A comprehensive thalamocortical projection map at the mesoscopic level

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The thalamus relays sensori-motor information to the cortex and is an integral part of cortical executive functions. The precise distribution of thalamic projections to the cortex is poorly characterized, particularly in mouse. We employed a systematic, high-throughput viral approach to visualize thalamocortical axons with high sensitivity. We then developed algorithms to directly compare injection and projection information across animals. By tiling the mouse thalamus with 254 overlapping injections, we constructed a comprehensive map of thalamocortical projections. We determined the projection origins of specific cortical subregions and verified that the characterized projections formed functional synapses using optogenetic approaches. As an important application, we determined the optimal stereotaxic coordinates for targeting specific cortical subregions and expanded these analyses to localize cortical layer–preferential projections. This data set will serve as a foundation for functional investigations of thalamocortical circuits. Our approach and algorithms also provide an example for analyzing the projection patterns of other brain regions.

Anatomical connections provide structural substrates for information processing in the brain, yet neuroanatomical maps in most model organisms are incomplete1. This is especially true in mouse, where there are few comprehensive characterizations of anatomical connectivity despite it being a primary model for studying neural function1. Anatomical connectivity at the mesoscopic level is critical for understanding of how circuits subserve behaviors and is necessary for investigation of circuit function using genetic manipulation1–3.

The thalamus is integral to the flow of information into and out of the brain via its extensive interconnection with the peripheral and central nervous systems4–9. Thalamocortical projections are the primary drivers of cortical activity in sensory areas5 and associative brain regions, such as the frontal cortex10–12. The thalamus contains ~40 nuclei13,14, each innervating a different combination of cortical areas. Thalamic inputs to the frontal cortex are poorly characterized compared with thalamic inputs to primary sensory cortices, and our knowledge of the thalamo-frontal pathway is based on an amalgam of tracing studies from primates, cats and rats spanning several decades4. Gaining a complete representation of each thalamo-frontal projection pathway from these studies has been difficult because of variability between techniques and inconsistencies in anatomical boundary definitions4. A systematic characterization of thalamo-frontal pathways is necessary for investigating the function of frontal subregions.

It remains challenging to create a comprehensive thalamocortical projection map from individual thalamic subdivisions in mouse. First, the potential target area spans the entire cortex, necessitating a high-throughput microscopic method that can image the projections throughout the cortex at sufficiently high resolution and sensitivity5. Next, demarcating the cytoarchitectural boundaries for mouse thalamic nuclei is difficult because they are less distinct than the boundaries in other mammalian brains4. Furthermore, a comprehensive neuroanatomical data set requires robust analysis methods to combine anatomical data across experimental animals15. Finally, it remains a major challenge to process, analyze, summarize and present large anatomical data sets.

To overcome these challenges, we developed a high-throughput approach using bilateral, two-color, anterograde, focal viral injections into mouse thalamus. We then injected injected brains at submicrometer resolution, providing single axon sensitivity. We developed algorithms to localize injections in a model thalamus, allowing us to compare injection and projection information across animals. We identified the origins of thalamic inputs to 19 cortical subregions in mouse, focusing on poorly understood thalamo-frontal pathways. We further localized the origins of layer-specific cortical projections to vibrissal motor cortex (vM1). On the basis of coordinates extracted from our analyses, we performed viral injections encoding channelrhodopsin, and optogenetically confirmed that the anatomically characterized projections formed functional synapses. Our data provide a practical guide for viral injection, imaging and manipulation of thalamocortical circuits in mice. This method and associated analyses can be adapted to develop comprehensive neuroanatomical connectivity maps in other brain regions.

RESULTS

Labeling and imaging thalamocortical projections

To visualize thalamic projections, we stereotaxically injected two recombinant adeno-associated viruses (serotype 2/1, AAV2/1)16–19 encoding eGFP and tdTomato, respectively, bilaterally into the mouse thalamus (Fig. 1a–c and Supplementary Fig. 1). Thalamic projections did not cross the midline in mouse20 (Supplementary Fig. 1a), which allowed us to inject, image and analyze each hemisphere independently. Bilateral, two-color viral injections quadrupled the

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throughput of subsequent data collection, consolidated the total amount of data (~0.5 TB per mouse) and eased computational demands for data processing. In addition, two-color labeling highlighted topographic projection patterns from adjacent thalamic volumes (Supplementary Fig. 2). By using a hydraulic apparatus to deliver ~10 nl of AAV, a consistently small infection volume was achieved (measuring 0.30 ± 0.23 mm³, corresponding to ~1.6% of the total thalamic volume, and of 630 ± 160 µm (n = 188 injections) wide in the medial-lateral axis; Fig. 1c). We detected fluorescent protein in 67.4 ± 10.3% of cells at the injection center, with an 88.8 ± 4.4% infection rate for neurons (Supplementary Fig. 1b–e).

Brains were paraformaldehyde fixed and cryostat sectioned coronally at 50 µm (Fig. 1a–c). All sections of each brain, from the start of the frontal cortex through the end of the thalamus, were fluorescently imaged in their entirety under identical conditions using a Hamamatsu Nanozoomer imaging system (0.5 µm per pixel),
providing sufficient sensitivity to detect single axons (Fig. 1b). The thalami were re-imaged to reduce saturation of the injection sites (Fig. 1c). We successfully imaged 75 mouse brains containing a total of 254 injections, resulting in ~40 TB of imaging data.

An overview of data analysis
To analyze and compare thalamic injections across animals, we developed a suite of custom algorithms using MATLAB (MathWorks). The goal of these algorithms is to align individual injection sites onto a model thalamus (Fig. 1d), such that injection and projection information can be compared across brains (Supplementary Fig. 3). We manually traced each thalamus from the section images to generate a binary thalamus mask. Injection sites were masked by applying an intensity threshold to the images using a threshold determined by Otsu’s method (Fig. 1c and Supplementary Fig. 3b). We then aligned and stacked each brain’s thalamus mask sections to create a three-dimensional volume mask (Fig. 1d, Supplementary Fig. 4 and Online Methods). We normalized the three-dimensional masks and their corresponding injection site masks, corrected them for variability in cutting angle, and aligned them using anatomical landmarks (Supplementary Figs. 4 and 5 and Online Methods). The aligned thalamic structures were then averaged to produce a model thalamus (Supplementary Figs. 3c and 6a), and each injection site was mapped onto the model (Fig. 1d).

We then determined the cortical projection targets for each injection and combined the injection and target information for all 254 injections to localize the precise thalamic origin of the cortical projections (Supplementary Fig. 3d). We aligned two widely used atlases to the model for nucleus-specific analysis (Supplementary Fig. 3e). Notably, we also used our comprehensive data set to create a nucleus-independent assessment of subdivisions in the thalamus (Supplementary Fig. 3e).

Assessment of thalamic alignment and injection coverage
After normalization and alignment (Online Methods), individual thalami were highly similar to each other, with 3.7% variability in the thalamic volume (percent s.d.) and 102 ± 51 µm (mean ± s.d.) variability in the thalamic border location (Fig. 2a and Supplementary Fig. 6b,c). This variability is nearly identical to that measured with alternative data collection methods such as serial block-face imaging (102.5 ± 45 µm) (24). The high degree of similarity between the individual masks and the model thalamus was confirmed using Dice’s coefficient (D = 0.94 ± 0.01; Supplementary Fig. 6d). To facilitate subsequent data analysis, we down-sampled thalamic masks to a voxel size of 36.4 × 36.4 × 50 µm (x, y, and z, respectively), which is more than twofold smaller than the variability across individual thalami.

We aligned our model thalamus to two atlases: the Allen Brain Atlas (ABA; http://mouse.brain-map.org/) and the Paxinos Mouse Brain Atlas (PMBA) (25) (Fig. 2b and Supplementary Fig. 3e; refer to Table 1 for all anatomical structure abbreviations). To verify this alignment, we traced four cytoarchitecturally identifiable structures (AD, AV, PT and fr) from five randomly selected experimental brains and compared them with their corresponding atlas structures (Fig. 2b,c). The overall shape, orientation and location of the thalamic structures were highly similar among the brains and atlases, as quantified using Dice’s coefficient (Fig. 2c and Supplementary Fig. 6e). Although variability across brains remained, the structures from experimental brains were as similar to the atlases (D = 0.53 ± 0.10, n = 4 thalamic structures) as the atlases were to one another (D = 0.60 ± 0.11; P = 0.35, t test). We concluded that the alignment of individual nuclei to our model was accurate.

We distributed the injections throughout the thalamus (Supplementary Fig. 7a) such that 93.4% of the thalamus was covered by at least one injection and 85.3% was covered by at least two injections (Fig. 2d–f and Supplementary Fig. 7c). The majority of thalamic nuclei are fully covered (Fig. 2f and Supplementary Fig. 7b); however, we excluded the geniculate nuclei from the data set. The center of the thalamus was more highly sampled because injections that extended beyond the lateral or ventral borders of the thalamus were excluded (Fig. 2d and Supplementary Fig. 7a,c).

Mapping the thalamic origins to cortical targets
Using this data set, we sought to identify the thalamic sources of projections to each of 19 cortical subregions of interest (ROIs), which were defined by their boundaries in the PMBA (Fig. 3a). We noted the strength and specificity of projections from each of our injections to all

| Table 1 Abbreviations of anatomical structures |
|-----------------------------------------------|
| Abbreviation | Definition | Paxinos location |
|----------------|------------|-----------------|
| Thalamic structures | | |
| AD | Anterodorsal nucleus | AD |
| AM | Anteromedial nucleus | AM + AMV |
| AV | Anteroventral nucleus | AV + AVDM + AVVL |
| CL | Central lateral nucleus | CL |
| CM | Central medial nucleus | CM |
| fr | Fasciculus retroflexus | fr |
| IAD | Interanterodorsal nucleus | IAD |
| IAM | Interanteromedial nucleus | IAM |
| IMD | Intermediodorsal nucleus | IMD |
| LD | Laterodorsal nucleus | LD + LVPL + LDDM |
| LP | Lateral posterior nucleus | LP + LPL + LPML + LPMC |
| MD | Mediodorsal nucleus | MDC + MDL + MDM |
| PCN | Paracentral nucleus | PC + OPC |
| Pf | Parafascicular nucleus | Pf |
| Po | Posterior nucleus | Po |
| Pr | Preriuneuni nucleus | vRe |
| PT | Parataenial nucleus | PT |
| PVT | Paraventricular nucleus | PVA + PV |
| RE | Reuniens nucleus | Re |
| RH | Rhomboid nucleus | Rh |
| SGN | Suprageniculate nucleus | SG |
| SMTr | Submedius nucleus | Sub |
| SPFp | Subparafascicular nucleus | SPFpc |
| VAL | Ventral anterior-lateral complex | VA + VL |
| VM | Ventromedial nucleus | VM |
| VPL | Ventral posterolateral nucleus | VPL + VPLpc |
| VPM | Ventral posteromedial nucleus | VPM + VPMpc |
| Cortical subdivisions | | |
| Al | Anterior insular cortex | pregenual (AI + AID + AIv + DI + G1) |
| Aud | Auditory cortex | Au1 + AuD + AuV |
| dACC | Dorsal anterior cingulate cortex | Cg1 |
| FrA | Frontal association area | FrA |
| IL | Infralimbic cortex | IL |
| Ins | Insular cortex | postgenual (AID + AIv + AIP + DI + G1) |
| LO | Lateral orbital cortex | LO + DLO |
| M1 | Primary motor area | M1 |
| M2 | Secondary motor area | M2 |
| MO | Medial orbital cortex | MO |
| PrL | Prelimbic cortex | PrL |
| Pt | Parietal association cortex | MPFA + MPFA + PTPR + PPTrD |
| Rhi | Rhinal cortex | Ect + Prh + Lent |
| RS | Retrosplenial cortex | RSA + RS |
| S1/2 | Sensory cortex | S1 (all sub-regions) + S2 |
| Tem | Temporal association cortex | TeA |
| vACC | Ventral anterior cingulate cortex | Cg2 |
| Vis | Visual cortex | V1 (all sub-regions) + V2 (all sub-regions) |
| vM1 | Vibrissal motor cortex | M2 |
| VO | Ventral orbital cortex | VO |
cortical areas using a manual scoring system (Supplementary Fig. 8). Independently, three experts blindly performed this analysis.

We then used the projection scores for each injection to perform a simple injection site grouping method, which allowed us to localize the thalamic sub-volumes projecting to each cortical ROI. First, to account for the alignment variability between thalami (102 ± 51 µm), we eroded each aligned injection site by 100 µm to produce the injection ‘core’ (Figs. 1c and 3b). The core, as compared with the periphery, represents the volume of an injection that we were more confident is accurately localized in the model thalamus.

Next, we combined the volumes of all injections that projected to a given ROI (positive injections) and then subtracted the volumes of the injections that did not project to that region (negative injections) from the combined total. This process resulted in a better localization of the thalamic volume projecting to each ROI than if the summed positive injection volume was used alone (Fig. 3b,c and Supplementary Fig. 8e–g). By employing this method, we localized volumes at a finer resolution than at the size of a single injection. The grouping method described above was expanded to assign higher confidence to injection site cores, as well as account for different projection properties,

Figure 3 Localization of the thalamic origins of cortical projections. (a) Illustration showing the 19 cortical areas examined (modified from PMBA25 and ref. 47): whole brain views (lower) and coronal sections (upper). (b) Simplified schematic of thalamic localization method. (i) The volume of each injection is eroded to generate an injection core (Fig. 1 and Online Methods). (ii) Injections that send projections to an ROI (positive injections) are summed and those that do not (negative injections) are subtracted. (iii) This results in the precise thalamic volume projecting to each ROI, which is separated into eight confidence levels (Supplementary Fig. 8). (c) Example thalamus section illustrating the injection grouping method summarized in b. (i,ii) injections that did project and did not project to a target. (iii) The refined thalamic volume projecting to that target (that is, the confidence map). (d) Diagram of the thalamus sections shown in e. Axes are distance relative to bregma (mm). Coronal section −1.16 mm posterior to bregma (top), horizontal section −3.52 mm ventral to bregma (bottom). (e) Confidence maps (gray scale) show the thalamic origin of projections to nine frontal brain areas (Supplementary Fig. 9). (f) Localization of exclusive and shared thalamocortical projections to PrL (green) and IL (magenta) through direct comparison of their confidence maps (right). Nuclear boundaries are shown on the left half of each thalamus section (PMBA). Coronal section −0.61 mm posterior to bregma (top), horizontal section −4.06 mm ventral to bregma (bottom). Scale bars represent 1 mm.
Figure 4  Localizing thalamic subdivisions on the basis of cortical projection patterns. (a) Summary of confidence maps to all cortical subregions, clustered on the basis of confidence map similarity (determined in c; Supplementary Fig. 9). (b) The thalamus was down-sampled into 150 × 150 × 150 µm voxels (left) and the average confidence level in each voxel was determined for each cortical projection (example, right). (c) All thalamic voxels (rows) were hierarchically clustered on the basis of their cortical projection patterns, and cortical subregions (columns) were clustered on the basis of which thalamic voxels innervated them. The average confidence level is indicated in gray scale, as in a. A threshold (gray dashed line, left) was applied to identify 11 distinct clusters. (d) Coronal thalamus sections showing the spatial location of clusters from c (left), with the corresponding atlas sections (PMBA, left; ABA, right) showing thalamic nuclear groups for comparison (right). (e) Schematic showing the convergence and divergence of projections for several clusters. (f) Overlap between voxel clusters (rows) and atlas-defined nuclear groups (columns). Colored boxes highlight the clusters that are dominant in (compose >10%) of the anterior and medial thalamic groups. Some nuclear groups were covered by relatively few clusters that have closely related projection patterns (for example, the anterior group mainly contains clusters 1 and 2), whereas other groups contained clusters with disparate projection patterns (for example, the medial group contains clusters 2 and 5–10). (g) Coverage of the anterior and medial nuclear groups by each voxel cluster.
such as strength and specificity (Supplementary Fig. 8 and Online Methods). This analysis resulted in confidence maps in which the value of each thalamic voxel, that is, volumetric pixel, indicates our certainty that the thalamic voxel projects to a particular ROI (Fig. 3d,e and Supplementary Fig. 8d–g), where a confidence value of 8 is highest and a confidence value of 0 indicates that no projections originated from that voxel. We determined confidence map summaries for the nine subregions of the frontal cortex: FrA, dACC, vACC, PrL, IL, MO, VO, LO and AI (Fig. 3d,e, Supplementary Movies 1–9 and Supplementary Fig. 9). Each confidence map contains a continuous positive volume, which is unique for each target region. To validate the projection sources predicted by our confidence maps, we performed injections of fluorescent retrograde beads in a subset of our characterized areas (Supplementary Fig. 10). We observed that all retrogradely transported beads were localized in the predicted confidence map.

One advantage of having confidence maps across many cortical subregions is that we could directly compare the thalamic origins of functionally related cortical subregions. For example, PrL and IL are both crucial in fear learning, but PrL is associated with the ‘high fear’ behavior state, and IL is associated with the ‘low fear’ state. By comparing the confidence maps for PrL and IL, we localized the shared and unique thalamic origins to PrL and IL (Fig. 3f), suggesting that differential thalamic inputs may contribute to their functional differences. Such comparisons allow for the selective targeting of thalamic projections to PrL and IL for future functional studies.

Defining thalamic subdivisions based on cortical targets
The thalamus is commonly subdivided into anatomically and functionally similar nuclear groups. Although useful, these divisions ignore ambiguity in nuclear borders, differences in projection patterns in a single nucleus and the possibility that cytoarchitecturally defined nuclei may not always be the relevant functional unit in the thalamus. Given that our confidence maps provide distinct topographic information (Fig. 4a), we determined whether the thalamus could be instead subdivided on the basis of cortical projection patterns alone.

The thalamus was divided into 150 × 150 × 150 μm voxels (Fig. 4b), which were then clustered (agglomerative hierarchical clustering, MATLAB) on the basis of their confidence values for all 19 cortical subregions (Fig. 4b,c). We applied a threshold to identify the 11 largest thalamic voxel clusters (Fig. 4c–e). Notably, the thalamic voxels comprising each cluster were spatially grouped and largely continuous, and similar to the thalamic nuclear groups (Fig. 4d). However, the voxel clusters and nuclear groups were not identical. Although several nuclear groups consisted of one or two closely related clusters (anterior and intralaminar nuclear groups; Fig. 4f,g), other nuclear groups contained several largely divergent clusters (for example, the medial and ventral groups; Fig. 4f,g), suggesting functional homogeneity in some nuclear groups, but substantial heterogeneity in others.

Optimal injection sites and functional confirmation
Stereotaxic viral delivery of optogenetic and pharmacogenetic reagents to manipulate neuronal activities has become an important method for dissecting functional circuitry. Currently, studies involving the mouse thalamus that employ these methods primarily rely on the empirical determination of the injection coordinates based on a small number of trials. Using our confidence maps, we simulated injections throughout our model thalamus and determined the optimal injection coordinates for targeting projections to a specific ROI (Fig. 5a and Supplementary Fig. 11).

Given that anatomical projections do not always guarantee functional connectivity, we sought to verify that the observed anatomical axonal projections form functional connections at each target region, which also allowed us to verify the validity of the optimal injection coordinates. We injected AAV2/1 expressing channelrhodopsin2 (ChR2) using our optimal injection coordinates to target thalamic projections to eight frontal subregions. Whole-cell recordings were made in each projection target area (dACC; Fig. 5b) and postsynaptic responses were observed following activation of the ChR2* thalamic axons with blue light stimulation (Fig. 5b–d and Online Methods). We observed excitatory responses in 48 of 50 recorded cells; specifically, we recorded responses from 4 of 4 cells in AI, 13 of 13 in VO/LO, 4 of 4 in MO, 3 of 5 in IL, 5 of 5 in PrL and 19 of 19 in dACC/vACC (Fig. 5d), indicating that the anatomically defined projections corresponded to functional thalamocortical synaptic connections.

Grouping thalamic nuclei on the basis of cortical targets
As described earlier, nucleus locations from both the ABA and PMBA were aligned to our model thalamus (Fig. 2b,c), allowing us to localize the origins of cortical projections to individual nuclei (Fig. 6a–c). To compute the fraction of each nucleus that projects to a given ROI,
we overlaid the nuclear boundaries aligned from the atlases onto the confidence maps (Fig. 6b) and averaged the coverage of a given nucleus between the two atlases. We determined the coverage distribution across nuclei for projections to select frontal subregions (Fig. 6a and Supplementary Fig. 12). We performed a cluster analysis using the nuclear localization of the confidence data for all 19 cortical subregions to identify projection patterns across thalamic nuclei (Fig. 6e). Functionally related cortical subregions formed tight clusters when grouped according to the origin of their thalamic inputs, suggesting that our comprehensive anatomical data set can be predictive of functional relationships, which validates our approach. It is important to note that there are limits to the resolution of this method: small (<300 μm wide) and intricately shaped nuclei are difficult to separate from their neighbors.

We compared our nuclear-localized thalamocortical projection data to previously described data for rat (Fig. 6a and Supplementary Figs. 12 and 13), as primary anatomical data for mouse is sparse. Overall, our nucleus projection data were largely consistent with the cumulative rat anatomical data, but we did find discrepancies between our findings and the rat literature (Fig. 6a and Supplementary Fig. 13a). Several factors may contribute to these discrepancies. First, the boundary definitions between cortical subregions vary across atlases; thus, the atlas used in each study affected their findings (Supplementary Fig. 13b). Second, localization of projection origins in specific thalamic nuclei can vary both as a result of the atlas used and the ability to precisely target individual nuclei, as demonstrated by the discrepancies reported in projections from CM (Supplementary Fig. 13a). To avoid anatomical bias, we averaged nucleus localization data between the two atlases (ABA and PMBA; Supplementary Fig. 13c) and created our confidence maps independent of nuclear boundaries (Fig. 3).

Figure 6 Nuclear localization of the thalamic origins of frontal projections. (a–c) Three representations of the nucleus origin data for LO (Supplementary Fig. 12). (a) The fractions of each thalamic nucleus projecting to LO are shown for confidence levels 3, 5 and 7, respectively (dotted lines, see Supplementary Fig. 13d), with their average (black line). The vertical gray line represents the inflection point (white) at 15%. Asterisks indicate potential differences between localized thalamocortical projection origins and literature data in rat (Supplementary Fig. 13a). (b) Single coronal section through the confidence map for LO (gray scale) overlaid with nuclear subdivisions from the ABA. The atlas is colored on the left to indicate the fraction of each nucleus covered by the average confidence trace (black line in a), with the inflection point (white) at 15%. (c) Spatial representation of all nuclei projecting to the LO. Circle diameters correspond to the relative size of each nucleus and their positions correspond to their relative center-of-mass location in the thalamus in the anterior-posterior and medial-lateral axes. Color scale is the same as in b. (d) The fractions of each thalamic nucleus projecting to AI, IL and FrA, shown for confidence levels 3, 5 and 7, respectively (dotted lines, see Supplementary Fig. 13d), with their average (black line), as described in a. (e) Aggregate nucleus coverage map for all cortical areas. Nuclei (rows) and cortical subregions (columns) are hierarchically clustered on the basis of output and input similarity, respectively. Color scale is the same as b.
Thalamic origins of layer-preferential projections in vM1

Different layers of the same cortical area have distinct roles in information integration. We analyzed the primary vibrissal motor cortex (vM1) to test whether our data set could be used to identify thalamic volumes preferentially innervating specific cortical layers. Previously, we found that the posterior ‘sensory’ thalamus is more likely to project to layers 2/3 and 5a (L2/3–5a) in vM1, whereas the anterior ‘motor’ thalamus projects to layer 5b (L5b) as well as L2/3–5a. However, we had to estimate the thalamic volumes responsible for these projections on the basis of subjective assessments of a small number of injections.

To accurately localize the thalamic origin of layer-specific projections to vM1, we quantified the fluorescence intensity of thalamic projections on the basis of subjective assessments of a small number of injections. We found that several clusters displayed strong preference to specific vM1 layers. For example, 81% of cluster 11 preferentially projected to L2/3–5a in vM1. An arbitrary 10% threshold is indicated (gray line). (g) Schematic showing layer preferential input from thalamus to vM1 in the context of a motor-sensory circuit diagram. Scale bars represent 1 mm.

DISCUSSION

Mesoscopic connectivity maps are crucial for studying interactions among multiple brain regions and for linking cellular circuit mechanisms to behaviors. In this study, by using anterograde viral tracing, high-throughput whole-brain imaging, and custom development of alignment and analysis software, we established a mesoscopic thalamo-centric projection map to the cortex in mouse, identified unique thalamic sub-volumes projecting to each cortical subregion, and determined the optimal injection coordinates for optogenetically
or anatomically targeting specific cortical regions. Our maps also permitted the identification of shared and unique thalamic sources to different cortical regions, such as PrL and II (Fig. 3f), providing an entry point for teasing out their common and distinct functions. In addition, our systematic approach allowed us to functionally subdivide the thalamus solely on the basis of cortical projection patterns (Fig. 4). We further identified the thalamic volumes that give rise to layer-preferential projections to vM1 (Fig. 7). Our results provide a foundation for understanding the function of the thalamic and frontal cortex, as well as for investigating and manipulating the microcircuits in and between thalamic and cortical subregions.

Historically, the extensive time and labor required to image and map long-range projections has limited the number of tracer injections used in anatomical studies and necessitated the reliance on subjective assessments to compare across experiments. Recent advances in high-throughput fluorescent imaging facilitate the generation of large anatomical image data sets22,38-40, allowing researchers to access vast amounts of anatomical information. However, extracting relevant biological information from these data remains a major challenge for several reasons: variability across experiments, both as a result of intrinsic size differences and experimental manipulation, makes it difficult to compare across experiments directly; the resolution is limited to the size and shape of the tracer injection site, and the tools needed for data analysis have not kept up with technological advances in data collection, impeding efforts to turn images into knowledge.

We addressed variability issues by tightly controlling the animal age (P30 ± 2), computationally correcting for angled sectioning and normalizing individual thalami to a standard volume. The variability among our thalamic mask boundaries is 102 ± 51 µm, comparable with that observed in the absence of mechanical sectioning24. By creating a comprehensive, age-matched thalamocortical projection map, we have provided a framework that others can build on to understand differences across age groups, cell types and species. These variances may explain some of the discrepancies seen across the 43 anatomical studies we evaluated in rat and our data in mouse (Supplementary Fig. 13).

Another major limitation of mesoscopic mapping is that the size of the tracer injection limits the resolution. To reliably identify the origin of the each mapped projection, tracer injections must target a single defined brain region. This is straightforward in cortical areas with large, superficial subregions39,40; however, this task is difficult, if not impossible, in the thalamus as a result of the complex shape and small size of many thalamic nuclei. We overcame this limitation by analyzing the intersectional areas of overlapping injections (Fig. 3b,c), which allowed us to localize volumes smaller than a single injection. We could only have obtained our confidence maps and optimal injection coordinates by integrating information from a large number of highly overlapping injections. Furthermore, to maximize our resolution, we used the smallest replicable viral infection volume (~0.3 mm^2, and laterally ~600 µm). We estimate our resolution to be larger than our variability (~100 µm) and smaller than our injection size (~600 µm), which is sufficient for most thalamic targeting, but small thalamic nuclei may require smaller injection volumes or more closely spaced injections to precisely discriminate their boundaries (Supplementary Fig. 10). Because of the heavy dependence on viral infection to deliver molecular reagents in systems neuroscience, our map, which is at the equivalent ‘operational scale’, can serve as a guide for targeting these tools.

By exploiting injections that both do and do not project to each cortical ROI, we were able to identify the entire thalamic volume that does not project to each cortical ROI. From an anatomical point of view, characterization of non-projecting regions is particularly important because it has been estimated that only ~10% of all possible connections in the rodent brain are fully characterized at the mesoscopic level, largely as a result of a lack of definitive information on non-existent projections1.

As stated previously5, “The concept of the thalamic nucleus as a single structural, functional and connectional entity has barely survived advancing techniques and new information. We stay with the thalamic nuclei as one of our prime analytical tools because, as yet, we have little to use in its place.” Here, our comprehensive projection map provided us with a unique opportunity to establish a nucleus-independent map of thalamic projections that transcends what we have learned from a nucleus-based framework (Figs. 3–5). Although we related our results to thalamic nuclei, we created our confidence maps independent of nuclear boundaries. This enabled us to unbiasedly identify the precise thalamic volumes responsible for projections to specific cortical subregions and cortical layers.

Our maps were obtained in adolescent mice, which is a dynamic period for prefrontal cortex (PFC) associated behaviors31-43. We found that thalamocortical projections from at least 25 nuclei reached PFC and formed functional synapses by P30 (Figs. 5 and 6). Given that the frontal subregions innervated by each nucleus were comparable to those seen in the adult rat (Supplementary Fig. 13), our data suggest that thalamocortical projections to PFC reach their final targets by P30 in mouse. We therefore propose that the behavioral changes that occur during adolescence are more likely a result of local refinements and synaptic pruning than larger rearrangements in thalamocortical projection distributions to PFC subregions.

In light of new tools for imaging, physiology and cell type-specific manipulations in mouse4, our mesoscopic data will serve as a critical reference for applying these tools to study circuit function. The results from over 43 disparate studies were necessary to summarize only a fraction of the thalamocortical projections in rat that are described here in mouse (Supplementary Fig. 13), which is a testament to the power of the high-throughput imaging and computational analyses that we used. The ability to directly compare across animals and experiments is a crucial step for extracting useful biological information from large anatomical data sets. Our results present an example for large-scale data integration and analysis, and will inform future studies in systems neurobiology.

METHODS

Methods and any associated references are available in the online version of the paper.

Note: Any Supplementary Information and Source Data files are available in the online version of the paper.

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AUTHOR CONTRIBUTIONS

All of the authors participated in designing the experiments. B.I.H., D.K., K.I.G. and T.M. performed the experiments. B.I.H., B.R.L., K.I.G., D.K., H.Z. and T.M. analyzed the data. B.I.H., B.R.L., K.I.G., H.Z. and T.M. wrote the manuscript.
COMPEITING FINANCIAL INTERESTS
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ONLINE METHODS

All animal experiments were conducted according to US National Institutes of Health guidelines for animal research and were approved by the Institutional Animal Care and Use Committee of the Oregon Health and Science University. All measurements are listed as mean ± s.d. unless otherwise indicated. All calculations were performed in MATLAB (MathWorks). The raw data and the analyzed data are publicly available at http://digitalcollections.ohsu.edu/projectionmap. The original resolution images are available upon request for hard drive format.

Stereotaxic viral injections. Injections were performed as described with optimizations and modifications. Briefly, C57BL/6 male and female mice were anesthetized (1–2% isoflurane, vol/vol) at P14–18 and stabilized in a custom stereotaxic apparatus (modified from a David Kopf system). A dental drill (Henry-Schein) was used to drill holes through the skull. A pulled glass micropipette (Drummond, tip diameter = 10–15 μm), beveled sharp, was back-filled with AAV (serotype 2/1) that expresses either eGFP (Addgene 28014) or tdTomato (a gift from J. Magee, Janelia Farm Research Campus). AAV2/1 is a hybrid serotype that has AAV2 inverted terminal repeats, AAV1 capsid proteins and widespread neuronal tropism. The transgenes were driven by CAG promoter and included a WPRE element to enhance the expression. The viruses were prepared by the University of Pennsylvania vector core and viral titers >5.0 × 10^12 GC per ml were used. Unless noted otherwise, a 10-μl volume of virus was dispensed at a speed of 5 nl s−1 using a hydraulic injector (Narishige), followed by a 5–10-min waiting period. The pipette was retracted 0.3 mm at 0.008 mm s −1, paused for 3 min, and then retracted at a rate of 0.008 mm s−1. This process minimized the undesirable infection of cells along the injection path. Up to four injections were performed in each animal (two colors and two hemispheres). Coordinates for injections ranged from: 0.5 to −1.6 anterior to posterior, 0–1.6 lateral, and 2.8–4.2 deep from the pia (in mm from bregma). Although no statistical methods were used to pre-determine sample sizes, we sought to insure that the final coverage of all thalamic labeling was >90%. We found that this could be achieved via 254 highly overlapping injections across 75 animals.

Sectioning and imaging. 14 d after viral infection, mice were perfused transcardially with 25 ml phosphate-buffered saline (PBS) followed by 50 ml of 4% paraformaldehyde (PFA, wt/vol). The brain was post-fixed in 4% PFA at 4 °C overnight and then placed in 30% sucrose (wt/vol) in PBS at 4 °C overnight. The brain was centered and aligned in a rectangular mold, embedded in Optimal Cutting Temperature medium (Tissue-Tek), and sectioned coronally on a cryostat (Thermo Scientific) at 50-μm thickness. The sections from the most anterior section of the cortex to the most posterior section of the thalamus were floated in PBS and then collected onto Superfrost-Plus microscope slides (FisherBrand). Slides were mounted using Fluoromount (Sigma) and covered with number 1.5 cover glass (Gold Seal, Fisher).

All sections on the slides were imaged with a 20x objective (0.5 μm per pixel) on the Nanozoomer slide scanner (Hamamatsu), at a fixed exposure time. Because injection sites were often overexposed under these settings, they were re-imaged at a lower exposure with either a 5x objective on a Zeiss Axio Imager or using shorter exposure times on the Nanozoomer. Axio images were matched to their corresponding Nanozoomer section images through rigid translation and rotation using manually selected anatomical landmarks visible in both images. After imaging, injections that extended beyond the lateral or ventral borders of the thalamus were excluded. Each brain was processed and imaged equivalently and randomly without any knowledge of the injection locations.

Cell counting. Confocal images were collected (Zeiss, LSM780) for DAPI (Vector Labs, catalog number H-1200) stained sections across the center of an AAV2/1-eGFP thalamic injection site from 17 mice. The fraction of cells found to be both DAPI and eGFP positive indicated the percentage of DAPI-positive cells infected. The transgenic infected, thalamic sections across the center of AAV2/1-eGFP injections from five mice were incubated with mouse antibody to NeuN (Millipore, catalog number MAB377, dilution 1:1,000), followed by Alexa-594 goat antibody to mouse (Life Technologies, catalog number A-11005, dilution 1:1,000) and DAPI. The fraction of DAPI-positive cells that were also NeuN- and eGFP-positive indicated the percentage of infected neurons at the injection site (Supplementary Fig. 1b–e). To confirm these results, the thalamic sections from three mice injected with eGFP-expressing AAV2/1 were stained with NeuroTrace (Life Technologies, catalog number N-21482, dilution 1:100) and DAPI and were analyzed in the same fashion (data not shown). To evaluate the viral tropism, eGFP and tdTomato expressing AAV were mixed (1:1) and co-injected into the thalamus in four mice. The same imaging process was used as with single viral injections (Supplementary Fig. 1f–h).

Thalamic and injection site segmentation. Individual sections were isolated from the full slide images by determining an intensity threshold that would distinguish tissue from background pixels. The outline of the thalamus was manually traced to generate a thalamus mask (Fig. 1c). The front of the thalamus was defined as the first slice posterior to the anterior commissure crossing the midline, and the back of the thalamus was defined as one slice posterior to the end of the lateral geniculate nucleus. In addition, the medial and lateral geniculate nuclei were not included due to their already well-characterized anatomy in the auditory and visual systems, respectively. Finally, the posterior portion of the reticular thalamic nucleus, which does not produce cortical projections and the posterior portion of the ventral medial nucleus were excluded from the traced masks due to technical difficulties in visualizing their borders. We segmented each injection site into a binary mask by applying independent intensity thresholds in green and red channels, using a supervised MATLAB routine based on Otsu’s method. Traveling axon bundles that were above threshold in the thalamus were manually excluded from the associated injection site.

Thalamic registration and alignment. The model thalamus and registered injections were created as described in Supplementary Table 1. First, two manually selected midline points were used to rotate and align the thalamus masks. Second, to align the masks in the correct y position and to correct for the cutting angle tilt about the x axis (that is, rotation around the x axis), we used anatomical landmarks to estimate the tilt angle (Supplementary Fig. 5). A separately traced ABA thalamus mask was rotated to the same tilt angle and the mask stack was resampled as 50 μm slices. Third, the centers of mass of these slices were used to direct the projection of experimental thalamus masks in y. The center of mass is defined as the unique point where the weighted relative position of the distributed mass sums to zero and was calculated as

\[ R = \frac{1}{M} \int \rho(r) r dV \]

where \( M \) is the sum of the masses of each point \( r \) in a volume \( V \) with constant density \( \rho(r) \).

The aligned thalamus masks were then rotated to a tilt angle of 0 degrees and resampled as 50 μm slices (Supplementary Fig. 5). The thalamus masks were downsampled to a 36.4 × 36.4 × 50 μm voxel size. Fourth, to control for the cutting angle tilt about the y axis, the aligned three-dimensional mask was sheared to maximize left and right asymmetry. Fifth, the overall size of the thalamus was scaled in z so that the midline distance from the beginning to the end of the thalamus matched the ABA thalamus, in x-y isotropically to match the total area of the central slices with that of the corresponding ABA thalamus, and the three-dimensional thalamus masks were scaled in z to match the average width of all thalamic masks. Finally, the masks were visually inspected and 18 of 75 brains underwent minor scaling or position adjustments in the z dimension. All brains were further aligned with each other in y based on their center of mass. All experimental masks were summed and then segmented according to a threshold that retains the volume of averaged thalamic mask volumes, producing the model thalamus.

We used Dice’s coefficient to assess the similarity between two thalamic structures (Fig. 2a and Supplementary Fig. 6d). Dice’s coefficient is defined as

\[ D(A, B) = \frac{2 \sum (A & B)}{\sum A + \sum B} \]

where \( A \) and \( B \) are two binary volumes, ‘&’ is the logical AND operator and \( \Sigma \) indicates the sum of all elements in \( X \). To further quantify the variability of thalamic masks (Fig. 2a and Supplementary Fig. 6), we overlaid the borders of each thalamus mask and measured the distribution of boundary points at 18 locations (6 locations per slice for 3 z slices).
Injection site masks were processed identically to their corresponding thalamic masks so that they are registered to the model thalamus. All injection site masks were summed to quantify the injection coverage at individual voxels (Fig. 2d–f). Data distribution was assumed to be normal, but this was not formally tested.

Atlas alignment. To register known thalamic nuclei within our model thalamus, we traced, scaled and re-sampled 25 nuclei from both the ABA and PMBA as three-dimensional volumes that are aligned with our model thalamus. Differences in animal age and tissue preservation techniques resulted in size and shape differences between the two atlases, so each atlas was scaled separately to best fit our model thalamus. The correspondence between the nuclei of individual experimental thalami and the atlas nuclei were assessed by manually tracing four cytoarchitecturally identifiable thalamic structures (nuclei AD, AV and PT and fiber tract fr) from five randomly selected brains (Fig. 2b).

The similarity between the atlas and experimental nuclei was assessed using Dice’s coefficient (Fig. 2c and Supplementary Fig. 6e). Notably, the values of Dice’s coefficients for comparing nuclei in were lower than those for comparing the thalami of all brains (Supplementary Fig. 6d,e) because this coefficient is inversely dependent on volume. For example, the average volume of the traced nuclei is 0.24 mm$^3$ (1.3% of the model thalamus volume), and our position variability (~100 nuclei) and similarities between traced nuclei and atlas nuclei (average $D = 0.53$ for comparing the same nuclei and $D = 0.02$ for comparing different nuclei) and similarities between traced nuclei and atlas nuclei ($D = 0.53 \pm 0.10$) are comparable to those of the atlases to each other ($D = 0.60 \pm 0.11$) (Fig. 2c and Supplementary Fig. 6e).

Confidence maps and thalamic origins of projections to the cortical sub-regions. For each thalamic injection, projection distributions were blindly scored by three independent experts. The presence/absence, strength (dense or sparse), coverage (full or partial ROI coverage) and specificity (whether the projection also goes to an adjacent ROI) were determined (Supplementary Fig. 8b). All final scoring decisions were reached by consensus. The cortical area boundaries were based on the PMBA. Injections are referred to as being positive or negative for a given cortical ROI, where positive indicates the presence of a projection and negative indicates the absence of a projection for this particular ROI. To control for our alignment variability (~100 µm) across thalamic masks (Figs. 2a and 3b), an injection core was produced by eroding each three-dimensional injection mask by 100 µm.

A confidence map, which defines the thalamic origin of cortical projections, was created for each target projection region. As shown in Supplementary Figure 8, a confidence map was developed by grouping injections (Fig. 3b) that met each of eight independent criteria. Meeting each criterion would give a thalamic voxel a score of 1 and meeting all criteria would result in a maximal confidence level of 8. For example, criteria (A) requires a voxel to be included in the core of an injection producing specific projections, but may not be in any negative injections (see Supplementary Fig. 8c for the remaining seven criteria descriptions). The binary masks produced by each grouping criteria were summed to create the confidence map (Supplementary Fig. 8d–g).

Overall, the confidence maps incorporate information about the intensity and specificity of projections, as well as the variability in thalamic transformation and alignment. The confidence map therefore represents the likelihood of a thalamic voxel projecting to a particular target.

Voxel clustering based on projection confidence maps. The model thalamus and individual confidence maps were down sampled to 150 x 150 x 150 µm voxels. The thalamic voxels were then subjected to agglomerative hierarchical clustering (MATLAB) based on their confidence map values across the 19 target regions in a 19-dimensional space using the city-block metric and average linkage with a set maximum of 11 clusters.

Quantifying the nuclear origins of thalamocortical projections. Atlas nuclei previously aligned to the model thalamus were overlaid onto our confidence maps. We calculated the fraction of injection-covered nucleus volume occupied by the confidence map at three confidence levels ($C \geq 3$, $C \geq 5$ and $C \geq 7$; Supplementary Fig. 8d). These values were averaged across the ABA and PMBA atlases to create the confidence threshold data (Fig. 6a,d and Supplementary Fig. 12a). The average nuclear fraction from these three thresholds (Fig. 6a,d and Supplementary Fig. 12a) gives the final nucleus projection data that forms the basis of our visualization and clustering results (Fig. 6e).

Clustering nuclei and projection regions. Each nucleus was assigned a point in a 19-dimensional space corresponding to the fraction of the thalamic nucleus volume occupied by projections to each cortical area (Fig. 6e). We performed a cluster analysis on the nuclei using a Euclidean distance metric and minimum linkage. The projection regions were similarly assigned a point in 25-dimensional space corresponding to the 25 nuclei, and clustered using the same method.

vM1 injection and projection analysis. The boundaries of vM1 were based on previous characterizations and were defined as follows: dorsally by the pial surface, medially by a line that connects the top arc of the cingulum to the point that the pia folds toward the midline, and laterally by a line from the cingulum to the pia that is parallel to the midline (Fig. 7a). vM1 was delineated independently for each hemisphere from three sections: the section where the corpus callosum merges plus one section anterior and one posterior.

Because layer depth and thickness varies depending on the position within vM1, we normalized all depths to that at the medial edge of vM1 (Supplementary Fig. 14). Specifically, for pixels at angle $\theta$, their depths are linearly transformed to the medial depth based on the layer boundaries at $\theta$ and at the medial boundary (Supplementary Fig. 14a). The normalized depths were separated into 100 bins, and fluorescence intensity values in each bin were averaged and normalized to the background fluorescence estimated from an unlabeled cortical region on the same section. These normalized fluorescence intensity traces were further background subtracted using the minimum values of the respective traces lying in the vicinity of cell body layer at the L1–L2/3 boundary.

A thalamic injection was considered to produce layer-preferential projections if it met two criteria. First, the average fluorescence intensity within either L2/3–5a or L5b had to be significantly greater than background fluorescence measured from the same depths in vM1 brains that did not contain projections to vM1. The threshold for each depth and each color was the median plus interquartile range of the background fluorescence levels in brain sections not containing vM1 projections. If ≥25% of the bins in either the L5b or L2/3–5a region were considered above threshold, then the second criteria would be evaluated. Second, after subtracting the layer specific background fluorescence, the intensity was averaged in L2/3–5a and L5b and a layer preference index, $\alpha$, was computed from these average intensities in L2/3–5a and L5b ($I_{L2/3-5a}$ and $I_{L5b}$, respectively)

$$\alpha = \log_2 \frac{I_{L2/3-5a}}{I_{L5b}}$$

$\alpha = 0$ indicates equal intensities in the two regions, whereas $\alpha > 0$ indicates higher fluorescence intensity in L2/3–5a and $\alpha < 0$ indicates higher fluorescence intensity in L5b. A threshold was set at 1.1: if $\alpha$ was greater than 1.1, the injection was classified as strongly L2/3–5a preferential, and if $\alpha$ was less than –1.1, the injection was classified as L5b preferential. Based on this classification, we created vM1 layer-preferential thalamus confidence maps by scoring each voxel against the following four criteria (each criterion gives a score of 1 and meeting all criteria gives a maximal confidence level of 4): 1) the voxel is in a layer preferential injection, 2) the voxel is in the core of a layer preferential injection, 3) the voxel is in the core of a layer preferential injection with strong intensity, and 4) the voxel is not in the core of an injection with opposite layer preference (Fig. 7).

Photostimulation and electrophysiology. Mice were injected at P14–16 with 20–40 nl of an AAV2/1 virus encoding ChR2-H134R-TaTomato (Addgene, 28017). Cortical brain slices were prepared 14 d later from mice anesthetized with an intraperitoneal injection of ketamine/xylazine (0.13 mg ketamine/0.01 mg xylazine per g of body weight) and perfused transcardially with ice cold artificial cerebrospinal fluid (ACSF) containing 127 mM NaCl, 25 mM NaHCO3, 25 mM NaH2PO4, 1 mM MgCl2, 2.5 mM KCl, 1 mM MgCl2, 2 mM CaCl2, and 1.25 mM NaH2PO4, pH 7.25–7.35, –310 mOsm, and bubbled with 95% O2 / 5% CO2. The brain was removed and placed into ice-cold cutting solution containing 110 mM choline chloride, 25 mM NaHCO3, 25 mM NaH2PO4, 13.5 mM sodium ascorbate, 7 mM MgCl2,
3 mM sodium pyruvate, 2.5 mM KCl, 1.25 mM NaH2PO4, and 0.5 mM CaCl2. 300 µm thick modified coronal slices were vibratome sectioned (Leica 1200S) at an angle to achieve a cut perpendicular to the pial surface for each recorded brain area. Slices were incubated in oxygenated ACSF for 45 min at 34 °C, and then maintained in an oxygenated holding chamber at 22–24 °C.

Subcellular channelrhodopsin-assisted circuit mapping (sCRACM) and electrophysiology were performed as previously described16,46. The excitatory postsynaptic currents (EPSCsCRACM) were recorded in voltage clamp (holding potentials were −70 mV or −75 mV) while blue light stimulated the thalamic axons transfected with channelrhodopsin. Each map was repeated 2–4 times. After sCRACM maps were obtained, a cell was counted as a positive responder if there was any excitatory postsynaptic current amplitude >6× the s.d. of the baseline.

Retrograde bead injections, imaging and analysis. Retrograde tracing was performed using fluorescent latex microspheres (LumaFluor: Red Microbeads IX and Green Microbeads IX) at a 1:2 dilution in PBS. Injections were performed similarly to the viral injections with P27 mice (tip diameter = 40–60 µm). 3 d later, mice were perfused as described above, with the exception that brains were not post-fixed following perfusion. Brains were sectioned coronally on a vibratome (Leica BT1200S) at 100-µm thickness. Sections were floated, collected, mounted and covered as described above. All sections on the slides were imaged (Olympus MVX10), at a fixed exposure time, using a Retiga 2000R camera. From these images, the cortical injection sites as well as the approximate distribution of fluorescent thalamic somas were manually mapped onto thalamic sections (Supplementary Fig. 10).

Statistics. Statistical comparisons were performed using a Student’s t test, where n indicates the number of independent brains. The significance level was set at P = 0.05.

A Supplementary Methods Checklist is available.

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