cre X LSL-H2B-mCherry labeling excitatory neurons. Total ROI count consists of neurons and apical dendrites for learning and volume imaging mice; for L4 axon imaging, it consists of bulk axon ROIs. For animals imaged during learning, the number of neurons imaged during learning is indicated in parentheses.