Wiring and Molecular Features of Prefrontal Ensembles Representing Distinct Experiences

Graphical Abstract

Highlights
- Cohort-scale CLARITY is enabled for whole-brain activity/projection measurements
- mPFC cells active during distinct-valence experiences exhibit distinct projections
- Positive-valence experience preferentially recruits an NPAS4+ population in mPFC
- Recruiting diverse experience-defined mPFC ensembles drives distinct behaviors

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In Brief
A quantitative analysis of the wiring and molecular properties of neurons in the prefrontal cortex that are associated with distinct behavioral experiences illuminates the logic of information processing in the brain.

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Wiring and Molecular Features of Prefrontal Ensembles Representing Distinct Experiences

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SUMMARY

A major challenge in understanding the cellular diversity of the brain has been linking activity during behavior with standard cellular typology. For example, it has not been possible to determine whether principal neurons in prefrontal cortex active during distinct experiences represent separable cell types, and it is not known whether these differentially active cells exert distinct causal influences on behavior. Here, we develop quantitative hydrogel-based technologies to connect activity in cells reporting on behavioral experience with measures for both brain-wide wiring and molecular phenotype. We find that positive and negative-valence experiences in prefrontal cortex are represented by cell populations that differ in their causal impact on behavior, long-range wiring, and gene expression profiles, with the major discriminant being expression of the adaptation-linked gene NPAS4. These findings illuminate cellular logic of prefrontal cortex information processing and natural adaptive behavior and may point the way to cell-type-specific understanding and treatment of disease-associated states.

INTRODUCTION

The many regions and layers of the mammalian prefrontal cortex are known to contain cells with a rich diversity of activity patterns. Indeed, otherwise-indistinguishable populations of principal cells exhibiting profoundly distinct changes in activity in response to stimulus have been characterized by electrophysiological recording and cellular-resolution fluorescence Ca2+ imaging (Hayden et al., 2008; Insel and Barnes, 2015; Ito et al., 2003; Matsumoto et al., 2007; Pinto and Dan, 2015; Warden et al., 2012). At the same time, datastreams of anatomical and molecular information on prefrontal cell typology have emerged from a variety of methods, also pointing toward a rich diversity of principal neurons despite the traditional view that these cells were more homogenous than the highly diverse and readily separable interneurons (Hof et al., 1996; Lee et al., 2014; Soloway et al., 2002). These findings have highlighted the morphological, wiring, and electrophysiological diversity of principal neurons even within individual layers and subregions.

However, the mapping and correspondences (if any) among these different domains of diversity (activity during behavior, long-range wiring, and molecular phenotype) have remained largely unclear within prefrontal cortex. Addressing this issue would have fundamental implications for elucidating the cellular logic of prefrontal cortex function; moreover, differences in wiring, role in behavior, and molecular signatures among differentially responsive cells could provide long-elusive insight into the mechanisms of action of current neuromodulation therapies and even lay the foundation for developing new kinds of cell-targeted disease treatment. But the unique and non-stereotyped nature of each mammalian nervous system has prevented experimental resolution of this fundamental question, since cellular-resolution activity in a brain during behavior would have to be directly linked to quantitative local and global wiring, as well as to molecular analysis of the very same cells in the same mammalian brain, which has not yet been possible. Here, we address this question by developing approaches to quantify long-range anatomy, molecular signatures, and the causal impact of prefrontal cortical cells, defined by activity during distinct experiences.

RESULTS

Behavioral Cohort-Scale Whole-Brain Activity Mapping

We began with development and integration of enhanced ArcTRAP and CLARITY technologies. The ArcTRAP transgenic
mouse line was previously shown to translate temporally delimited neuronal activity into fluorescence via CreER/tamoxifen-mediated recombination (Figure 1A) (Guenthner et al., 2013). While the ArcTRAP line had been reported to have high background recombination, we developed an aqueous formulation of the fast-acting tamoxifen (4-hydroxytamoxifen, 4TM) enabling dosage reduction by an order of magnitude; this approach to ArcTRAP was found to eliminate the background recombination (Figures 1B and S1B). ArcTRAP mice can then be exposed to a rewarding or aversive experience (e.g., cocaine dosing at 15 mg/kg or a series of 0.5 mA/2 s foot shocks, respectively) with resulting robust and faithful labeling of neurons (Figure S1A); for example, histology confirmed that both experience types (with comparable but opposite conditioning value; Figures S1C and S1D) recruited neurons in basolateral amygdala (BLA; which is known to be involved in both reward and aversion processing) (Hsiang et al., 2014) (Figure 1B).

This approach brings untapped potential for obtaining deeper perspective on the cells activated, including insight into brain-wide patterns and molecular properties of cells recruited. Traditional thin-section histology would be prohibitively labor-intensive for whole-brain analysis in large behavioral-scale cohorts and also subject to sampling loss and bias, but the recent emergence of tissue transparency techniques, such as CLARITY, has, in principle, enabled lossless high-resolution brain-wide imaging compatible with molecular analysis (Chung et al., 2013; Tomer et al., 2014), if not yet in large cohorts of subjects. We therefore developed an enhanced CLARITY process suitable for this challenge of large-cohort analysis linked to behavioral experience;
key capabilities achieved included much faster speed and lower cost, as well as automated analysis algorithms adapted to both TRAP labeling and widely available commercial light-sheet microscope hardware (Figures 1A, 1C, 1D, and S1E–S1H; see the Experimental Procedures).

Employing this integrated TRAP/CLARITY approach, multiple cohorts (with temporally delimited exposure to appetitive cocaine dosing, aversive foot shock, or neither experience; n = 5 mice per group) were processed using two-color imaging on the light-sheet microscope. Image volumes taken in the green (autofluorescence) channel were iteratively co-registered, using an intensity-based non-linear registration algorithm, to generate an average reference brain. The registration based on the green channel from each brain was then used to warp the red channel (consisting of the TRAP signal) for each brain into this reference space (Figure S1I).

We first manually delineated potentially relevant regions in the reference brain, including mPFC, NAc (nucleus accumbens), BLA, BNST (bed nucleus of the stria terminalis), LH (lateral hypothalamus), PVN (paraventricular nucleus of hypothalamus), and CeA (central nucleus of the amygdala). At this level of inspection, the singly defined regional levels of neuronal activation were found to be largely similar between the two very different kinds of experience (Figures 2A–2D and S2H–S2J; Table S1), highlighting the need for a more refined analysis. To search for patterns in cell labeling that could differentiate the two

![Image](image_url)
conditions despite this apparent similarity, we manually registered the Allen Brain Atlas (ABA) (Oh et al., 2014) to the CLARITY reference. Locations of experience-recruited cells were also mapped into the reference, so that the number of recruited cells in every brain region could be quantified in an unbiased fashion (Menegas et al., 2015); changes in TRAP cell numbers (relative to home cage-animal mean values) in ~200 brain regions were then automatically quantified (Figures S2A–S2G).

Individual single-region activation patterns consistent across subjects within a group largely agreed with the manual analysis. For example, robust increases in TRAPped cells were observed in both settings within mPFC, NAc, BLA, pallidal regions (containing BNST), and hypothalamus (Figure S2K). However, the change in active cell count across all regions revealed common multi-region correlation patterns in mice from the same experimental group, suggesting experience-specific brain-wide patterns (Figure 2E). To investigate these differences, we applied principal component analysis (PCA) to the matrix of activated cell-count changes in cocaine- or shock-labeled mice. The first two principal components (PCs) comprised >75% of the variance for the dataset (Figure S2L) and thus were examined more closely. Plotting the position of each mouse in the space defined by the first two PCs revealed a clear separation between the behavioral conditions (Figure 2F).

To better understand the source of this separation, we examined the PC loadings across all of the included brain areas (Figure 2G). The first PC was found to reflect more general differences in activated cell recruitment across all regions (e.g., higher mean change across all regions in shock versus control mice; p < 0.05, t test). The second PC, by contrast, revealed a specific correlated group of DORpm (polymodal association cortex-related dorsal thalamus) regions, containing the lateral habenula (LHb), a key nucleus for aversive behaviors (Li et al., 2011; 2013). These regions were more strongly recruited by shock than by cocaine, selecting a responsive selectivity to the aversive experience. Consistently, manual quantification also found that shock more than cocaine, activated the LHb (p < 0.01 for shock versus home-cage experience; p < 0.05 for shock versus cocaine experience; Figure 2H). Thus, this behavioral cohort-scale TRAP/CLARITY labeling revealed brain-wide patterns of neuronal activation, as well as quantitative region-specific activation differences at the cellular level between different experimental conditions.

**Resolving mPFC Populations and Projections Activated by Appetitive or Aversive Experience**

Similarity in activation pattern by appetitive and aversive experiences has been reported in individually selected brain regions (verified broadly, though not in all regions, by the brain-wide analysis here) (Johnson et al., 2010; Xiu et al., 2014). A falsifiable hypothesis would be that the same neuron-type distribution was recruited by the two stimuli. In mPFC, the existing literature does not support or falsify this hypothesis, though mPFC is associated with specific reward and aversion processes (including cocaine-conditioned place preference, (Tzschenkte and Schmidt, 1998, 1999), fear and anxiety behaviors (Burgos-Robles et al., 2009; Lammel et al., 2011; Shah et al., 2004; Shah and Treit, 2003), and more general functions potentially relevant to the single-population hypothesis (including attention, salience- and novelty-detection, and working memory) (Dalley et al., 2004; Miller, 2000). The region-specific brain-wide analysis here may open the door to testing a distinct hypothesis—that appetitive and aversive experience recruit distinct neuron-type populations. Axonal projection anatomy is one of the most important features that might resolve principal cell population types involved in such distinct processes (Jennings et al., 2013; Kim et al., 2013; Lammel et al., 2012; Namburi et al., 2015), but this feature has been difficult to explore in a brain-wide fashion, while remaining linked (at the cellular level) to function during behavior.

A very strongly expressed activity-dependent cell-filling label (unlike traditional nuclear c-fos immunostaining or transiently or transgenically expressed fluorophores) in principle might allow for acquisition of this crucial wiring information as well from the same experimental subjects, provided that axon tracts of labeled neurons could be robustly imaged and quantified in this context. We therefore developed a CLARITY-optimized axonal-filling fluorescent protein, engineered in part, by inserting the 3’ UTR of neuritin (NRN) RNA (Akten et al., 2011) at the C terminus of EYFP (Figures S3A and S3B). We found that this construct could be packaged into high-titer adeno-associated virus (AAV) capsids that indeed enabled focal injection-defined projection labeling in CLARITY: for example, efferent mPFC projections could be readily followed throughout the entire adult mouse brain after a single stereotoxic injection (Figures 3A and 3B). Visualizing axonal tracks in 3D revealed key topographical features that were difficult, if not impossible, to detect in thin 2D sections (Figures 3A and S3C); for example, a prominent axon bundle traveling from mPFC to ventral medial thalamus was observed to carry out a sharp U turn near the VTA (Figures 3C and 3D), a potentially important feature that has not been described in existing atlases (Figure S3D).

Quantifying these tracts was also an unresolved challenge, and we therefore developed a method to compute 3D structure tensors from CLARITY images for tractography (Figures S3E–S3H). Faithful reconstruction of calculated streamlines was achieved (using tools adapted from diffusion tractography; Supplemental Experimental Procedures); these streamlines mapped onto fibers from CLARITY images (Figures 3E and 3F), and importantly, the streamline count in each bundle tightly correlated with the ground-truth physical diameter of the axonal bundles (Figure S3I). Using this method, we reconstructed whole-brain projections based on CLARITY images (Figure 3G); connectivity between a seed region (defined by stereotoxic injection site) and any specified target could be readily visualized and assessed by counting streamlines (Figures 3H and S3J–S3M).

To integrate this capability, with the needed additional capability of projection-labeling in cells defined by their use during experience, we sought to develop a viral CreER/4TM strategy to translate time-locked activity to sustained transgene expression. We therefore engineered a c-Fos promoter combining minimal promoter and regulatory elements in intron-1 (Figure S4A) (Barth et al., 2004; Schilling et al., 1991; Smeyne et al., 1992) that can be packaged into AAV particles and is specific enough to capture neuronal activity (Figures S4B–S4D). We also inserted a destabilized ER-Cre-ER-PEST...
Figure 3. CLARITY Enables Brain-wide Origin/Target-Defined Projection Mapping

(A) 2D orthogonal views of a mouse brain. Insert shows schematic for location of viral injection. D, dorsal; V, ventral; A, anterior; P, posterior; L, lateral; M, medial.

(B) 3D rendering of CLARITY hemisphere, visualizing outgoing mPFC projections (imaged by 2 × objective at 0.8 × zoom with a single FOV, step size: 4 μm, 1,000 steps).

(C) 3D visualization of the axonal bundle projecting from mPFC to VM (ventral medial thalamus), showing tracts turning near the VTA (indicated by arrows).

(D) Visualizing the same projection in (C) with sparse labeling by injecting a smaller volume of the same virus (25 nL).

(E) Raw image from a CLARITY volume. Orange: user-defined “seed region” so that only the fibers passing this region were tracked.

(F) Streamlines reconstructed from (E), using structural tensor-based tractography. Note that fibers in the CLARITY image that did not pass the user-defined seed region were excluded in the reconstruction (indicated by the magenta arrows).

(G) Reconstructed brain-wide streamlines from CLARITY image in (B). The streamlines are color-coded for orientation. A-P, red; D-V, green; L-M, blue.

(H) Representative computational isolation of mPFC fibers that project to VTA (yellow) or BLA (green).

Scale bars, 500 μm.

See also Figure S3.
cassette under this promoter (Kawashima et al., 2013; Li et al., 1998; Matsuda and Cepko, 2007); when injected into a reporter mouse, this viral CreER/4TM system reliably enabled activity- and tamoxifen-dependent cell-body and projection labeling (Figures S4E and S4F).

A final essential feature for quantitative activity-dependent projection mapping was normalization on an individual level to the absolute tract labeling strength independent of activity; this normalization is crucial in a virus-based approach to control for variation in injection efficacy. We enabled this feature (Figure 4A) by building in simultaneous two-color activity-independent (structural, EYFP) labeling and activity-dependent (tdTomato) labeling of projections from the same injection site, in which these two viruses infected cells with a stable relative ratio across.

Figure 4. Distinct Projection Targets of Cocaine and Shock-Activated mPFC Populations

(A) Summary of CAPTURE workflow (described in text).
(B) Representative CLARITY images of the structural projections (green, EYFP) and activity-dependent projections (white, tdTomato) from cocaine- and shock-labeled mice in NAc (top), LHb (middle), and VTA (bottom). Arrowheads indicate axon bundles terminating in the circled region. Scale bar, 200 μm.
(C) Reconstructed streamlines from (B), showing streamlines terminating in the 3D brain regions (purple). Green streamlines, reconstructed from EYFP fibers; red streamlines, reconstructed from tdTomato fibers. Scale bars, 200 μm.
(D–F) Quantification of projection intensity from cocaine- and shock-activated mPFC populations in three regions. Behavior-specific projection intensity was quantified using the ratio between red and green fibers (i.e., the number of red streamlines, divided by the number of green streamlines) terminating in indicated regions and plotted as fold change (relative to the cocaine group, n = 6 per group; ns, p > 0.05, *p < 0.05, **p < 0.01, unpaired t test). Error bars, mean ± SEM. See also Figure S4.
animals (Figures S4G and S4H). Dual-color quantification of projections to multiple downstream regions is then achieved by counting the number of streamlines ending in these regions, and the activity-dependence is corrected for anatomical and injection variability from the red/green streamline ratio at the individual animal level. This quantification of projection use across the brain from behaviorally defined neuronal populations is (for brevity) termed here CLARITY-based activity projection tracking upon recombination, or CAPTURE (Figure 4A).

Distinct Projection Patterns among Behavioral Experience-Defined mPFC Populations

We applied CAPTURE to quantify projections from cocaine- and shock-recruited mPFC populations. Ai14 reporter mice were co-injected with CaMKIIz-EYFP-NRN and cFos-ER-Cre-ER-PEST AAVs and subjected to 4TM-mediated cocaine and shock labeling. With CAPTURE, projections from all CaMKIIz (principally excitatory glutamatergic) neurons are labeled with EYFP and projections from behaviorally recruited populations are labeled with tdTomato. Importantly, EYFP fibers in the NAc, BLA, and VTA were found to be indistinguishable between the cocaine- and shock-labeled animals, indicating minimal variation in viral injection, transduction, and expression between the two groups (Figure 4B).

In the very same animals, significantly more projections from behaviorally activated mPFC neurons were observed targeting the NAc in cocaine-exposed animals compared to shock-exposed animals. Conversely, we observed significantly more behaviorally activated mPFC fibers to the LHb in shock-exposed animals (Figures 4C–4F). No significant difference in red/green (activity/structure) ratio was observed between the two groups in the VTA, showing no detectable systematic difference in efficiency or targeting of viral labeling. The cocaine-activated mPFC population thus preferentially projects to the NAc, whereas the shock-activated population projects more strongly to LHb, revealing that the neurons recruited in mPFC by distinct-activity experience are not simply different in terms of the patterns of input that they happen to receive, but represent anatomically distinct cell populations in terms of projection pattern across the brain.

Molecular Signature of Experience-Defined mPFC Neurons: Pre-existing NPAS4 Population

Next, we sought to identify molecular distinctions among cocaine- and shock-responsive populations in mPFC. We began by crossing ArcCreER mice to a Cre-dependent ribosome-tagged mouse (Long et al., 2014) to allow for labeling the ribosomes from neuronal populations activated by well-defined stimuli (Figure 5A). With this approach, the tagged ribosome can be immunoprecipitated (IP) from tissue lysate using an anti-GFP antibody, isolating translating mRNA for analysis (Ekstrand et al., 2014; Heiman et al., 2008; Sanz et al., 2009). These mice were exposed to either cocaine or shocks with 4TM. After labeling, mice were returned to the home cage for 2 weeks before initiation of ribosome purification, a design focused on identifying intrinsic signatures of these populations rather than on immediate mRNA expression changes associated with the exposure to stimuli.

Equal enrichment of GFP transcripts under these conditions was first confirmed by qPCR (Figure 5A). Four groups of mRNA (“IP fraction” for ribosome-associated transcripts from cocaine and shock and “input” for whole-tissue lysate from cocaine and shock; see the Experimental Procedures) were then analyzed by microarray. For both stimuli, IP fraction groups were enriched 3-fold in the ribosomal-component mRNA (Rpl10a) compared with input groups (Figures 5C and 5D), further demonstrating similar levels of mRNA enrichment between treatments. Gene expression profiles from cells recruited under the two conditions were highly similar upon initial examination (<10 genes showed >2-fold difference; Figure S5C); however, a transcript of the activity-dependent transcription factor Npas4 (Bloodgood et al., 2013; Lin et al., 2008; Shamloo et al., 2006) was uniquely enriched in cocaine-activated cells compared to shock-activated cells (Figure 5B). This enrichment was confirmed by comparing the IP fraction to the input from the cocaine-labeled cells (Figure 5C) and finally validated by qPCR (Figure 5E); in contrast, this enrichment was not observed by comparing the IP and input fractions from the shock-labeled population (Figure 5D).

Because Npas4 itself is an immediate early gene that can be transiently induced by neuronal activity, we first tested if its enrichment in the cocaine-recruited population could relate to a lasting general cocaine-elevated Npas4 transcription. Both cocaine and shock caused a rapid increase of Npas4 expression in mPFC within 30 min, as expected; however, this induction under both conditions, (also as expected) completely returned to baseline after 10 days before the ribosome was profiled (Figure S5B). Indeed, no difference was observed when we examined Npas4 expression in whole-tissue lysates from cocaine- and shock-labeled mPFC (Figure 5F), indicating that exposure to cocaine did not lead to an overall long-term increase of Npas4 expression in mPFC.

At the single-cell level, we discovered that the number of mPFC cells with NPAS4 immunoreactivity did not differ between behavioral groups or between the home-cage and behavior-challenged animals (Figures 5F–5H). However, TRAP/NPAS4 double-positive cell counts were significantly higher in cocaine-exposed mice (Figure 5I), consistent with activity recruitment of an NPAS4 population. Similarly, the percentage of NPAS4+ cells among TRAP+ cells was also significantly higher in cocaine-exposed mPFC than in shock-labeled mPFC (Figure 5J). Importantly, we found that another positive-valence experience (highly palatable food consumption), but not another negative-valence experience (restraint stress), also selectively recruited this mPFC NPAS4+ population (Figures S5D and S5E), further illustrating that a pre-existing NPAS4+ population is preferentially activated by salient stimuli associated with positive valence rather than negative valence. Finally, the molecular overlap between the positive valence-associated population and the NPAS4+ population led us to further examine the spatial distribution of valence-specific populations, since the NPAS4+ cells appeared to be enriched in superficial layers of the mPFC (Figures 5F and S5E); indeed, the positive-valence experiences were associated with a significantly larger fraction of TRAP+ ed layer 2/3 neurons in the mPFC than in the negative-valence experiences (Figure S5F).
Cocaine- and Shock-Activated Populations Control Appetitive and Aversive Behaviors

Next, we tested if electrical activity in these behavioral activity-defined populations had distinct positive or negative conditioning valence for the same animals that had experienced the stimulus, assessed by causal impact on behavior during the place preference task. To maintain robust, low-background, and direct region-specific control in mPFC, we designed a viral approach employing channelrhodopsin tagged with EYFP (ChR2-EYFP) under the control of the AAV-cFos backbone described earlier (termed fosCh; Figure 4A). Animals injected with this virus (in mPFC) were exposed to daily cocaine dosing or shock over 5 days. 

**Figure 5. Cocaine Preferentially Activates the NPAS4+ Population in mPFC**

(A) Schematic of activity-dependent ribosome profiling. Green, activated neurons; gray, non-activated neurons.

(B–D) Scatter plot of the most-enriched genes comparing cocaine- versus shock-activated cells (cocaine IP versus shock IP) (B), cocaine IP versus cocaine input (C) and shock IP versus shock input (D). The bottom right quadrant (pink) of each scatter plot denotes genes with p < 0.05 and fold change >2 for the indicated comparisons. As a positive control, the enrichment of RpL10a is highlighted in green. Black dots denote genes that were non-specifically enriched by IP (as shown enrichment in both cocaine and shock IP fraction). In (D), Npas4 (light-gray dot, fold change = 1.35, p value = 0.385) did not approach the statistically significant (pink) quadrant.

(E) Quantitative PCR analysis of Npas4 mRNA expression in the input and IP fractions.

(F) Representative images showing the overlap between TRAP+ and NPAS4+ cells in the mPFC. Scale bar, 100 μm. Arrowheads indicate double-positive cells.

(G–I) Quantification of numbers of TRAP+ (G), NPAS4+ (H), and TRAP+&NPAS4+ (I) cells in the mPFC.

(J) Percentage of NPAS4+ cells in TRAP+ cells under three conditions. n = 4 per group, *p < 0.05, **p < 0.01, unpaired t test. Error bars, mean ± SEM.

See also Figure S5 and Movie S1.
consecutive days, after which time robust increases in the number of fosCh-labeled cells and in EYFP expression were observed (Figure 6A–6C).

We first quantified *Npas4* expression in fosCh cells, hypothesizing that cocaine-labeled fosCh cells would exhibit higher *Npas4* expression compared with shock-labeled fosCh cells. This was indeed the case (Figure 6D); importantly, expression of general excitatory or inhibitory neuronal markers did not differ between those two populations (Figures S6A and S6B). Consistent with CAPTURE findings, cocaine-labeled fosCh cells were found to project strongly to NAc, while the LHb contained significantly denser fibers arising from the shock-labeled cells (Figures 6E–6G). Crucially, this method of targeting was sufficient to enable optical control over the resulting sparsely distributed neuronal subsets; fosCh-labeled cells displayed robust light-evoked firing assessed by in vivo electrophysiological recording (Figures 7A and 7B). Together, these data demonstrated resolution with the fosCh strategy of the same pattern that had been characterized molecularly and anatomically and enabled the final test of whether these neuronal subsets were capable of differentially controlling behavior.

To address this question, we employed the real-time place preference paradigm in which 10 Hz light-pulse trains were automatically triggered on entry into one side of a behavioral chamber. Place preference was monitored over three consecutive trials: before, during, and after light delivery for reactivation of fosCh-defined neuronal ensembles (Figure 7C). To control for the possibility that behavior could be biased by randomly labeled neurons, an additional group of animals was included in which comparable expression of ChR2 was driven by the CaMKIIα promoter without link to prior activity (Figures S7A and S7B). Optogenetic stimulation of these non-activity-specific neuronal populations did not influence place preference, nor were homecage-recruited fosCh-population animals observed...
to exhibit preference or aversion for the chamber in which the cells were optically activated. Remarkably, however, reactivation of the shock- or cocaine-defined fosCh populations induced significant (and opposite-direction) shifts in place preference, with cocaine-exposed mice demonstrating preference, and shock-exposed mice demonstrating aversion, for the photostimulation-paired side (Figures 7D and 7E; mean preference change at post-test for cocaine: 1.3x ± 0.1, Wilcoxon p = 0.0006; for shock: 0.8x ± 0.1, Wilcoxon p = 0.002). These data reveal that the activity-defined mPFC neural populations differ not only anatomically and molecularly but also in functional impact in modulating behavior.

**DISCUSSION**

It has been remarked (building on Dobzhansky’s comment that “nothing in neurobiology makes sense except in the light of evolution” (Dobzhansky, 1973) that “nothing in neurobiology makes sense except in the light of behavior” (Shepherd, 1988). This perspective may be directly relevant to efforts in multiple subfields of neuroscience to describe the cellular diversity in nervous systems. While these efforts are pioneering and informative, it may be limiting to collect cellular typology datastreams relating to anatomy, genetics, and membrane biophysics, without knowledge of the importance of activity patterns in those same cells during behavior. However, forging such direct linkages has been difficult, especially in vertebrates. Indeed no system may be more challenging in this regard than the mammalian mPFC, which does not have well-defined direct sensory input or motor output pathways, and wherein principal cells with similar spike properties and anatomical positioning exhibit divergent or even opposing responses during behavior. Here, we have addressed these challenges, focusing on the specific problem of whether mPFC cells that respond differently to positive or negative...
valence experiences are, in fact, separable cell types as definable by molecular or anatomical measures, as well as by causal impact on behavior.

We identified not only brain-wide projection patterns separating mPFC cells responding to positive- and negative-valence experience but also a unique and unexpected enrichment of rating mPFC cells responding to positive- and negative-valence experiences (Figure 4; Movie S1) that was not observed in cells responding to salient aversive experiences with comparable conditioning value. In illustrating that a pre-existing NPAS4+ population is preferentially activated by salient stimuli associated with positive valence rather than with negative valence, these mPFC results may be contrasted with elegant memory-encoding models in which cells expressing a key activity marker may be recruited into diverse stimulus representations, as with CREB-expressing cells in amygdala fear conditioning (Han et al., 2007). NPAS4 has been linked to cellular and microcircuit processes relating to the balance of excitation and inhibition, including a form of adaptive plasticity in which inhibitory synapses are formed on cells in response to high levels of excitation (Bloodgood et al., 2013; Lin et al., 2008; Maya-Vetencourt et al., 2012; Spiegel et al., 2014). This previously known role of NPAS4, together with the observation here that the positive-valence, but not negative-valence, experience recruited the mPFC NPAS4 population may bear relevance to previous observations among behaviorists that appetitive-learning experiences may be more vulnerable to adaptation or extinction, whether with natural stimuli or in pathological situations seen in drug dependence (Bouton, 2004).

The CAPTURE approach (the integration of enhanced CLARITY for behavioral cohort-scale work with cell and projection quantification linked to activity during behavior) is provided here in the form of tools, software, and datasets that may be readily applied across systems. The intact-brain approach is not only scalable in a practical manner to include large cohorts of experimental subjects and enabling of quantitative projection and population analysis but also helps address key confounds relating to selection bias and loss of continuity in traditional sectioning approaches widely used across neuroscience and biology. Moreover, improved 3D understanding of the path of behaviorally relevant axon bundles through the intact brain may be critical for the expanding field of neuromodulation, which is increasingly dependent on accurate targeting of white matter tracts (Henderson, 2012).

These tools are also readily adaptable for broader applications. For example, the NRRN construct allowing rapid axonal projection labeling may be useful along with recombinase- or tetracycline-controlled (Liu et al., 2012) promoters for driving axon-filling fluorophores in approaches to multiplexed cell-type-specific CLARITY projection mapping or for driving opsins (Gore et al., 2015) to enhance functional assays of long-range projections (for example, to test synaptic release at different points along an axonal projection in slice assays or to test behavioral significance of cells defined by projection target). The fos-CreER virus (Figure 4) might be used in conjunction with future generations of high-potency opsin expression systems to enable optogenetic control based on experience temporally defined with 4TM, in strategies that could be eventually developed to reduce the number of repeated exposures currently required for sustained expression using the fosCh construct (Figure 7C).

Lastly, recombinase-dependent constructs expressing Ca2+ indicators (such as G/RCaMPs) may be used to resolve temporal dynamics and individual-neuron activity magnitudes within experience-defined cells; these features are not accessible by conventional binary IEG-based methods but may be important, for example, in positive versus negative reinforcement in the PFC (Pi et al., 2013; Pinto and Dan, 2015).

In future work, it will be important to build on these results with additional investigation of the mPFC populations. For example, exploring long-term changes in gene expression and structural properties in these assemblies after experience (continuing to leverage the activity-tagging and intact-brain methods) may help to identify cell-population-resolved processes important in plasticity of adaptive or maladaptive circuit dynamics. And identification of these separable mPFC populations, along with analysis of their interactions and modulations during adaptive and pathological conditions (in the setting of rapid parallel advances in high-content molecular phenotyping, targeted drug screening and design, and genome editing) may point the way toward cell-population-targeted interventions in diseases of addiction, anxiety, and fear.

**EXPERIMENTAL PROCEDURES**

**Animals**
Male and female C57BL/6J mice were group-housed on a reverse 12-hr light/dark cycle. Mice were 6–8 weeks old at viral infusion. Ai14 and wild-type C57BL/6 mice were purchased from JAX. Rosa26lox-stop-lox-eGFP-L10 mice were from Dr. Evan Rosen at Harvard Medical School. Male mice were used in behavioral assays. Both male and female mice were used for anatomy assays. All experimental protocols were approved by the Stanford University Institutional Animal Care and Use Committee and were in accordance with the guidelines from the NIH.

**CAPTURE Labeling**
Ai14 mice were injected with a 1-μl mixture of AAV8-CaMKIIz-EYFP-NRN and AAV8-Cfos-ER-Cre-ER-PEST in the mPFC. 2 weeks later, the mice were given 15 mg/kg cocaine (IP) or 20 random foot shocks (2 s/0.5 mA, two shocks/minute on average) for 2 consecutive days. The control group remained in their home cage for the whole period. 10 mg/kg 4-TM was given to all mice 3 hr after the last behavior section to enable CreER-mediated recombination. The mice were returned to their home cages for an additional 4 weeks to allow the full expression of fluorophores.

**CLARITY and Whole-Brain Imaging**
Three key features of the new CLARITY platform were (1) accelerated clarification through parallelized flow-assisted clearing crucial for large cohorts and independent of specialized equipment, such as electrophoresis or perfusion chambers, (2) >90% cost reduction using a new refractive index-matching process, and (3) optical properties such that the whole mouse brain can be imaged using a commercial light-sheet microscope (Lavision) under a single field of view and as a single stack in less than 2 hr with single-cell resolution (this speed and simplicity is also useful for large behavioral cohorts). The COLM system (described earlier in Tomer et al., 2014; Lerner et al., 2013) is optimized for high-resolution imaging and is most useful for studies requiring sub-cellular information, while the approach outlined here is optimized for speed and practicality in cellular resolution work, most useful for robust application to large cohorts in CLARITY (see the Supplemental Experimental Procedures). All custom software, including the scripts used for atlas building, registration, and analysis, as well as code for CLARITY-based tractography, are freely available in a unified package at http://capture-clarity.org. The Web
site also provides detailed protocols, sample images, and other CLARITY-related resources.

SUPPLEMENTAL INFORMATION

Supplemental Information includes Supplemental Experimental Procedures, seven figures, one table, one movie, and one data file and can be found with this article online at http://dx.doi.org/10.1016/j.cell.2016.05.010.

AUTHOR CONTRIBUTIONS


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