Time course of homeostatic structural plasticity in response to optogenetic stimulation in mouse anterior cingulate cortex

Abbreviated title: Time course of homeostatic structural plasticity

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Abstract

Plasticity is the mechanistic basis of development, aging, learning and memory, both in healthy and pathological brains. Structural plasticity is rarely accounted for in computational network models, due to a lack of insight into the underlying neuronal mechanisms and processes. Little is known about how the rewiring of networks is dynamically regulated. To inform such models, we characterized the time course of neural activity, the expression of synaptic proteins, and neural morphology employing an *in vivo* optogenetic mouse model. We stimulated pyramidal neurons in the anterior cingulate cortex of mice and harvested their brains at 1.5 h, 24 h, and 48 h after stimulation. Stimulus-induced cortical hyperactivity persisted up to 1.5 h and decayed to baseline after 24 h, indicated by c-Fos expression. The synaptic proteins VGLUT1 and PSD-95, in contrast, were upregulated at 24 h and downregulated at 48 h, respectively. Spine density and spine head volume were also increased at 24 h and decreased at 48 h. This specific sequence of events reflects a continuous joint evolution of activity and connectivity that is characteristic of the model of homeostatic structural plasticity. In computer simulations, we thus showed that this model was able to corroborate the observed empirical evidence in our animal experiments.

Introduction

Neural circuits in the mammalian brain are highly plastic (Holtmaat and Svoboda 2009). **Synaptic plasticity comes in two different flavors.** Functional plasticity means that chemical synapses change their strength by modifying signal transmission based on neurotransmitters and receptors (Bear and Malenka 1994; Malenka and Bear 2004). Structural plasticity, in contrast, refers to a variety of changes including the branching of dendrites, the geometry of dendritic spines, and number of dendritic spines and axonal boutons, and the connectivity between specific pairs of neurons (Trachtenberg et al. 2002; Caroni et al. 2012; Pfeiffer et al. 2018). Both forms of plasticity are underlying network assembly during development, use-dependent adaptation and learning in the adolescent and adult brain, but also network decay during aging and disease (Lamprecht and LeDoux 2004). Memory depends on plasticity. For instance, fear conditioning has been shown to increase both the synaptic strength and connection probability among a subgroup of granule cells in the dentate gyrus (Ryan et al. 2015). The resulting memory engram encodes a distinct episodic memory (Liu et al. 2012; Ramirez et al. 2013). Plasticity caused by injury, such as synaptic potentiation and network remodeling triggered by stroke or brain lesion, is likely to involve both activity perturbation and neuroinflammation (Keck et al. 2008; Murphy and Corbett 2009). In brain diseases, pathological plasticity may affect several brain regions. Acute and chronic stress, for instance, has been shown to induce different functional and structural alterations in the hippocampus, anterior cingulate cortex (ACC), amygdala (Lucassen et al. 2014), and elsewhere. Given this wealth of phenomena, the question arises how functional and structural plasticity is regulated.

The rules underlying experience-dependent plasticity need to be investigated further. Experiments in different brain regions with different plasticity-inducing paradigms have given rise to a host of different phenomena (Bear and Malenka 1994; Malenka and Bear 2004; Keck et al. 2017; Gainey and Feldman 2017). Correlation-based Hebbian plasticity, summarized as “neurons that fire together wire together” (Hebb 1949), was proposed to account for homosynaptic strengthening observed in animals minutes to hours after arti-

ficial high-frequency stimulation (Lowel and Singer 1992). Despite its great potential in explaining learning and memory, Hebbian plasticity in computational network models was shown to increase the risk of excessive excitation or silencing, respectively (Sejnowski 1977; Miller and MacKay 1994). The same lack of network-level stability is implied by spike timing dependent plasticity (STDP), which implements either homosynaptic strengthening or weakening based on the relative timing of presynaptic and postsynaptic spikes (Markram et al. 1997; Bi and Poo 1998). The discovery of heterosynaptic plasticity and synaptic scaling, however, hinted that the modulation of a synapse may also depend on its neighbors (Lynch et al. 1977; Chater and Goda 2020) and the activity of the postsynaptic neuron (Turrigiano 2012). Chronic *in vivo* recordings have indeed revealed a robust cell-by-cell firing rate homeostasis across days and weeks (Hengen et al. 2016; Ma et al. 2019; Pacheco et al. 2019). New models of homeostatic plasticity (Turrigiano 2012, 2017), possibly in combination with Hebbian plasticity rules, are now being evaluated for their ability to solve the aforementioned network stability issues. Preliminary conclusions posit that the time scales of homeostatic control should be much faster than those observed in experiments (Zenke and Gerstner 2017). Rarely, however, was a possible role of structural plasticity explored in these theoretical studies.

Structural and functional plasticity do not represent independent processes. Changes in spine number and individual spine head volumes were observed after synaptic potentiation or depression *in vitro* (Engert and Bonhoeffer 1999; Matsuzaki et al. 2004; Zhou et al. 2004). Moreover, both spine density and synaptic strength were shown to compensate for input loss caused by entorhinal denervation in organotypic slice culture (Vlachos et al. 2012b; Lenz et al. 2019). In parallel, theoreticians began to reflect over possible functional aspects of structural plasticity in a network (Fauth and Tetzlaff 2016). The homeostatic structural plasticity (HSP) model seems particularly promising in reconciling robust development and associative learning (Butz et al. 2009; Van Ooyen 2011; Butz and van Ooyen 2013; Gallinaro and Rotter 2018; Lu et al. 2019; Gallinaro et al. 2020). Still, the empirical data justifying such activity-dependent structural plasticity models are sparse. Most studies report changes in spine density after massive manipulation of activity, or in

brain diseases, see review by Chidambaram et al. (2019). Unfortunately, time-resolved neural activity and connectivity was not included in any of them. Yusifov et al. (2021) revealed the elaborate temporal dynamics of spine density during monocular deprivation but only apical dendrites were monitored. The time course of structural changes while the neuronal activity recovers, however, is of great importance to disambiguate structural plasticity models.

The optogenetic stimulation paradigm employed in the present study evolved from our previous studies. We activated the pyramidal neurons in mice ACC for four consecutive days (30 minutes per day) until they developed depressive-like behaviors. Using the same optogenetic stimulation paradigm, we sampled mouse brains at 1.5 h, 24 h and 48 h after chronic stimulation. We stained and quantified the relative abundance of neuronal activity marker c-Fos, and general synaptic markers the vesicular glutamate transporter 1 (VGLUT1) and the postsynaptic density scaffold protein PSD-95 (De Gois et al. 2005; Ehrlich et al. 2007). As expected, optogenetic stimulation triggered depressive-like behaviors and hyperactivity in mice (Barthas et al. 2015). Hyperactivity of pyramidal neurons, evidenced by a robust c-Fos expression at 1.5 h, eventually diminished to baseline 24 h after the end of the stimulation, while VGLUT1 and PSD-95 showed strong delayed up-regulation at 24 h and again downregulated after 48 h in the stimulated mice. Similar to the temporal expression profile of VGLUT1 and PSD-95, dendritic spine density and spine head volume of the stimulated mice were increased at 24 h and restored to the control levels, or even slightly below control at 48 h. Microglia and astrocytes are known to be involved in structural plasticity (Weinhard et al. 2018) and glutamate homeostasis (Schousboe and Waagepetersen 2005). Therefore, we also checked whether glial markers, GFAP and IBA1, were overexpressed throughout 48 h after stimulation. We argue that, compared to other candidate theories, the homeostatic structural plasticity model could explain the biphasic changes of synaptic proteins and dendritic morphology consistently at 24 h and 48 h after the chronic stimulation.

Materials and Methods

Animals

3–5 months old genetically modified mice expressing ChR2 and yellow fluorescent protein (Thy1-ChR2-YFP) in a subset of pyramidal neurons (MGI Cat# 3719993, RRID:MGI:3719993) as well as C57BL/6J male adult mice (IMSR Cat#JAX:000664, RRID:IMSR_JAX:000664; Charles River, L’Arbresle, France) were used in the current study. All mice were kept in a reversed day-night cycle, with food and water provided *ad-libitum*. Mice were firstly group caged and then single housed after the optic fiber implantation. The Chronobiotron animal facilities are registered for animal experimentation (Agreement A67-2018-38), and protocols were approved by the local ethical committee of the University of Strasbourg (CREMEAS, n° 02015021314412082).

Animal experimental design

The animal experiments’ objective was to determine the time course of plastic phenomenon triggered by external stimulation. We adopted an established optogenetics mouse model from our laboratory, in which the pyramidal neurons in ACC (24a/24b) were activated for four consecutive days (details see Optogenetic stimulation section below). We studied the temporal dynamics with discrete time points, by harvesting the mouse brain tissue at 1.5 h, 24 h, or 48 h after the last stimulation. As shown in our previous studies (Barthas et al. 2015, 2017), sustained stimulation of the ACC induces depressive-like behavior in naïve mice. So in the current study, we used splash test and novelty-suppressed feeding (NSF) test to verify the behavioral effects of the optogenetic stimulation. In the 24 h-post groups, we conducted splash test on the fifth day, while in the 48 h-post groups, we performed both NSF test and splash test on the fifth and sixth day and sacrificed the mice afterwards. We later evaluated the temporal evolution of neural activity by quantifying the expression of c-Fos at the three aforementioned time points. To capture when and where synaptic alterations may occur, the expression of pre- and postsynaptic proteins

were evaluated. Following the pattern shown by preliminary molecular screening, we examined if structural changes accompany molecular alterations by estimating the spine morphology of ACC pyramidal neurons harvested at 24 h and 48 h post-stimulation. To further confirm the involvement of glial cells, the expression of glial markers at 1.5 h, 24 h, and 48 h were respectively inspected. Before we performed all the experiments in Thy1-ChR2-YFP mice, we compared the efficacy of the transgenic approach with viral transfection. For the latter, we injected bilaterally AAV-CaMKII-ChR2 (H134R)-EYFP (Addgene plasmid #26969; <http://n2t.net/addgene:26969>; RRID: Addgene_26969) into the ACC (details see Virus injection section in the Supplementary Materials) of C57BL/6J mice. As we observed no differences in the c-Fos activity and behavioral outcomes in two approaches after optogenetic activation, we decided to perform all the experiments in transgenic mice to reduce the number of surgery that animals go through. All mice group information was summarized in Supplementary Table 1.

Stereotactic surgery

Stereotactic surgery was conducted to inject virus and implant optic fiber into ACC. During the surgery, mice were deeply anesthetized with a mixture of zoletil (25 mg/kg tiletamine and 25 mg/kg zolazepam) and xylazine (10 mg/kg) (Centravet, Taden, France; i.p. injection) and locally anesthetized by bupivacaine (Mylan, The Netherlands; 0.5 mg/mL; subcutaneous injection, 1 mg/kg). The coordinates of the injection/implantation site are +0.7 mm from bregma, lateral: ± 0.3 mm, dorsoventral: -1.5 mm from the skull (Barthas et al. 2015; Sellmeijer et al. 2018).

Optic fiber implantation

We inserted 1.7 mm-long LED optic fiber (MFC_220/250-0.66_1.7mm_RM3_FLT, Doric Lenses, Canada) unilaterally (left or right) in C57BL/6J mice two weeks after the virus injection or directly in naïve Thy1-ChR2-YFP mice. The fiber was inserted into ACC for 1.5 mm deep with reference to the skull. The metal end was fixed onto the skull by

superglue and dental cement, and then the skin was stitched. For stimulation, we used blue light (460 nm wavelength) and the light intensity of optic fibers used in the current study ranged from 1.7 mV/mm² to 6 mV/mm².

Optogenetic stimulation

After the optic fiber implantation, we individually housed the mice to avoid possible damage to the implant. After seven days of recovery, we started the optogenetic stimulation protocol on freely moving mice in their home cages. Optogenetic stimulation took place on four consecutive days. We applied repetitive pulses for a total duration of 30 minutes, in which the pulses were organized into multiple 10 seconds long pulse trains, each consisting of 8 seconds at 20 Hz with 40 ms pulse duration and 2 seconds without stimulation. We did not observe the effects of light on the behaviors in gene-matched wild type mice (Supplementary Figure 1-2). We used transgenic mice for all experiments and kept the light off for the sham groups. At the end of the stimulation, all mice were handled again and unplugged from the cable.

Behavioral tests

We performed all the behavioral tests during the dark phase under red light. Splash test (Nollet et al. 2013) and novelty-suppressed feeding (NSF) test (Samuels and Hen 2011) were used to evaluate depressive-like behaviors. In the splash test, we sprayed 15% sucrose solution onto the coat of the mice and recorded the total grooming time for each mouse during the following 5 min. The NSF test was conducted on a different day of splash test and we removed the food pellets 24 h before testing. During the test, we put each mouse into an open field, where a food pellet was placed in the middle, and recorded the time delay necessary for each mouse to touch and eat the pellet (within 5 min).

Verification of injection site and tissue harvesting

Mice were perfused with cold 4% paraformaldehyde (PFA) in $1\times$ phosphate buffer (PB, 0.1 M, pH 7.4), under Euthasol Vet (intraperitoneal injection, $2\mu\text{L/kg}$; TVM, UK) overdose anesthesia. The details of timing and pump speed can be found in the Supplementary Materials. Frontal sectioning of the brains ($40\mu\text{m}$ -thick for immunohistochemical staining and $300\mu\text{m}$ -thick for microinjection) was performed on a vibratome (Leica-VT1000s, Rueil-Malmaison, France). The injection or implantation site of each perfused mouse was checked under the microscope.

Immunohistochemical staining

We did fluorescent staining to examine the expression of c-Fos, VGLUT1, PSD-95, Neurogranin, and GFAP (see Supplementary Table 2 for the antibody concentrations). We used sections ranging from 1.42 mm to -0.23 mm away from Bregma, with a distance $160\mu\text{m}$ in between. The sections were firstly washed in $1\times$ PBS ($3\times 10\text{ min}$) and then blocked at room temperature (RT) with 5% donkey serum in 0.3% PBS-T (1 h). Later the sections were incubated at 4°C with corresponding primary antibody and 1% donkey serum in 0.3% PBS-T overnight. Sections were rinsed with $1\times$ PBS ($3\times 10\text{ min}$) in the next morning, incubated with secondary antibody in 0.3% PBS-T at RT (2 h), and rinsed again with $1\times$ PBS ($3\times 10\text{ min}$). Sections were mounted on gelatin-coated slides, air-dried, and coverslipped with Vectashield H-1000 (Vector Laboratories, Germany).

We stained IBA1 with 3,3'-Diaminobenzidine (DAB, Sigma, US). Sections were selected, washed, blocked, and treated with primary and secondary antibodies as described above. Then the sections were rinsed with $1\times$ PBS ($3\times 10\text{ min}$) and incubated with avidin-biotin-peroxidase complex (ABC Elite, Vector Laboratories, Germany; 0.2% A and 0.2% B in $1\times$ PBS) at RT (1.5 h). Later the sections were rinsed with 0.05 M Tris-HCl buffer (TB; pH 7.5; $3\times 10\text{ min}$). Peroxidase revelation was achieved by incubation shortly (20 s) with a mixture of 0.025% DAB and 0.0006% H_2O_2 in 0.05 M TB. Sections were carefully rinsed with TB ($2\times 10\text{ min}$) and $1\times$ PBS ($2\times 10\text{ min}$) to cease the reac-

tion. All sections were mounted and air-dried, then dehydrated in graded alcohol baths (1×5 min in 70%, 1×5 min in 90%, and 2×5 min in 100%), cleared in Roti-Histol (Carl Roth, Germany), and coverslipped with Eukitt.

Microinjection

We used microinjection and confocal microscope (Dumitriu et al. 2011) to visualize and quantify the neural morphology at 24 h and 48 h post-stimulation. The sections were selected within the distance of approximately ± 0.4 mm anterior-posterior (AP) away from the optic fiber. The injection was done only into the pyramidal neurons from layer 2 – 3 of ACC (24a/24b) from both hemispheres. The injection pipettes were pulled from glass capillaries with filament, with a final resistance around $150 \text{ M}\Omega$. We filled the pipette with red fluorescent dye solution Alexa 568 hydrazide (#A10441, Thermo Fisher, USA) in filtered $1 \times \text{PBS}$ (1 : 40). We performed microinjection under the microscope of a patch-clamp set-up. During injection, we penetrated the pipette tip into the soma and switched on the current to -20 pA to drive the dye diffusion for 20 min. Later we switched off the current but left the pipette tip inside the soma for another 5 min to fill the dendrite and spines. All the sections were retrieved and covered with Vectashild H-1000 (Vector Laboratories, Germany) for confocal microscope imaging. We checked all the injected neurons for YFP signal; only neurons with YFP signal were identified as pyramidal neurons and selected for further analysis.

Microscope imaging

To quantify the expression of c-Fos, VGLUT1, PSD-95, neurogranin, and GFAP in the ACC, we imaged epifluorescent signals of stained sections with Morpho Strider on Zeiss Imager2 (Carl Zeiss, Germany) with $2.5\times$ objective. To achieve better resolution of representative images, we also imaged the sections at the middle focal plane with a confocal microscope Leica SP8 (Leica Microsystems, Germany; software Leica SP8 LAXS 3.5.6) with objectives HCX PL Fluotar $5 \times /0.15$, $20\times$, and HC PL APO CS2 $63 \times /1.40$.

The bright-field images of DAB-stained IBA1 were acquired with a NikonEclipse E600 microscope with 4 $\times$ and 40 $\times$ objectives (MBF Bioscience, USA; software Neurolucida 2019).

To analyze the morphological features, we took z-stacked images of microinjected neurons (with step size 0.2 μm -0.3 μm sampled by the software Leica SP8 LAXS 3.5.6) with confocal microscope Leica SP8 (Leica Microsystems, Germany). The entire neuron structure was imaged with the objective HC PL APO CS2 63 $\times$ /1.40. If not stated otherwise, we used a pulsed laser for excitation (White Light; 488 nm). Segments of apical and basal dendrites of each neuron were imaged with the objective HC PL APO CS2 63 $\times$ /1.40 as well. Second and third level dendrite segments with little overlap and clear background were selected.

3D reconstruction and analysis of dendritic morphology

Firstly, after imaging, we deconvolved our confocal z-stack images with Huygens Professional 19.04 (Scientific Volume Imaging, The Netherlands) to restore the object from the acquired image through knowledge of the point spread function (PSF) and noise. 3D reconstruction and morphological analysis were later performed on the deconvoluted images.

For each pyramidal neuron, we reconstructed the soma and its dendritic tree with Imaris 9.5.1 (ImarisXT, Bitplane AG, Switzerland). Based on the reconstructed data, the dendritic tree structure was represented by Sholl intersections (Sholl 1953) at different radiuses. The order of each dendrite segment and its corresponding length and average diameter were also estimated. We further used Fiji (ImageJ, Fiji) to measure the soma size of each neuron on its z-projected image.

We reconstructed the dendritic shafts and spines with Imaris 9.5.1 again for selected dendrite segments at high resolution. We also classified the spine classes (filopodia, long-thin, stubby, and mushroom) based on their morphological features with the Imaris Spines Classifier package. The criteria of spine classification were summarized in Supplementary

Table 3. We harvested the overall spine density of each segment and the spine density of each spine class based on the reconstructed data. The spine head volume of individual spines was also estimated.

Quantifying immunohistochemical staining images

Visually inspection showed the expression of marker proteins was not homogeneously distributed in ACC but constrained to the vicinity around the optic fiber. To reflect such a pattern, we systematically analyzed the expression of c-Fos, VGLUT1, PSD-95, Neurogranin, GFAP, and IBA1 in both hemispheres at different distances to the optic fiber at 1.5 h, 24 h, and 48 h post-stimulation.

We firstly organised the corresponding epifluorescent images (obtained under $2.5\times$ objective) or bright-field images (obtained under $4\times$ objective) of each marker for each mouse in sequential order. The Bregma level of each section was identified in reference to the Mouse Brain Atlas (Franklin et al. 2008). Later we checked the implantation site for each section. Sections with a clear trace of implantation were marked as *distance zero*. Sections at a more anterior position than the distance zero were labeled with a negative sign ($-$), while posterior sections were labeled with a positive sign ($+$). In the end all sections were classified into five distance groups and their average distances were noted as -0.4 mm, -0.2 mm, 0 mm, $+0.2$ mm, and $+0.4$ mm. Both hemispheres were also carefully identified as the ipsi- or contralateral side in reference to the implantation site.

To quantify the signal intensity of markers on each section, we created two same-sized masks ($700\text{ }\mu\text{m} \times 700\text{ }\mu\text{m}$) on both hemispheres with Fiji. For c-Fos and IBA1, we counted signal positive cell numbers within each section’s masks, while for VGLUT1, PSD-95, neurogranin, and GFAP, we quantified the fluorescent intensity within the masks. The quantified fluorescent intensity or cell count at each discrete time point were respectively normalized by the averaged intensity or cell count from the ipsilateral side of the *zero distance* sections of the sham mice.

For neurogranin, in addition to the overall signal intensity, we also quantified its

relative intensity in the soma to infer its cellular translocation (FI_{soma}). Sections within 0.1 mm anterior-posterior to the optic fiber were selected for this analysis. It was easy to identify the neurogranin-positive neurons for sections from all three sham groups and the two stimulated groups, respectively, from 90 min-post and 48 h-post stimulation. Signals from these neuronal somas are often clearly organized in round shapes with brighter intensity against the salt-and-pepper pattern in the background. In contrast, sometimes the soma intensity is darker for the stimulated group at 24 h-post stimulation. To select the neurons without a bias, we firstly checked if any round cell-like shape is detectable by zooming into the micrographs. We further examined all round-circled signal clusters: soma area with signal dots were included, while empty somas were excluded to rule out non-pyramidal neurons. The selected somas were also verified with the channel-merged micrograph of YFP signals. Since YFP was sparsely tagged with pyramidal neurons, we finally noted down the soma size for post-hoc examination. In all the identified neurons, we first drew the soma shape and measured its fluorescent signal intensity and area size. Then we moved the mask to the neighboring area around the soma and measured the fluorescent intensity of the same-sized area as a reference. Five random selections were measured in the adjacent regions and averaged to serve as the reference. We normalized the signal intensity of soma by the signal intensity of its neighboring area as the relative soma intensity. We found that soma size and soma signal intensity were uncorrelated, so all the data points were used for statistical analysis.

Statistical analysis

We have different types of data in the current study, non-clustered independent data and nested data. Independent measurement, such as behavioral data, was contributed only once by each mouse. We used the non-parametric Mann Whitney U test to examine if the optogenetic stimulation triggered significant behavioral alterations.

Some datasets, such as the signal intensity of immunohistochemical staining and the neural morphology, are highly nested. In the staining experiments, each mouse con-

tributed multiple brain sections in five distance groups; for the morphological data, each mouse contributed several neurons, and each neuron further contributed multiple dendrite segments. In such conditions, using multiple measurements from each mouse as independent measurements artificially inflates sample size N and risks our study for achieving inappropriate conclusions. Indeed, we observed highly significant results for all quantified data obtained in IHC and microinjection experiments, when applied tests such as Mann Whitney U test or Kruskal Wallis test (Supplementary Materials: Comparing LMM with other tests). We therefore used a more conservative analysis, linear mixed effects model (LMM) in R (R Core Team 2019), to assess the effects of stimulation, while accounting for the nested residual structure. We used the lme4 package (Bates et al. 2015) and applied glmer function (GLMM) to model cell counting data. Our null hypothesis is that there is no significant difference between the sham and the stimulated mice, neither between the ipsi- and contralateral hemisphere of each mice. So in the model, we set the main effects of stimulation, implantation site, and their interaction effects as fixed effects. On the other hand, neuron ID, animal ID, and distance level are random effects (Zuur et al. 2009). Three discrete time points were separately analyzed. All models were checked in terms of homogeneous and normally distributed residuals, using diagnostic plots. We further checked final models for over-dispersion. Detailed model structures and the model selection and validation processes were described in the Supplementary Materials. All the R scripts of LMM and GLMM could be found under the following link: <https://github.com/ErbB4/LMM-GLMM-R-plasticity-paper>

The significance of fixed effects was tested by extracting effect strengths of each parameter, including their confidence intervals; $p < 0.05$, $p < 0.01$, and $p < 0.001$ were used to indicate 95% CI, 99% CI, or 99.9% CI of the estimated coefficient does not cross zero. If not stated otherwise, * denoted the main effect of optogenetic stimulation (sham/stimulated), # denoted the main effect of stimulation side (ipsi/contra to the optic fiber). Significant interaction effects were not denoted but stated in the main text. “n.s.” denoted neither main effects nor interaction effects were significant.

Neuron, synapse, and network models

Numerical simulations of networks with homeostatic structural plasticity were used as a framework to interpret the outcome of our various measurements in mouse experiments. We used the same neuron model, synapse model, and network architecture, as published in our previous paper on transcranial electric stimulation (Lu et al. 2019). All the plastic neuronal network simulations of the current study were performed with the NEST simulator using a MPI-based parallel configuration (Linssen et al. 2018).

The current-based leaky integrate-and-fire (LIF) neuron model was used for both excitatory and inhibitory neurons. We employed an inhibition-dominated recurrent network with 10 000 excitatory and 2 500 inhibitory neurons to represent the local network of ACC (Brunel 2000). All neurons in the network receive Poisson drive at a rate of $r_{\text{ext}} = 30$ kHz to reflect external inputs. All connections involving inhibitory neurons in this network were established randomly with 10% connection probability and then kept fixed. Only excitatory to excitatory (E-E) connections were subject to homeostatic structural plasticity (HSP) (Gallinaro and Rotter 2018; Lu et al. 2019). Each excitatory neuron monitors its own firing rate using its intracellular calcium concentration and grows or retracts its spines and boutons to form or break synapses (Figure 6A, see Supplementary Materials for details). Initially, the network has no E-E connections at all, but spontaneous growth goes on until an equilibrium between network activity and structure is reached. In this equilibrium state, the E-E connectivity is at 9%, and all excitatory neurons fire around the rate of 8 Hz imposed by the controller. Spiking activity is generally asynchronous and irregular. Detailed parameters of the neuron model, synapse model, and network model can be found in Supplementary Tables 4, Table 5, and Table 6. The methods employed to measure neuronal activity and connectivity in numerical experiments are described again in more detail in the Supplementary Materials.

Modeling optogenetics

Optogenetics uses microbial opsin genes to achieve optical control of action potentials in specific neuron populations (Yizhar et al. 2011). In our current study, we used humanized channelrhodopsin-2 (hChR2), a fast light-gated cation-selective channel, to depolarize mouse pyramidal neurons (Nagel et al. 2003). The kinetics of ChR2 activation is a complicated light-dark adaptation process: Light activates and desensitizes the channels, while they recover in the dark phase (Bruun et al. 2015; Zamani et al. 2017). These state transitions have been studied in detail in computational models of ChR2 (Nikolic et al. 2013; Williams et al. 2013). It did not seem necessary, however, to include the detailed kinetics of ChR2 in our large spiking neural network with homeostatic structural plasticity. To reduce the complexity of the model and save computational power, we conceived optogenetic stimulation as an extra Poisson input of rate $r_{\text{opto}} = 1.5 \text{ kHz}$ and weight $J_{\text{opto}} = 0.1 \text{ mV}$. Neurons responded with an increased firing rate to this stimulation, as observed in optogenetic stimulation experiments.

Numerical experimental design

In mouse experiments, the ACC of animals were optogenetically stimulated for four consecutive days at the same time, with a duration of 30 min per day. In our computational model, we started the optogenetic stimulation after the network had reached its structural equilibrium. To avoid excessively long simulation time, we accelerated the remodeling process by employing relatively fast spine and bouton growth rates, see Figure 2 of Gallinaro and Rotter (2018) for details. The relative duration of stimulation vs. relaxation was left unchanged, however. Since the optogenetic stimulation in experiments activates a large fraction of all pyramidal neurons in ACC, we chose to stimulate half of the excitatory neurons ($f_{\text{opto}} = 50 \%$) in the model. All the model parameters are summarized in Supplementary Table 7.

Results

Optogenetic activation of ACC (24a/24b) pyramidal neurons triggered cortical hyperactivity and behavioral alterations

To study the time course of neural structural plasticity, we adopted the optogenetic mouse model previously published in our laboratory (Barthas et al. 2015, 2017) in which we repetitively activated ACC pyramidal neurons for four days. Using electrophysiological recordings and optogenetic approaches, we have previously shown that the hyperactivity of anterior cingulate cortex drives chronic pain-induced anxiodepressive-like behavior. Therefore, we wondered whether activating ACC pyramidal neurons via optogenetic stimulation can trigger the same molecular and behavioral changes as seen in depressive-like behaviour. Indeed, the functional validation of ChR2-YFP expression using *ex vivo* electrophysiological recordings confirmed that the optogenetic stimulation at 20 Hz reliably triggers action potential firing of the ACC pyramidal neurons. We also observed *in vivo* that ACC optogenetic stimulation leads to a robust induction of c-Fos, compared to controls. In addition, four consecutive days of stimulation (30 minutes per day) were sufficient to trigger an MPK-1 upregulation in the ACC, as well as depressive-like phenotypes in naive animals (Barthas et al. 2017). Although a single stimulation might have been sufficient to trigger structural plasticity, in our structural plasticity analyses we focused on the effect of chronic optogenetic stimulation to induce robust and reproducible behavioural and molecular alterations.

First, we compared the viral transfection and transgenic approaches (Supplementary Figure 1-1A-E). We have previously shown with the transgenic approach that there was an increased c-Fos expression at 1.5 h after the stimulation (Barthas et al. 2015) and here we also reproduced the results with viral injection approach (Supplementary Figure 1-1F-G; Figure 1G, left panel). Besides the cortical hyperactivity, both approaches induced a depressive-like phenotype in mice at 24 h and 48 h post-stimulation (Supplementary Figure 1-1H-I; Figure 1E-F) as published before. To avoid double surgeries, we decided

to continue with transgenic mice throughout the current study. We also confirmed that light did not trigger behavioral alterations (Supplementary Figure 1-2).

In two other batches of transgenic mice, we also examined the c-Fos expression at 24 h and 48 h post-stimulation and observed no difference between the stimulated and sham mice (Figure 1G, middle and right panels). These data suggested that optogenetic stimulation triggered hyperactivity in the ACC was restored to baseline level at 24 h and 48 h post-stimulation.

To capture the temporal evolution of neural plasticity, we stimulated transgenic Thy1-ChR2-YFP male adult mice and harvested their brains at 1.5 h, 24 h, and 48 h post-stimulation for further experiments (Figure 1A-C). Mice used for immunohistochemical staining experiments showed depressive-like behaviors at 24 h and 48 h post-stimulation, as shown by decreased grooming behaviour in splash test and increased latency to eat in the NSF test (Figure 1D-F). All mice group information and experimental design were summarized in Supplementary Table 1.

Figure 1: Consecutive optogenetic stimulation triggered hyperactivity in ACC and depressive-like behavior in mice. **A-C** Experimental design of the current study. **D-F** Mice used for immunohistochemical staining experiments showed depressive-like behavior at 24h and 48h post-stimulation. **D** Splash test results of mice sacrificed at 24h post-stimulation ($p = 0.003$, Mann Whitney U test; $N = 9$ for sham, $N = 10$ for stimulated). **E-F** Results of NSF and splash tests for mice sacrificed at 48h ($p = 0.01$ and $p = 0.005$, Mann Whitney U test; $N = 5$ for sham, $N = 5$ for stimulated mice). **G** c-Fos expression was elevated by the optogenetic stimulation at 1.5h ($p < 0.001$) and specifically on the ipsilateral side ($p < 0.01$), decayed to baseline at 24h and 48h post-stimulation. The x ticks label the mean distance of stained sections (–, anterior; +, posterior) from the optical fiber. The y axis labels the c-Fos+ cell number within the masks of stained sections. GLMM was used for the statistical analysis. $p < 0.05$, $p < 0.01$, and $p < 0.001$ means 95% confidence interval (CI), 99% CI, and 99.9% CI does not cross zero respectively.

VGLUT1 and PSD-95 in ACC showed time-dependent regulation by optogenetic stimulation

Structural plasticity could be reflected in changes of the number or size of synaptic contacts, but also in changes of network connectivity. Quantifying changes in network connectivity, however, is an unsolved challenge in the field. Before zooming directly into the microscopic analysis of synaptic contacts, using immunohistochemical staining against synaptic proteins, we determined when and where structural plasticity occurred. We first quantified the overall expression of presynaptic VGLUT1 and postsynaptic PSD-95 in the ipsi- and contralateral hemispheres of ACC sections with reference to the hemisphere where the optic fiber was implanted, at 1.5 h, 24 h, and 48 h (Supplementary Table 1). Frontal-sectioned brain slices in both sham and the stimulated mice were organized based on their distance away from the optic fiber.

The representative fluorescent staining of VGLUT1 was organized by distance and by time in Supplementary Figure 2-1. Intensity quantification summarized in Figure 2A-B showed that optogenetic stimulation did not trigger significant alteration at 1.5 h (95% CI = $[-0.083, 0.070]$, LMM), while significant upregulation was observed in the stimulated mice compared to sham mice at 24 h (99% CI = $[0.009, 0.477]$, LMM). Further examination of interaction effects confirmed stronger upregulation in the ipsilateral side (99.5% CI = $[-0.344, -0.116]$, LMM) in the stimulated mice. At 48 h, no more significant difference was detected between the stimulated and sham mice (95% CI = $[-0.012, 0.138]$, LMM). Our data at discrete time points suggested that optogenetic stimulation altered VGLUT1 expression in a time-dependent manner. Indeed, the upregulation was observed after 1.5 h, peaked around 24 h, and returned to baseline at 48 h post-stimulation, while the stimulation effects were constrained to areas around the optic fiber.

Similar expression pattern was observed with PSD-95 staining (Supplementary Figure 2-2, Figure 2C-D). At 1.5 h, no significant changes were induced by the stimulation (95% CI = $[-0.051, 0.018]$, LMM). At 24 h, enhanced expression of PSD-95 in ACC was observed in the stimulated mice (95% CI = $[0.008, 0.55]$, LMM) and specifically in the

ipsilateral hemisphere (99.5%CI = $[-0.335, -0.109]$, LMM). At 48 h, although the effect size was small, the PSD-95 expression of the stimulated mice declined to a lower level than sham (99% CI = $[-0.171, -0.005]$, LMM). Our data suggested a similar time-dependent manner of PSD-95 upregulation as VGLUT1 after the optogenetic stimulation: upregulation at 24 h and downregulation at 48 h.

Neurogranin was not upregulated by optogenetic stimulation

Since PSD-95 is expressed in the postsynaptic membrane of glutamatergic synapses in both excitatory and inhibitory neurons (Zhang et al. 1999), we studied another postsynaptic protein, neurogranin, which is exclusively expressed in the pyramidal neurons (Singec et al. 2004). Despite the fact that the same type of quantification and analysis procedures were applied to neurogranin stained ACC sections, no time-dependent or side-dependent alterations of neurogranin was observed (Supplementary Figure 2-3, Figure 2E-F; 95% CI = $[-0.111, 0.031]$, 95% CI = $[-0.170, 0.277]$, 95% CI = $[-0.102, 0.266]$, respectively, LMM).

Considering that neuronal stimulation could drive the translocation of neurogranin from soma to dendrites (Huang et al. 2011), we suspected that the optogenetic stimulation might fail to trigger neurogranin upregulation but induced the cellular translocation. Consequently, we selected sections within 0.1 mm anterior-posterior to the optic fiber and quantified the relative fluorescent intensity of neurogranin in the soma (insets in Figure 2E-F). Pyramidal neurons from the three sham groups all showed a high soma concentration. After the stimulation, the relative signal intensity of soma was slightly increased at 1.5 h ($p < 0.001$, Mann Whitney U test), decreased to a level lower than 1 at 24 h ($p < 0.001$, Mann Whitney U test), and recovered to a level above 1 but lower than the sham group at 48 h ($p < 0.001$, Mann Whitney U test). These data suggested that the optogenetic stimulation may not trigger neurogranin upregulation, but induce translocation with time: concentrated in soma at 1.5 h, translocated away from soma at 24 h, and recovered at 48 h.

Since neurogranin is essential for the calcium concentration dependent translocation of

calmodulin (Huang et al. 2011), the translocation of neurogranin hinted at the fact that calcium based plastic changes mostly occur in non-somatic compartments, most probably the dendrites.

Figure 2: Continued on the following page.

Figure 2: Optogenetic stimulation altered the expression of VGLUT1 and PSD-95 and the cellular translocation of neurogranin. **A, C, E** Representative images of VGLUT1, PSD-95, and neurogranin staining on the ipsilateral hemisphere of sections within 0.1 mm anterior-posterior (AP) to the optic fiber from both sham and stimulated mice. The insets in panel **E** display the example neurons. **B** Normalized VGLUT1 fluorescent intensity at different times and different distances to the fiber optic. The main effect of optogenetic stimulation (sham/stimulated) was significant at 24 h. **D** The PSD-95 fluorescent intensity at different times and distances. At 24 h and 48 h, the main effect of stimulation was significant. **F** The overall fluorescent intensity of neurogranin was not changed by stimulation (LMM), while the relative intensity of neural soma (FI_{soma}) from the stimulated mice was slightly increased at 1.5 h, greatly dropped below 1 at 24 h, and recovered to a level lower than sham at 48 h ($p < 0.001$, $p < 0.001$, $p < 0.001$, respectively, Mann Whitney U test).

Dendritic tree structure was not drastically affected by optogenetic stimulation at 24 h and 48 h

The analysis of the expression of VGLUT1, PSD-95, and neurogranin suggest dendritic changes on the ipsilateral side in sections close to the optic fiber from the stimulated mice, at 24 h and 48 h after the stimulation. The amount of VGLUT1 and PSD-95 correlates with the total synaptic strength for all synapses. To specify whether the change of synaptic proteins corresponds to an alteration of synapse numbers or modifications of individual synaptic strength, we injected a fluorescent dye to visualize and analyze the neuronal morphology.

We then stimulated mice the same way as described above and harvested their brains at 24 h or 48 h post-stimulation. As shown in Figure 3A, mice showed depressive-like behavior as expected ($p = 0.004$ for 24 h-post group, $p = 0.012$ and $p = 0.006$ for 48 h-post group, Mann Whitney U test). We injected red fluorescent dye (Alexa 568) into pyramidal neurons selected from the area around the optic fiber to visualize and analyze the neuronal morphology.

Neural dendritic structure at 24 h and 48 h was visualized as in Figure 3B. Pyramidal neurons from both ipsi- and contralateral ACC were collected (Figure 3C). The soma size and dendritic tree structure evaluated by Sholl intersections were not changed by the optogenetic stimulation (Figure 3D-E). No remarkable changes were detected in nei-

ther dendritic length nor average dendritic diameter, except that some dendritic segments showed a reduction or increase in dendritic diameter (Supplementary Figure 3-1). Our data suggested, except for local dendrite diameter changes, no drastic dendritic tree structure and soma size inflation or shrinkage of pyramidal neurons in the vicinity of the optic fiber were induced by the optogenetic stimulation.

Figure 3: Neuronal dendritic tree structure was not drastically affected by the optogenetic stimulation at 24 h and 48 h. **A** Mice sacrificed at 24 h and 48 h for microinjection experiments showed depressive-like behavior in splash and NSF tests (for both batches, $N = 6$ for sham and $N = 6$ for the stimulated; $p = 0.006$, $p = 0.012$, and $p = 0.004$ respectively, Mann Whitney U test). **B** Representative example of neurons filled with red fluorescent dye. **C** Overall distribution of pyramidal neurons injected in layer 2-3 of ACC from both hemispheres for both batches. Black dots are from sham mice, and red dots are from the stimulated mice. For the 24 h-post group, we selected 32 well-injected neurons in total and 15 neurons were from sham mice; each mouse contributed 2.67 neurons on average (SD = 1.31). For the 48 h-post group, we selected 24 neurons in total and 12 neurons were from sham mice; each mouse contributed 2 neurons on average (SD = 2.12). **D** The soma size was not changed by stimulation (LMM). **E** Dendritic tree structure was not altered by stimulation (GLMM).

Optogenetic stimulation induced the opposite spine morphological changes at 24 h and 48 h

To further analyze morphological changes at dendritic level, we sampled several secondary to third level apical and basal dendritic segments from each neuron and did the 3D reconstruction of spines (Figure 4A-B). Besides spine density, we also evaluated the spine head volume and classified different types of spines such as filopodia, long-thin, stubby, and mushroom.

As shown in Figure 4C-D, the overall spine density was increased at 24 h ($p < 0.05$, LMM) and decreased at 48 h ($p < 0.05$, LMM) post-stimulation. The apical spine density showed the same tendency but the changes were not statistically significant; the basal dendrites showed significant spine density alterations ($p < 0.05$ and $p < 0.05$, LMM). Analysis by spine type suggested subtle changes in different spine types (Figure 4E-F). At 24 h, the spine density of filopodia and stubby type was increased in both apical ($p < 0.05$ and $p < 0.05$ respectively, LMM) and basal dendrites ($p < 0.05$ and $p < 0.05$, LMM). At 48 h, the spine density of long-thin type was reduced in apical dendrites ($p < 0.05$, LMM), while the stubby and mushroom type were reduced in basal dendrites ($p < 0.05$ and $p < 0.05$ respectively, LMM). These data suggested optogenetic stimulation triggered spinogenesis and spine retraction in both apical and basal dendrites at 24 h and 48 h post-stimulation respectively.

In addition, spine head volume data (Figure 4G-H) showed different regulation in apical and basal dendrites. At 24 h, the head volume distribution of long-thin spines in the apical dendrites was right-shifted to larger mean values by the optogenetic stimulation ($p < 0.05$, LMM) while no changes were detected in the head volume of basal dendrite spines. At 48 h, the spine head volume of mushroom spines in basal dendrites was left-shifted to smaller mean values by the optogenetic stimulation ($p < 0.05$, LMM), whereas the apical dendrites showed no significant difference. Our data suggested in addition to spine density changes, optogenetic stimulation induced spine enlargement and shrinkage at 24 h and 48 h post-stimulation respectively. The changes of overall spine density and

spine head volume align with the evolution of PSD-95.

Figure 4: Spine density and head volume showed the opposite changes at 24 h and 48 h post-stimulation. **A-B** Representative example of filled dendritic segments. **C-D** Spine density at 24 h (84 dendritic segments from sham and 111 segments from stimulated mice) and 48 h (101 segments from sham and 61 segments from stimulated mice). **E-F** Spine density of each class. **G-H** Cumulative distribution of spine head volume. LMM was used for statistical analysis.

Glial responses were involved in homeostatic plasticity induced by the optogenetic stimulation

Although glial cells were often linked with pathological brain states involving inflammation and injury, their involvement in regular synaptic plasticity has been frequently highlighted in recent studies (Dissing-Olesen et al. 2014; Haydon and Nedergaard 2015; Weinhard et al. 2018). The time course during the evolution of structural plasticity, however, has not been reported before. We thus stained glial fibrillary acidic protein (GFAP) as the markers for activated astrocytes (Hol and Pekny 2015) and ionized calcium-binding adaptor molecule 1 (IBA1) for both inactive and active microglia (Ohsawa et al. 2004) at 90 min, 24 h, and 48 h after the stimulation.

The fluorescent staining of GFAP was organized by distance and by time in Supplementary Figure 5-1. Our statistics analysis showed optic fiber implantation triggered astrocytes reactivation in both sham and stimulated mice, but the stimulation further enhanced the reactivation in the ipsilateral hemisphere throughout 48 h post-stimulation (Figure 5A-B). Similar results were observed with IBA1 staining (Supplementary Figure 5-2) as optogenetic stimulation induced significant enhancement of IBA1 expression at 24 h and 48 h post-stimulation ($p < 0.001$ and $p < 0.001$, GLMM) in the ipsilateral hemisphere and in the sections close to the optic fiber (Figure 5C-D). Since IBA1 labels microglia regardless of its activation state, the upregulation of IBA1 suggested microglia proliferation. Our data suggest that optogenetic stimulation triggered simultaneous astrocytes reactivation and microglia proliferation, and they displayed a time course that is different from the expression of synaptic proteins and neural morphological changes.

Figure 5: Expression of GFAP and IBA1 were upregulated by optogenetic stimulation at 24h and 48h post-stimulation. **A**, **C** Representative images of GFAP and IBA1 staining on the ipsilateral hemisphere of sections within 0.1 mm AP to the optic fiber from both sham and the stimulated mice. **B** The normalized GFAP fluorescent intensity at different time and distance. At 1.5h, the main effects of side of stimulation were significant (99.5% CI = $[-0.0923, -0.0356]$, LMM). At 24h, the main effects of optogenetic stimulation and side of stimulation were again significant (99.5% CI = $[0.0646, 0.399]$, 99.5% CI = $[-0.182, -0.0628]$ respectively); their interaction effects were also significant (99.5% CI = $[-0.264, -0.109]$). At 48h, the main effects of stimulation, side of stimulation, and their interaction effects were significant (95% CI = $[0.009, 0.094]$, 99.5% CI = $[-0.115, -0.043]$, 95% CI = $[-0.079, -0.006]$ respectively). **D** The IBA1+ cell counting at different time and distance to the optic fiber. At 1.5h, only the interaction between stimulation and side of stimulation were significant (95% CI = $[-0.188, -0.00016]$, GLMM). At 24h, the main effect of optogenetic stimulation was significant (95% CI = $[0.025, 0.237]$); their interaction effects were also significant (99.5% CI = $[-0.533, -0.105]$). At 48h, the main effects of optogenetic stimulation was significant (99.5% CI = $[0.073, 0.193]$); their interaction was also significant (99.5% CI = $[-0.208, -0.0369]$).

A computational model of stimulation-induced homeostatic structural plasticity

To achieve a clearer picture of the ongoing network remodeling dynamics in the current study, we interpolated the time course of synaptic protein expression and neural morphological changes within 48 h after four stimulation sessions (Figure 6A). The stimulation triggered immediate hyperactivity in ACC pyramidal neurons, as represented by c-Fos over-expression. Neural activity was restored to baseline at 24 h and 48 h. However, although neural activity was restored, we observed a delayed upregulation of synaptic proteins (VGLUT1 and PSD-95) and spine density at 24 h and a decrease to or below baseline at 48 h. These data suggest that the elevated synaptic proteins and spine density at 24 h do not contribute to sustaining high spontaneous neural activity. Three possibilities arise. (i) The upregulation of VGLUT1 and PSD-95 and spinogenesis failed to increase functional synaptic transmission among the stimulated pyramidal neurons. (ii) Optogenetic stimulation indeed increased the glutamatergic transmission, but additional mechanisms such as rapid E/I balance masked its effect on neural activity (Van Vreeswijk and Sompolinsky 1996; Shu et al. 2003; Zenke and Gerstner 2017). (iii) The upregulation of synaptic proteins and increase in spine number and volume are a consequence of firing rate homeostasis. Although we cannot directly reject options (i) and (ii) without electrophysiological recordings, it has indeed previously been shown that repetitive transcranial direct current stimulation (tDCS, 20 min * 3 days) triggered enhanced synaptic transmission and increased spine density of pyramidal neurons 24 h after the stimulation in mice (Barbati et al. 2020). Besides, many previous studies have suggested that strength and morphology of excitatory synapses are homeostatically regulated (Turrigiano et al. 1998; Konur et al. 2003; De Gois et al. 2005; Ehrlich et al. 2007; Van Ooyen 2011) with or without changes of inhibitory synapses (Lenz et al. 2019; Knott et al. 2002) after activity perturbation. Therefore, it is highly possible that in our case optogenetic stimulation triggered homeostatic regulation.

The question now is which neuronal mechanism can account for the observed time

course. Inhibitory STDP, inhibitory plasticity, synaptic scaling, and the Bienenstock-Cooper-Munro (BCM) model are commonly known homeostatic rules complementing Hebbian plasticity. These rules do not include synapse rewiring. In some cases, enhanced spontaneous neural activity emerges with enhanced synaptic weight, which does not fit what we observed at 24 h post-stimulation (Litwin-Kumar and Doiron 2014; Toyozumi et al. 2014; Lazar et al. 2009; Zenke and Gerstner 2017). We thus selected the model of homeostatic structural plasticity (HSP), which assumes structural changes regulated by firing rate homeostasis. We simulated an inhibition-dominated spiking neural network to represent ACC (Figure 6B), in which optogenetic stimulation was introduced to half of the excitatory population. Transient stimulation perturbed the neural activity and triggered synapse turnover as a result of homeostatic structural plasticity (blue curves in Figure 6B), as described in a previous publication (Lu et al. 2019). As a result of synaptic reorganization, the connectivity among the stimulated neurons remained elevated after stable firing rates were achieved (Figure 6C). In a repetition protocol based on our *in vivo* experiments, simulation results show that the connectivity among the stimulated neurons increased after each repetition. Within 48 h after the final stimulation (S4), the firing rate of the stimulated neurons rapidly returned to baseline. The connectivity, however, remained elevated and decayed only slowly (insets in Figure 6D). Although we could not differentiate stimulated and non-stimulated neurons in the mouse experiments, the neurons selected for morphological analysis were close to the optic fiber and, therefore, had a higher chance to be stimulated. The HSP model, thus, provided a fully consistent explanation for the observed c-Fos expression and spine morphology.

Figure 6: The homeostatic structural plasticity model. **A** The interpolated time course of the expression of c-Fos, VGLUT1, PSD-95 and spine morphology compared to sham (baseline). **B** Temporal evolution of neural activity and network connectivity in response to optogenetic stimulation. Light blue shaded areas indicate the stimulation period. Blue and dark gray curves in the upper panel represent the firing rate of stimulated and non-stimulated neurons. Blue, dark gray, and light gray curves in the lower panel represent the synaptic connectivity within or between populations. The blue and light gray curves finally coincided due to identical population sizes. **C** The connection matrix of all excitatory neurons at two time points before and after stimulation, labeled by orange dots in panel **B**. Columns are the presynaptic neurons, and rows are the postsynaptic neurons. Color indicates the average connectivity. The white square labels the intra-group connectivity of the stimulated neurons. **D** Repetitive optogenetic stimulation boosted the connectivity among the stimulated neurons. Small insets display the dynamics within 48 h after the last stimulation session.

Discussion

In the current study, we combined both mouse experiments and computer simulations to study structural plasticity. We first systematically investigated the neural activation and neural morphology of the neocortical region anterior cingulate cortex (ACC) after chronic optogenetic stimulation in an *in vivo* mouse model. We found that the activation of a subset of excitatory neurons in ACC over four consecutive days triggered substantial alterations. In fact, the temporal profiles of specific molecular and morphological changes over 48 h post-stimulation were intertwined in a specific way. The expression of VGLUT1 and PSD-95, as well as the spine density and spine head volume, were above baseline at 24 h and restored to baseline or slightly below at 48 h. Intriguingly, although such changes seem to suggest altered synaptic transmission, neural activity estimated by c-Fos expression did not show any change at 24 h and 48 h. After neural activity has rapidly returned to baseline, synaptic protein expression and spine density undergoes a rise and a decay as compared to the control (Figure 7, red and blue curves). Altogether, it appears as if synaptic plasticity regulated by firing rate homeostasis can explain the time course of events described above quite well. In fact, we verified with the help of computer simulations that the homeostatic structural plasticity (HSP) model, in principle, recapitulates the observed biphasic dynamics (Gallinaro and Rotter 2018; Lu et al. 2019; Gallinaro et al. 2020).

Figure 7: Joint summary of the results of mouse experiments and the computational model. Light blue shaded areas indicate four sessions of optogenetic stimulation (S1 to S4, 30 min per session). Red and blue curves represent the temporal profile of neural activity and connectivity of stimulated neurons, as predicted by the model. Colored symbols in the right panel represent the expression level of c-Fos, synaptic proteins, and glial markers, respectively. Measurements were made at 1.5 h, 24 h, and 48 h post-stimulation, shown are the values from stimulated mice relative to the corresponding sham group. For Neurogranin, we show its relative expression in dendrites. The gray dashed line represents the baseline. Symbols above or below the baseline indicate that values in stimulated mice are increased or decreased as compared to sham, respectively. The illustration of a neuron with dendrites and spines depicts that spine density and spine head volume were both increased at 24 h, but decreased at 48 h post-stimulation.

Our experiments elucidated how structural plasticity of pyramidal neurons evolves in time after optogenetic stimulation in the mouse experiments. Analysis of the expression of synaptic proteins clearly indicates that robust synaptic changes occurred at 24 h and 48 h after stimulation. VGLUT1 is the glutamate transporter protein that controls the quantal glutamate content of individual synaptic vesicles (Freneau et al. 2004; Wilson et al. 2005). Therefore, the upregulation and downregulation of VGLUT1 observed in our current study hints at a change of glutamate release in synapses, corresponding to the presynaptic strength accumulated over many neurons in a given tissue volume. PSD-95 is a scaffold protein in the postsynaptic density which transforms rapidly (Gray et al. 2006) and which is actively redistributed between the synaptic sites and the cytoplasmic pool (Bresler et al. 2001; Gray et al. 2006). Functionally, PSD-95 organizes the distribution of AMPA receptors (Zhang et al. 1999; Chen et al. 2011) and modulates spine morphology (El-Husseini et al. 2000; Pak and Sheng 2003; Fossati et al. 2015). We observed similar

biphasic changes in the expression of PSD-95 and VGLUT1 after stimulation.

However, the expression of neurogranin, a calmodulin-binding protein exclusively expressed in the soma and dendrites of excitatory neurons (Singec et al. 2004), behaved differently. Neurogranin has a molecular weight less than 10 kDa and thus can swiftly translocate within pyramidal neurons upon synaptic stimulation from the cell plasma to the nucleus (Garrido-García et al. 2009) or from the soma to the dendrites (Huang et al. 2011). We found that optogenetic stimulation failed to trigger upregulation or downregulation of neurogranin, but changed somatic signal intensity over time, which may suggest a translocation away from the soma in parallel with the upregulation of VGLUT1 and PSD-95. Functionally, neurogranin bidirectionally modulates synaptic plasticity by affecting the phosphorylation pattern of postsynaptic density proteins via calmodulin (Xia and Storm 2005; Zhong and Gerges 2012; Svirsky et al. 2020; Hwang et al. 2021). Upon activity perturbation, neurogranin also coordinates the conversion of AMPAR-silent synapses to AMPAR-active synapses, as well as synapse elimination (Han et al. 2017). Based on this evidence, we propose that translocated neurogranin may interact with postsynaptic density proteins. This could work in synergy with PSD-95 to modulate functional and plasticity. In addition, due to the dependence of translocation on calcium concentration (Huang et al. 2011) and the temporal profile of intracellular calcium dynamics, neurogranin might not be stored in spines and exert its impact on synaptic plasticity only at a later time.

All these observations point to a biphasic regulation of synaptic strength within 48 h and an orchestrated regulation of presynaptic and postsynaptic plasticity after stimulation, as reported before by others (Ehrlich et al. 2007; Letellier et al. 2019; Sanderson et al. 2020). The results of neural morphology analysis were in line with our observations in synaptic proteins. Optogenetic stimulation did not alter dendritic branching structure or soma size, as it was observed in the case of diseases (Chidambaram et al. 2019). Rather, stimulation induced biphasic changes at the level of dendritic spines. The density and volume of spines in the stimulated pyramidal neurons increased at 24 h and slightly decreased at 48 h, as compared to controls. This is highly interesting, as spine volume

correlates with synapse strength (Matsuzaki et al. 2004) and PSD-95 clustering (Cane et al. 2014). Increased spine density increases the chances to form new synapses. All things considered, we conclude that synaptic transmission and connectivity is increased at 24 h and restored to baseline at 48 h after stimulation.

Given the biphasic temporal profile of changes in synaptic proteins and dendritic spine morphology, we hypothesized that they might be a result of homeostatic regulation. In fact, the time evolution of synaptic protein expression reflects the accumulation of effects of all four stimulation sessions. The time course of PSD-95, for example, is around baseline at 1.5 h, upregulated at 24 h, and decayed slightly below baseline at 48 h after the last stimulation. Classical Hebbian plasticity depends on positive feedback, and it would systematically increase PSD-95 expression upon stimulation. Homeostatic plasticity, in contrast, depends on negative feedback, and it would at least transiently decrease the PSD-95 level. As the low level of PSD-95 immediately after the fourth stimulation is also 24 h after the third stimulation, a pure Hebbian mechanism is out of question, and a contribution of homeostatic control is likely (see Supplementary Figure 6-1 for a graphical illustration of the argument). A second independent argument can be derived from the temporal profile of c-Fos expression. Indeed, the hyperactivity expected by optogenetic stimulation in pyramidal neurons is not visible 24 h and 48 h after stimulation, possibly because homeostatic regulation has brought it back to its set point. Although c-Fos expression is not a very accurate indicator of neural activity, other *in vivo* electrophysiological recordings did confirm the robust firing rate homeostasis in the cortex (Pacheco et al. 2019). As a result, it seems possible that the observed changes in synaptic proteins and spine morphology reflect the dynamic process of homeostatic control to restore neural activity after activity perturbation. Theoretically, an alternative explanatory scheme may be linked to inhibitory plasticity (Vogels et al. 2011, 2013), although we found no evidence for this in our data.

We proposed the model of homeostatic structural plasticity to explain the observed biphasic changes of spine morphology at 24 h and 48 h post-stimulation, when the neural activity indicated by c-Fos expression was already back at baseline. This model explains

how neuronal firing rates are stabilized using structural plasticity linked with a homeostatic controller (Butz et al. 2009; Van Ooyen 2011; Butz and van Ooyen 2013). We have previously shown in computer simulations how external stimulation can trigger cell assembly formation by deleting connections and forming new synapses controlled by firing rate homeostasis (Gallinaro and Rotter 2018; Gallinaro et al. 2020; Lu et al. 2019). In computer simulations of the optogenetic experiment, we observed a very similar cell assembly formation process. Specifically, we showed that in this model the connectivity among stimulated neurons remained at a high level although the firing rate had already returned to baseline. Although we could not record the connectivity among the stimulated neurons in mouse ACC like we did in computer simulations, the changes of spine density in the pyramidal neurons sampled in the area close to the optic fiber served as a proxy and seemed to fit the computer simulations. Due to firing rate homeostasis, application and termination of the external stimulation should trigger slow homeostatic responses of opposite sign. Typical experiments, however, record a mix of changes occurring during perturbation and after perturbation, which may be of opposite sign. For instance, plastic changes observed as a result of a persistent lesion or denervation occur during input deprivation. In contrast, plastic alterations observed after stimulation, as in the current study, are mixed on and off effects. So it is critical to use an experimental design that includes both phases and measure during time periods that are long enough to re-establish the neural activity homeostasis. In addition to spine turnover and spine density, specific changes in connectivity represent another crucial feature that influences network function.

Nevertheless, the question arises whether similar effects on neural morphology would occur after a single stimulation already, rather than only after four stimulation sessions. Some alterations, such as developing a depressive-like phenotype, indeed requires chronic stimulation, as reported in our previous work (Barthas et al. 2017). The same rule, however, does not necessarily apply to changes in spine size and density. Chronic *in vivo* imaging has revealed ongoing spine turnover and fluctuation in spine head volume even under normal conditions (Holtmaat et al. 2006). However, external perturbation of

neural activity can accelerate network remodeling considerably. The magnitude of such changes depends on circumstantial factors, such as the developmental stage, but also on stimulation parameters, including the type, intensity, and duration of the perturbation. Changes in spine head volume generally seem to come first. Both weak stimulation (e.g. using glutamate uncaging) and supra-threshold stimulation (e.g. using repetitive magnetic stimulation) were reported to induce considerable changes in spine volume within a few hours post-stimulation (Vlachos et al. 2012a; Matsuzaki et al. 2004; Noguchi et al. 2019). Longer and/or stronger stimulation is required to induce changes in spine density, which involves protein synthesis and *de novo* spinogenesis (Trachtenberg et al. 2002; Holtmaat et al. 2006). In experiments, the question arises whether a single stimulation can trigger neural morphological changes at all, and whether potentially subtle changes can be detected statistically. It is very well possible that changes in spine head volume occur already after a single stimulation, but they might be masked by the inevitable inter-neuron and inter-animal variance. Repetitive imaging of one and the same dendritic segment might be necessary to answer this question. Assuming that a single stimulation already triggers the structural remodeling process, we have shown in computer simulations in a previous study (Lu et al. 2019) and this manuscript that repetitive stimulation can be accumulated, increasing synaptic connectivity in an incremental way. This may also explain why, in our experiments, we could observe elevated c-Fos expression in the contralateral hemisphere of the stimulated mice, but no significant changes in neural morphology. Neural activation of the contralateral hemisphere could either be due to inter-hemispheric projections, or due to light penetrating from the ipsilateral side. Both effects are certainly weaker than the effects of direct stimulation. As already mentioned above, subtle changes triggered by weak stimulation might not be detectable by the analysis methods used in our current study.

Our study also casts light on the relation between ACC hyperactivity, synaptic plasticity, and depressive-like behavior. ACC is a hub for negative affects, pain, and their comorbidity (Humo et al. 2019), **accompanied by different forms of synaptic plasticity** (Bliss et al. 2016). Chronic pain can induce hyperactivity and synaptic potentiation in

ACC, along with anxiodepressive behavior in mice (Sellmeijer et al. 2018; Koga et al. 2015). ACC hyperactivity artificially induced by optogenetic stimulation also generates depressive-like behavior in naive mice (Barthas et al. 2015, 2017). It is unclear, however, whether changes in neuronal activity, spine morphology, and depressive-like behavior develop in parallel due to a common condition, or whether there are causal links between individual factors (Gipson and Olive 2017). In our experiments, neural activity quickly decayed to baseline after stimulation, but the mice exhibited sustained depressive-like behavior, which can last for around two weeks after the stimulation was terminated (Barthas et al. 2017). So the alterations in depressive-like behavior seem to always lag behind changes in ACC neural activity. This evidence suggests that depressive-like behavior may be mediated by persistent changes which depend on the accumulated effects of neural activity. On the other hand, we observed that synaptic plasticity (synaptic proteins and spine morphology) exhibit transient changes during depressive behavior. This suggests that the time scales of structural plasticity and behaviour differ from each other.

Although glia cells are not the main focus of this study, we nevertheless showed that the time course of astrocyte activation and microglia proliferation matches the time course of homeostatic structural plasticity. Indeed, we observed increased numbers of IBA1-positive cells and enhanced GFAP expression throughout 48 h after stimulation, which suggests that both microglia cells and astrocytes were active at least between 1.5 h and 48 h post-stimulation. Microglia and astrocyte activation is often regarded as a sign of neural inflammation. This also raises the question whether the phenomena observed in our present study reflect a pathological state, or whether they should be interpreted as reflecting regular structural plasticity. We cannot fully exclude the possibility of neuroinflammation, as glial activation is heterogeneous, and there is no clear distinction between normal and pathological conditions anyway. Besides, accumulating evidence emphasizes the critical role of glial cells in cleaning up extracellular chemicals, fostering spinogenesis, and facilitating synaptic plasticity in normal brain development and plasticity (Eroglu and Barres 2010; Zamanian et al. 2012; Liddelow and Barres 2017). Glia-secreted pro-inflammatory cytokine tumor necrosis factor α (TNF - α), for instance, has been shown

to also regulate synaptic transmission and homeostatic synaptic scaling (Stellwagen and Malenka 2006; Steinmetz and Turrigiano 2010). According to our data, the structure of dendritic trees and soma size were only weakly affected by optogenetic stimulation. Therefore, it seems unlikely that glial activation is caused by excitotoxicity and neural inflammation. Further studies are of course needed to explore if microglia and astrocyte activation triggered by optogenetic stimulation are involved in maintaining chemical homeostasis (Jo et al. 2014) and inducing morphological changes of spines (Weinhard et al. 2018).

In conclusion, our joint experimental-theoretical efforts provide evidence that, in response to supra-threshold optogenetic excitation, neurons modulate their synaptic connectivity to restore neural activity in a homeostatic way. The homeostatic structural plasticity model was able to qualitatively explain the observed time course of neural activity and spine morphology. Further joint work is needed to capture the effects of activity perturbation on specific network connectivity and include functional aspects to structural plasticity models.

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Author contributions

HL, JG, CN, SR, and IY conceived the project and designed the experiments. HL, JG, and SR established the computer model. HL performed the network simulation and analysis. HL performed all the mice experiments, collected data, and performed the analysis. CN and IY helped with the experiments. IY proposed the quantification strategy. HL wrote the manuscript, and all the authors revised and approved the paper.

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Supplementary Materials

Virus injection

To compare the effects of optogenetic stimulation between the transgenic animals and viral injected ones, we also transfected C57BL/6J mice with virus pAAV-CaMKIIa-hChR2(H134R)-EYFP (Addgene plasmid #26969; <http://n2t.net/addgene:26969>; RRID: Addgene_26969). We injected the same virus bilaterally into the ACC (0.5 μ L per side, with a speed of 0.05 μ L per 30 seconds) in both stimulated and sham mice. 15 min after the injection, We removed the Hamilton syringe from the brain. The same stimulation protocol and behavioral tests were used for both transfected mice and transgenic mice.

Tissue harvesting

For immunohistochemical staining, the perfusion protocol contains 2 min wash with cold $1\times$ phosphate buffer (PB, 0.1 M, pH 7.4) solution and 18 min perfusion with 4% paraformaldehyde (PFA) in $1\times$ PB (with the pump speed at 10 mL/min). Then brains were extracted and kept in 4% PFA in $1\times$ PB overnight for post-fixation. Brains were later transferred and stored in $1\times$ PBS with 0.02% sodium azide at 4 °C for further experiments.

For microinjection experiments, we used a fast perfusion method to protect the fine dendritic structure (Dumitriu et al. 2011). We washed the blood with 1% PFA in $1\times$ PB for 2 min, and perfused animals for another 12 min with 4% PFA in $1\times$ PB with 0.125% glutaraldehyde. The pump speed was 5 mL/min. Then we extracted the brains and put them in 4% PFA in $1\times$ PB with 0.125% glutaraldehyde for another 6 hrs for post-fixation. All brains were later transferred and stored in $1\times$ PBS with 0.02% sodium azide at 4 °C for further experiments.

Linear mixed models

In the current study, we aimed to analyze the difference between the sham and stimulated mice, in terms of the expression of markers and neural morphology. Besides, we wanted to

specify if the difference between the stimulated and sham groups showed various patterns on the ipsi- and contralateral hemispheres. In principle, these analyses are analyses of variance (ANOVA) or two-way ANOVA, depending on the number of explanatory variables. Applying ANOVA, however, demands the satisfaction of certain assumptions. The relationship between the explanatory variables and the response variable should be linear. The independence of observations ought to be ensured. Apart from that, the variance of observations should be normally distributed and remain homogeneous for different groups. In the current study, however, each mouse either contributed several sections for each distance group or provided several neurons, which violated the independence assumption. The distribution of data for each mouse or each group is not necessarily homogeneous. In such conditions, initial trials with Mann Whitney U test or Kruskal Wallis test returned appropriate highly significant results and risked our study for achieving inappropriate conclusions. So we used a new method to deal with the heterogeneity and dependence of our data points, the linear mixed model (LMM).

IHC quantification and LMM

In an LMM model, we specified the effects of fixed and random components. Fixed effects refer to the effects explained by the explanatory factors, while random effects refer to the nestedness of data. With visual inspection, we used being stimulated or not (sham/stimulated) and stimulation side (ipsi/contra) and their interaction effect as the fixed term to explain their difference. Animal ID or section ID and distance to the optic fiber are the random term. With this model, we can approximate the effects of explanatory variables as much as possible with all the data points collected from multiple mice and meanwhile control the discrepancy between mice.

In our study, we used LMM to analyze the results of two experiments, the immunohistochemical staining and microinjection, with different fixed effects and random effects structures. Here we showed the analysis of immunohistochemical staining results as a detailed example. The analysis of morphological data was similar and we will skip the details.

We quantified the protein signal intensity of both hemispheres of ACC sections from five distance groups in both sham and stimulated mice groups. We **hypothesized** that the overall signal intensity is different in the stimulated mice from the sham group. Moreover, the ipsilateral hemisphere of the stimulated mice should be different from the contralateral hemisphere, too, since the ipsilateral side is closer to the optic fiber and receives more stimulation. The contribution of stimulation side (ipsi/contra) to the signal intensity of the sham mice, in theory, would not be as robust as in the stimulated group. Thus, we proposed two explanatory variables and their interaction effects as the master structure of the fixed effects. As for random effects, we used mice and distance group as the group factor for sections. As showed in Equation S.1,

$$\text{Signal_intensity} \sim \text{Sham} * \text{Ipsi} + (1|\text{AnimalID}/\text{Distance2}) + \epsilon, \quad (\text{S.1})$$

Signal_intensity is the response variable. The first part follow the symbol $\sim$ is the fixed effects structure where *Sham* and *Ipsi* are two nominal explanatory variables and $*$ denoted the interaction effects between them. $(1|\text{AnimalID}/\text{Distance2})$ is the random effects structure, grouping sections per mouse and per distance group. ϵ is the residual.

After fitted the model, we plotted the fitted values' residuals against each explanatory variable to inspect if the model fits well visually. The significance of fixed effects was tested by profiling the confidence interval of each estimated parameter. $p < 0.05$, $p < 0.01$, and $p < 0.001$ means 95% CI, 99% CI, or 99.9% CI of the estimated coefficient does not cross zero.

Counting data and GLMM

In the case of counting data, such as IBA1+ and c-Fos+ cell counting, Poisson distribution of data points was assumed, so LMM was no longer suitable. We used the generalized linear mixed model (GLMM) to fit the counting data. The same process, as described above, was used to estimate the parameters. In the last step, in addition to visual inspection, we also applied the over-dispersion test to validate our model. In the case of over-dispersion

(> 2.0) or underdispersion (< 0.6), we adjusted the model back and forth. Significance notification for the coefficients estimated by GLMM models remained the same as LMM models.

Morphological analysis and LMM

Different from immunohistochemical staining data, we did not specify distance information for each neuron. All the neurons were selected from sections within ± 0.4 mm anterior-posterior (AP) away from the optic fiber and treated as homogeneous. So regarding the random effects structure, we used two types depending on the response variable. If quantifications were taken per neuron, such as soma size and Sholl intersections, we used mice as the group factor (model1). If quantifications were taken per dendrite segment of individual neurons for each mouse, such as the dendritic length, average dendritic diameter, and spine density, we used a two-layer group factor with both neurons and mice (model2).

```
model1 <- lmer(Soma_size~Sham*Ipsi + (1|AnimalID),REML=TRUE,data=time24h)
model2 <- lmer(Spine_density~Sham*Ipsi+(1|AnimalID:NeuronID),REML=TRUE,data=time24h)
```

Comparing LMM with other tests

Here in this section, we showed an example dataset and compared the results of different analysis. Supplementary Figure 4-1 showed the distribution of stubby spine head volume from the apical dendrites for each mouse sacrificed at 24 h post-stimulation. Visually we saw each mouse contributed several spines from various segments. If we ignored the independence assumption and pooled all the data points, the difference between sham and the stimulated mice was significant ($p = 0.0002$, the Kolmogorov-Smirnov test; $p = 2.8 * 10^{-7}$, Mann Whitney U test). If we specified the random effects and grouped the data points by the dendrite segment and by mouse, the sham and the stimulated mice were no longer significantly different from each other ($p > 0.05$, LMM) as presented in Figure 4G (upper panel). So breaking the independence assumption could result in inappropriate conclusions.

Neuron model

The membrane potential dynamics of linear integrate-and-fire (LIF) neuron model is explained by Equation S.2,

$$\tau_m \frac{d}{dt} V_i(t) = -V_i(t) + \tau_m \sum_j J_{ij} S_j(t-d) + J_{\text{opto}} S_{\text{opto}}(t-d), \quad (\text{S.2})$$

where $V_i(t)$ is the membrane potential of neuron i , with a resting value at 0 mV. τ_m is the membrane time constant. $S_j(t) = \sum_k \delta(t - t_j^k)$ represents the spike train generated by neuron j , where t_j^k represents the individual spike times, and d is the synaptic transmission delay. The amplitude of the postsynaptic potential that induced in neuron i upon the arrival of a spike from neuron j is governed by matrix J_{ij} . $S_{\text{opto}}(t-d)$ represents the depolarizing inputs from optogenetic stimulation to a subset of excitatory neurons with synaptic weight J_{opto} . When the membrane potential $V_i(t)$ reaches the threshold, V_{th} , an action potential is emitted and the membrane potential is reset to $V_{\text{reset}} = 10$ mV. All parameters of neuron model are summarized in Supplementary Table 4.

Network model

ACC was modeled as an inhibition-dominated recurrent network (Brunel 2000), including 10 000 excitatory and 2 500 inhibitory neurons. The excitatory synapses onto inhibitory neurons (E-I) and all inhibitory connections (I-I and I-E) were static with a fixed weight. All excitatory synapses are at $J_E = 0.1$ mV, while all inhibitory synapses are at $J_I = -0.8$ mV. These connections were beforehand randomly formed with a 10% connection probability. On the other hand, the excitatory-to-excitatory (E-E) connections were grown from zero based on the rule of homeostatic structural plasticity (Diaz-Pier et al. 2016; Gallinaro and Rotter 2018). Each neuron in the network connected to a Poissonian external input at a rate of $r_{\text{ext}} = 30$ kHz. E-E connections formed during the growth period, and the network automatically entered an asynchronous-irregular state (Brunel 2000) when the equilibrium state were reached. All network parameters are listed in Supplementary Table 5.

Homeostatic structural plasticity model

Homeostatic structural plasticity rule governed the growth of presynaptic elements (boutons) and postsynaptic elements (spines) via calcium concentration, $C(t) = [\text{Ca}^{2+}]$. As showed in Equation S.3, whenever the neuron emits a spike, the intracellular calcium concentration experiences an influx by the amount β_{Ca} . Between spikes, the calcium concentration decays exponentially with time constant τ_{Ca} .

$$\frac{d}{dt}C(t) = -\frac{1}{\tau_{\text{Ca}}}C(t) + \beta_{\text{Ca}}S(t). \quad (\text{S.3})$$

We applied a linear growth rule to both pre- and postsynaptic elements (Equation S.4). Each neuron tends to maintain its activity as well as calcium concentration at a homeostatic level, called *set-point*. When the firing rate (or calcium concentration) falls below its set-point, the neuron will grow new synaptic elements and form functional synapses. When the firing rate (or calcium concentration) goes above the set-point, the neuron will randoly break synapses and retract synaptic elements. The respective counter-parts of deleted elements are added to the pool of free synaptic elements. New synapses can form only if free synaptic elements are available. Pairs of neurons can form multiple synapses between them, and each individual functional synapse has the same weight $J_E = 0.1 \text{ mV}$.

$$\frac{d}{dt}z(t) = \nu \left[1 - \frac{1}{\epsilon}C(t)\right], \quad (\text{S.4})$$

where $z(t)$ is the total number of pre- or postsynaptic elements a neuron has, ν is the growth rate, and ϵ is the target level of calcium concentration. All the parameters about the structural plasticity rule are summarized in Supplementary Table 6.

Measurements for the numerical experiments

Firing rate

The firing rate of a neuron was calculated from its spike count, in a 5 s activity recording. The mean firing rate of a population was taken to be the arithmetic mean of firing rates across neurons in the population.

Connectivity

Let (A_{ij}) be the $n \times n$ connectivity matrix of a network with n neurons. Its columns correspond to the axons, its rows correspond to the dendrites of the neurons involved. The specific entry A_{ij} of this matrix represents the total number of synapses from the presynaptic neuron j to the postsynaptic neuron i . The mean connectivity of this network is then given by $\Gamma(t) = \frac{1}{n^2} \sum_{ij} A_{ij}$, where t is the observing time point.

Supplementary Figure 1-1: No differences were observed between viral transfection and transgenic mice after the optogenetic stimulation at the behavioral and c-Fos expression level. **A** C57BL/6J mice bilaterally infected with virus vector and expressed hChR2 or Thy1-ChR2-YFP mice were used. **B-E** Both mice models received the same optogenetic stimulation protocol in ACC and behavior tests. **D** Virus-injected mice received activation again on the same day after splash test and sacrificed 1.5h later to measure c-Fos expression. **F-G** c-Fos expression was elevated in the stimulated mice (99.5% CI = [0.287, 0.687], GLMM). The c-Fos data was presented again in Figure 2G, left panel. **H-I** Mice in both approaches showed depressive-like behavior at 24 h and 48 h post-stimulation ($p = 0.012$ and $p = 0.032$ for the virus-injected group, $p = 0.018$ and $p = 0.0059$ for the transgenic group, Mann Whitney U test). The behavioral data from the transgenic mice (panel **I**) was presented again in Figure 1E-F.

Supplementary Figure 1-2: Light did not trigger changes in behavior. **A** When Thy1-ChR2-YFP mice were used in both sham and stimulated groups, we kept the light off for sham mice during stimulation. Splash test showed stimulated mice presented depressive-like behavior after four days of stimulation ($p = 0.003$, Mann Whitney U test). **B** When Thy1-ChR2-YFP mice were used in the stimulated group while gene matched wild-type mice were used for sham, we kept the light on for the sham group during stimulation. Significant difference was observed in the splash test after four days of stimulation ($p = 0.009$, Mann Whitney U test). To keep the mice gene background consistent, we used Thy1-ChR2-YFP mice throughout our study and kept light off for the sham group.

Supplementary Figure 2-1: Representative images of VGLUT1 staining at different distance to the optic fiber at 1.5 h, 24 h, and 48 h post-stimulation. High magnification images of the ipsilateral hemispheres in the middle panel were presented in Figure 2A.

Supplementary Figure 2-2: Representative images of PSD-95 staining at different distance to the optic fiber at 1.5 h, 24 h, and 48 h post-stimulation. High magnification images of the ipsilateral hemispheres in the middle panel were presented in Figure 2C.

Supplementary Figure 2-3: Representative images of neurogranin staining at different distance to the optic fiber at 1.5 h, 24 h, and 48 h post-stimulation. High magnification images of the ipsilateral hemispheres in the middle panel were presented in Figure 2E.

Supplementary Figure 3-1: Dendritic length and average dendritic diameter were not drastically altered at 24 h and 48 h post-stimulation. **A-D** Length of apical and basal dendrites were not altered by stimulation at 24 h and 48 h. **E-H** At 24 h, the average diameter of primary apical dendrites were reduced while the fourth level basal dendrites were increased by stimulation ($p < 0.05$ and $p < 0.05$). At 48 h, only the average diameter of third and fourth level apical dendrites were reduced by stimulation ($p < 0.05$ and $p < 0.05$). LMM was used for statistical analysis.

Supplementary Figure 4-1: The distribution of spine head volumes of stubby spines in the apical dendrites from each mouse at 24 h post-stimulation. The black histograms represent six sham mice respectively, while the red histograms represent stimulated mice. The fifth sham mouse and the fourth stimulated mouse did not contribute apical dendrite segments.

Supplementary Figure 5-1: Representative images of GFAP staining at different distance to the optic fiber at 1.5 h, 24 h, and 48 h post-stimulation. High magnification images of the selected regions (white squares) in the middle panel were presented in Figure 5A.

Supplementary Figure 5-2: Representative images of IBA1 staining at different distance to the optic fiber at 1.5 h, 24 h, and 48 h post-stimulation. High magnification images of the selected regions (white squares) in the middle panel were presented in Figure 5C.

Supplementary Figure 6-1: The expected expression of PSD-95 without and with homeostatic regulation. **A** If no homeostatic regulation occurred, PSD-95 expression at 1.5 h after the stimulation should remain upregulated due to accumulated effects. **B** The hypothesized time course of PSD-95 upon each stimulation session.

Supplementary Table 1: Animals and experiments used in the current study

Group	Sample size & genotype	Behavioral tests	Experiments
1.5 h-post	12 Thy1-ChR2-YFP (6 shams)	No test	IHC
24 h-post	19 Thy1-ChR2-YFP (9 shams)	Splash test	IHC
24 h-post	11 Thy1-ChR2-YFP (5 shams)	Splash test	Microinjection
48 h-post	10 Thy1-ChR2-YFP (5 shams)	NSF + Splash tests	IHC
48 h-post	12 Thy1-ChR2-YFP (6 shams)	NSF + Splash tests	Microinjection
Virus-injected*	12 transfected C57BL/6J (6 shams)	NSF + Splash tests	IHC (c-Fos only)

NSF test, novelty-suppressed feeding test; IHC, immunohistochemical staining.

* Mice were stimulated again after the splash test and sacrificed 1.5 h later.

Supplementary Table 2: Primary and secondary antibodies and their concentration used for immunohistochemical staining

Target Proteins	Primary AB	Primary AB concentration	2nd AB	2nd AB concentration
VGLUT1	Millipore, Cat#MAB5502 (RRID: AB_262185)	1 : 300	Cy5 donkey anti mouse	1 : 200
PSD-95	Millipore, Cat#MAB1596 (RRID: AB_2092365)	1 : 500	Cy5 donkey anti mouse	1 : 200
Neurogranin (Ng)	Millipore, Cat#AB5620 (RRID: AB_91937)	1 : 1000	Cy3 donkey anti rabbit	1 : 200
GFAP	US biological, Cat#2032-27N (RRID: AB_2232293)	1 : 1000	Cy3 donkey anti rabbit	1 : 200
IBA1	Wako, Cat#019-19741 (RRID: AB_839504)	1 : 1000	biotinylated anti rabbit	1 : 400
c-Fos	Synaptic Systems, Cat#226003 (RRID:AB_2231974)	1 : 1000	Cy3 donkey anti rabbit	1 : 400

AB, antibody

Supplementary Table 3: Spine classification criteria

Spine class	Criteria
stubby	$\text{length}_{\text{spine}} < 1$
mushroom	$\text{length}_{\text{spine}} < 3 \ \& \ \max(\text{width}_{\text{head}}) > \text{mean}(\text{width}_{\text{neck}}) * 2$
long-thin	$\text{mean}(\text{width}_{\text{head}}) \geq \text{mean}(\text{width}_{\text{neck}})$
filopodia	spines that are not stubby, mushroom, long-thin

Supplementary Table 4: Parameters of neuron model

τ_m	t_{ref}	V_0	V_{reset}	V_{th}
10.0 ms	2.0 ms	0.0 mV	10.0 mV	20.0 mV

Supplementary Table 5: Parameters for network model

N_E	N_I	Γ_{E-I}	Γ_{I-E}	Γ_{I-I}	J_E	J_I	r_{ext}
10 000	2 500	10%	10%	10%	0.1 mV	-0.8 mV	30 kHz

Supplementary Table 6: Parameters for the homeostatic structural plasticity model

ϵ	ν	τ_{Ca}	β_{Ca}
0.008	0.004 s^{-1}	10 s	0.0001

Supplementary Table 7: Parameters used in the model of optogenetic stimulation

Figure	r_{opto} (kHz)	f_{opto}	T_{opto} (s)	T_{int} (s)	Repetition	T_{relax} (s)
6B	1.5	50%	75	-	no	675
6E	1.5	50%	75	3 600	4	7 200