Optical TrkB activation in Parvalbumin interneurons regulates intrinsic states to orchestrate cortical plasticity

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23 **Graphical Abstract**

33 **Abstract**

34 Activation state of Parvalbumin (PV) interneurons regulates neuronal plasticity, driving the closure
 35 of developmental critical periods and alternating between high and low plasticity states in response
 36 to experience in adulthood. We now show that PV plasticity states are regulated through the
 37 activation of TrkB neurotrophin receptors. Activation of an optically activatable TrkB (optoTrkB)
 38 specifically in PV interneurons switches adult cortical networks into a state of elevated plasticity
 39 within minutes by decreasing excitability of PV neurons. OptoTrkB activation induces changes in
 40 gene expression related to neuronal plasticity and excitability, and increases the phosphorylation of
 41 Kv3.1 channels. OptoTrkB activation shifted cortical networks towards a low PV configuration,
 42 promoting oscillatory synchrony and ocular dominance plasticity. Visual plasticity induced by
 43 fluoxetine was lost in mice lacking TrkB in PV neurons. Our data suggest a novel mechanism that
 44 dynamically regulates PV interneurons configuration state and orchestrates cortical networks
 45 during adulthood.

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52 Introduction

53 Brain plasticity is a key process to allow learning and adjust maladapted networks throughout life.
 54 The brain is particularly plastic during critical periods of early postnatal life (Hensch, 2005), and
 55 transition to a state of more limited plasticity in the adult brain coincides with the maturation of
 56 GABAergic parvalbumin-expressing (PV) interneurons. The role of the PV interneuron network has
 57 been best described for the critical period in the primary visual cortex, which is typically determined
 58 through ocular dominance plasticity (Huang et al., 1999; Jiang et al., 2010; Pizzorusso et al., 2002) .
 59 PV interneuron maturation is promoted by brain-derived neurotrophic factor (BDNF) signalling
 60 (Huang et al., 1999) and the assembly of perineuronal nets (PNNs) (Pizzorusso et al., 2002),
 61 extracellular matrix components rich in chondroitin sulphate proteoglycans preferentially encasing
 62 PV interneurons. PV interneurons remain intrinsically plastic during adulthood, and external stimuli
 63 can switch the configurations of PV interneurons between plastic/immature and consolidated states
 64 as defined by low- and high-expression, respectively, of PV in these cells (Donato et al., 2013).

65 Increasing evidence has shown that a critical period-like state of plasticity can be evoked during
 66 adulthood by interventions, such as environmental enrichment (Sale et al., 2007) and
 67 antidepressant treatment (Karpova et al., 2011; Maya-Vetencourt et al., 2008; Mikics et al., 2018).
 68 Studies in the visual cortex have shown that adult ocular dominance plasticity is associated with
 69 decreased inhibitory activity thought to be driven by the PV interneuron network (Harauzov et al.,
 70 2010; Lensjø et al., 2017). However, the mechanisms underlying the switch in network states have
 71 remained a knowledge gap.

72 Considering that brain plasticity is an activity-dependent process that involves neurotrophin
 73 signaling, we hypothesized that the activity of the BDNF receptor neurotrophic receptor tyrosine

74 kinase B (TrkB) within PV interneurons could regulate metabolic processes to mediate PV plasticity
75 states during adulthood. TrkB is expressed in the majority of neurons, therefore, BDNF application
76 to study TrkB functions in specific neuron populations holds considerable limitations (Gorba & Wahle,
77 1999; Kuczewski et al., 2009).

78 Here, we modified and optimized an optically activatable TrkB (optoTrkB) (Chang et al., 2014; Fenno
79 et al., 2014) that can be expressed and activated in PV interneurons in a cre-dependent manner. This
80 tool then allowed us to study the mechanisms underlying TrkB activity in PV interneurons and its
81 effects on visual cortex plasticity as a well-defined model network. This optogenetic approach differs
82 substantially from traditional channelrhodopsin experiments as the activation of optoTrkB allows us
83 to control a relatively slowly developing, physiologically relevant molecular pathway.

84 We show here that optoTrkB activation specifically in PV interneurons in the primary visual cortex
85 during 7 days of monocular deprivation (MD) was sufficient to reinstate ocular dominance plasticity.
86 More strikingly, a single 30 seconds blue light stimulation of optoTrkB in PV interneurons resulted in
87 LTP inducibility within 30 minutes and enhanced oscillatory synchrony in the visual cortex of adult
88 mice. Subsequent single-nuclei sequencing revealed that activation of optoTrkB in PV interneurons
89 rapidly induces changes in the expression of genes related to neuronal excitability and plasticity,
90 such as *Grin1* and *Grik3*, subunits of the NMDA and kainate receptor, respectively. Surprisingly, also
91 *PV* expression itself appeared to be regulated through TrkB activation. Consistently, we found a rapid
92 decrease in PV intrinsic excitability and spontaneous excitatory postsynaptic current (sEPSC)
93 frequencies after optoTrkB activation, as well as increased phosphorylation of the PV specific Kv3.1
94 channels. In line with our single-nuclei sequencing data, we found that TrkB activation in PV
95 interneurons dynamically regulates PV and PNN configuration states, resetting the neuronal network
96 into a plastic, immature-like state. Conversely, deleting TrkB from PV interneurons blocked these

effects when induced pharmacologically using fluoxetine treatment. These findings demonstrate that TrkB activation in PV interneurons rapidly orchestrates cortical network plasticity by regulating intrinsic states and provide new evidence for its role in intracortical inhibition and plasticity modes during adulthood.

Results

TrkB activation in PV interneurons induces visual cortex plasticity

Using BDNF to study cell type-specific functions of TrkB has critical caveats as it can activate TrkB receptors in any neuron expressing it. To be able to specifically activate TrkB in PV interneurons, we optimized and modified an optically activatable TrkB (optoTrkB) where addition of a plant-derived photolyase homology region (PHR) domain into the C-terminus of full-length TrkB mediates light-induced dimerization of TrkB monomers (Chang et al., 2014) (Fig. 1A) (Figure S1A). We infected the V1 of adult mice specifically expressing cre recombinase in PV interneurons (PV-cre) with a lentivirus expressing a double-floxed inverted open reading frame of opto-TrkB (DIO-optoTrkB) to specifically express optoTrkB in PV interneurons (Figure S1B). Acute light activation of optoTrkB in V1 resulted in increased phosphorylation of TrkB and CREB (Figure S2A-S2E), suggesting successful activation of optoTrkB and downstream signalling *in vivo*.

We then infected the V1 of adult PV-cre mice with DIO-optoTrkB lentivirus and subjected these mice to the standard protocol of ocular dominance (OD) plasticity induced by 7 days of monocular deprivation (MD) (see Material&Methods). As expected, mice infected with the lentivirus but having their transparent skull (Steinzeig et al., 2017) painted black to prevent light stimulation, failed to show any OD plasticity. In contrast, when the DIO-optoTrkB infected adult primary visual cortex (V1)

119 was stimulated twice daily for 30 seconds by blue light through a transparent skull (Fig. 1B) during 7
 120 days of MD, we observed a shift in ocular dominance index (ODI) towards the non-deprived
 121 ipsilateral eye (Fig.1C). The shift persisted for a week in the absence of visual deprivation or light
 122 stimulation (Fig. 1C). We then closed the eye that had been left open during the first MD session
 123 and again exposed the cortex to light twice daily for 7 days and again observed a shift towards the
 124 non-deprived eye (Fig. 1C). Light stimulation alone without DOI-optoTrkB virus infection during 7
 125 days of MD had no effect on visual cortex plasticity (Figure. S2F).

126 A shift in ocular dominance takes several days to occur but we reasoned that direct activation of
 127 optoTrkB through light stimulation might have more immediate effects. We therefore studied the
 128 induction of long-term potentiation (LTP) as another proxy for visual cortex plasticity. LTP occurrence
 129 through theta-burst stimulation (TBS) in V1 is normally restricted to critical periods (Kirkwood et al.,
 130 1996). While TBS stimulation (Kirkwood & Bear, 1994) of acute V1 slices expressing optoTrkB in PV
 131 interneurons but kept in darkness did not induce LTP in slices, a robust LTP was induced in slices
 132 exposed for 30 seconds to blue light 30 minutes before the TBS stimulation (Fig.1D). These results
 133 suggest that direct activation of TrkB receptors in PV interneurons promotes excitatory transmission
 134 and plasticity within minutes.

135 Plasticity in the visual cortex is controlled locally by synchronized neuronal oscillations, which are
 136 regulated by the activity of PV interneurons (Galuske et al., 2019; Lee et al., 2012). *In vivo* local field
 137 potential (LFP) recordings of neuronal activity from V1 revealed a progressive increase in the
 138 broadband (4–112 Hz) LFP power in response to a 30-second long optoTrkB activation (Fig. 1E). The
 139 broadband magnitude at 100 minutes after stimulation showed a significant increase when
 140 compared to the pre-stimulation baseline. Furthermore, no increase in the broadband magnitude
 141 was observed in optoTrkB infected control animals stimulated with infrared light (780nm) (Fig. 1F,

H). We also found increases in the magnitude at the separate frequency bands, particularly in alpha (α) and gamma (γ) range at 100 minutes after optoTrkB activation compared to baseline (Fig. 1G). Together, these results suggest that optoTrkB activation in PV interneurons promotes synchronized oscillations, which renders the network state of the visual cortex responsive and permissive for plasticity.

TrkB regulates plasticity-related genes in PV interneurons

To identify the molecular mechanisms regulated by optoTrkB activation in PV interneurons, we performed single nuclei RNA sequencing (snRNA-seq) of visual cortical samples from DIO-optoTrkB-TdTomato (Figure S1C and S1D) transfected PV-cre mice 60 minutes after a 30-second light stimulation (Figure S3A). Single cell RNA sequencing was not feasible due to potential light-induced optoTrkB activation during cell extraction and processing (Figure S3B). Integrated data analysis (see Material&Methods) identified 18 clusters of neurons, including a cluster containing PV interneurons (Fig. 2A; Figure S4 and S5; Table S1-S3), which was further divided into four PV interneuron clusters using PCA analysis (Fig. 2B and 2C; Figure S6 and S7; Table S4). *Arc*, a known downstream signal of TrkB (Messaoudi et al., 2002, 2007; Yin et al., 2002; Ying et al., 2002), was only up-regulated in PV cluster 3 of the light-stimulated group, along with other immediate early genes, such as, *Jun* and *Egr1* (Fig. 2E), indicating that this cluster harbours optoTrkB transfected PV cells. In this cluster, genes related to dendrite morphogenesis, synaptic plasticity, and regulation of excitatory postsynaptic currents (EPSC) were differentially regulated in the light-exposed group (Fig. 2D) (Table S5). Strikingly, we found a decrease in *PV* expression itself among them (Fig. 2E), and this decrease was confirmed by qPCR (Figure S10). Particularly interesting is the downregulation of *Grin1* and *Grik3* that code for subunits of N-methyl-D-aspartate receptor (NMDA) and kainate receptors, respectively, and

165 *Homeobox protein cut-like 2 (Cux2)* (fold value, 0.63, Table S6), all being essential components of
 166 excitatory synapses (Fig. 2E). Furthermore, *Leucine Rich Repeat And Ig Domain Containing 2 (Lingo-*
 167 *2)* that is broadly expressed in neurons was reduced after optoTrkB activation (fold value, 0.46, Table
 168 S6). Finally, we found an upregulation of the insulin-like growth factor 1 (IGF-1) receptor (*Igf1r*)
 169 expression (fold value, 1.29, Table S5), which shares many components with the BDNF pathway
 170 (Zheng & Quirion, 2004)

171 These results suggest that optoTrkB activation in PV interneurons changes the gene expression
 172 profile to particularly regulate the PV intrinsic properties.

173

174 **TrkB activation in PV interneurons reduces cell excitability**

175 We next sought to validate the functional effects of optoTrkB activation on the intrinsic properties
 176 of PV interneurons. We therefore obtained patch clamp recordings from optoTrkB-positive PV
 177 interneurons in the V1 co-expressing TdTomato (Figure S1C and S1D). OptoTrkB was activated 10-60
 178 minutes before the recordings using a 30-second blue light stimulation. We first recorded the
 179 intrinsic excitability by injecting current steps ranging from -100 to 600 pA. Strikingly, the intrinsic
 180 excitability of PV cells in optoTrkB stimulated slices was significantly lower 30-60 min after activation
 181 when compared to the non-activated controls (Fig. 3A and 3F) (Table S6A), and was accompanied by
 182 trends towards increased action potential (AP) half-width (Fig. 3B and 3F). We subsequently
 183 recorded spontaneous EPSCs (sEPSC) and found that the frequency of sEPSCs was also decreased in
 184 optoTrkB activated PV interneurons compared to controls (Fig. 3C and 3F), confirming functional
 185 changes in excitatory transmission onto PV interneurons. Neither the sEPSC amplitudes nor the
 186 frequency or amplitudes of spontaneous inhibitory postsynaptic currents (sIPSC) were changed

(data not shown), suggesting that optoTrkB activation specifically affects excitatory inputs.

Potassium currents are known to regulate intrinsic excitability of neurons and Kv3 channels that are highly expressed in cortical PV interneurons regulate the fast-spiking properties (Chow et al., 1999; Du et al., 1996). Furthermore, Kv3.1 channels are directly inhibited by phosphorylation through PKC (Song & Kaczmarek, 2006), a downstream target of TrkB signaling. We therefore hypothesized that optoTrkB activation could result in enhanced phosphorylation or reduced expression of Kv3.1 channels. First, we immunohistochemically examined the expression intensities of phospho-Kv3.1 within PV interneurons and found that optoTrkB activation resulted in increased phospho-Kv3.1 expression in PV interneurons in visual cortical slices (Fig. 3D and 3E), which is consistent with reduced excitability. We then used qPCR with tissue samples from the V1 of optoTrkB-infected PV-cre mice 15 min, 30 min or 60 min after light stimulation to measure mRNA expression of Kv3.1 and Kv3.2 channels. The expression of Kv3.2 mRNA was significantly reduced and there was a progressive trend toward reduction for Kv3.1 mRNA ($p = 0.0923$) at 60 min after light stimulation (Fig. 3G and 3H). These data suggest that optoTrkB activation inhibits Kv3.1 potassium channels through phosphorylation and regulates Kv3.1 and Kv3.2 mRNA expression, thereby reducing the intrinsic excitability of PV interneurons.

203

204 **TrkB activation regulates PV-plasticity**

PV expression itself is plastic and regulated by experience (Donato et al., 2013, 2015; Karunakaran et al., 2016), low PV expression being associated with plastic and high PV with consolidated networks. In addition, PV interneurons become gradually enwrapped by PNN during critical periods (Jiang et al., 2005) and PNN removal during adulthood reinstates a juvenile-like plasticity and reduces

intracortical inhibition (Lensjø et al., 2017; Pizzorusso et al., 2002). As our snRNA-seq data revealed that optoTrkB activation decreases PV expression, we hypothesized that optoTrkB activation could directly mediate PV and PNN plasticity states. Indeed, activation of optoTrkB in the V1 PV interneurons for 30 seconds twice daily for 7 days reduced the PV intensities particularly in the subgroup of PV interneurons expressing high levels of PV (Fig. 4A, B) and the PNN intensities were also reduced in the same high-PV subgroup (Fig. 4C), suggesting a switch towards a low PV expressing network state. We next reasoned that a reduction in PNN intensities could correlate with a decrease in PNN expression and quantified the numbers of PNNs. We found that optoTrkB activation reduced the numbers of PNN-positive (PNN⁺), PV/PNN-double-positive (PV⁺PNN⁺) cells, and PV⁺PNN⁺ cells within the PV population (Fig. 4D-G), suggesting that PV interneurons switch to a critical period-like state after optoTrkB activation.

TrkB activation in PV interneurons is necessary for induction of visual cortex plasticity

Together, our findings demonstrate that TrkB activity in PV interneurons is sufficient to render the cortical network towards a plastic configuration state. We wondered, however, whether TrkB activity is also necessary for it. To test this hypothesis, we used a “loss-of-function” approach using conditional heterozygous mice with PV-specific TrkB knockout (hPV-TrkB CKO) and pharmacologically induced plasticity using chronic fluoxetine treatment. Using again the standard protocol for ocular dominance plasticity, we confirmed our previous findings that chronic fluoxetine treatment induced a shift of ocular dominance in combination with MD in the adult V1 of wild-type mice (WT) (Maya-Vetencourt et al., 2008; Steinzeig et al., 2017). However, no OD shift was seen in hPV-TrkB CKO mice in response to MD during chronic fluoxetine treatment (Fig. 5A). Consistently, chronic fluoxetine treatment also permitted LTP induction after TBS stimulation, as previously reported (Maya-

232 Vetencourt et al., 2008), but this effect was absent in hPV-TrkB CKO mice (Fig. 5B). These data suggest
233 that TrkB activation in the visual cortex is necessary for fluoxetine-induced adult plasticity.

234

235 **TrkB expression is necessary for fluoxetine-induced changes in PV intrinsic properties**

236 We then sought to validate whether the underlying mechanisms of fluoxetine-induced plasticity are
237 similar to those induced by optoTrkB activation. Indeed, similarly to what was observed after
238 optoTrkB activation in PV interneurons, chronic fluoxetine treatment reduced intrinsic excitability
239 (Fig. 6A, Fig.S7) (Table S6B), increased AP half-width (Fig. 6B, Fig.S7) and produced a trend towards
240 a decrease in sEPSC frequency ($p = 0.5397$) (Fig. 6C, Fig.S7) in PV interneurons in WT mice, but none
241 of these responses to fluoxetine were seen in hPV-TrkB CKO mice.

242 Finally, again in alignment with responses to optoTrkB, we found that chronic fluoxetine treatment
243 reduced PV and PNN intensities among the high PV expressing population (Fig. 6D, E) and PNN⁺
244 abundance within the PV population in WT (Fig. 6F, G, H) but not in hPV-TrkB CKO mice (Fig. 6F-H).
245 Taken together, these results strongly suggest that TrkB activation in PV interneurons is both
246 necessary and sufficient to regulate PV intrinsic properties and switch the visual cortex to a state of
247 high plasticity.

248

249

250 Discussion

251 Our data show that TrkB signalling powerfully regulates the intrinsic activity of PV interneurons and
 252 thereby orchestrates cortical plasticity. TrkB activation in PV interneurons switches the intrinsic state
 253 of PV interneurons towards low excitability resulting in disinhibition and induction of critical period-
 254 like plasticity in cortical networks. Conversely, inhibition of TrkB signalling prevents induction of
 255 plasticity produced by fluoxetine treatment. Using optoTrkB as a methodological advancement we
 256 could show that TrkB particularly affects the expression of genes involved in synaptic transmission
 257 and intrinsic excitability as well as *PV* itself. The functional consequences of these changes are
 258 expressed in an increase of excitatory transmission, synchronized oscillations and visual cortex
 259 plasticity.

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261 TrkB activation in PV interneurons induces plasticity in adult brain

262 We previously demonstrated that chronic fluoxetine treatment can reactivate ocular dominance
 263 plasticity in the adult visual cortex (Maya-Vetencourt et al., 2008; Steinzeig et al., 2017). This effect
 264 was accompanied by a reduction in intracortical GABAergic transmission as measured by decreased
 265 extracellular basal GABA levels; induction of LTP after TBS stimulation and increased BDNF protein
 266 levels, strongly suggesting a regulatory role of intracortical inhibition in adult visual cortex plasticity
 267 (Maya-Vetencourt et al., 2008).

268 Although PV interneurons cover only a small percentage of the total neuronal population, their
 269 extensive axonal arborization enable strong inhibitory control over pyramidal cells (Hu et al., 2014).
 270 As BDNF promotes the maturation of PV cells during critical periods, we hypothesized that TrkB
 271 activation in PV interneurons could also affect plasticity processes during adulthood. Considering

272 that administration of BDNF and fluoxetine activate all TrkB receptors expressed in the majority of
 273 neurons, we developed a cre-dependent optoTrkB to circumvent this problem and to be able to
 274 specifically activate TrkB signaling in PV interneurons only. Strikingly, while BDNF-induced TrkB
 275 activation during critical periods promotes the maturation of PV interneurons (Huang et al., 1999),
 276 TrkB activation in PV interneurons during adulthood reinstates a critical period-like plasticity
 277 machinery through rejuvenating these interneurons. These findings suggest that the plasticity
 278 machinery remains intact in the adult brain but needs the right tools to be reactivated.

279

280 **Differential effects of TrkB activation in pyramidal neurons and interneurons?**

281 Reactivation of critical period-like plasticity in the adult brain is associated with reduced cortical
 282 inhibition and subsequent disinhibition of cortical networks (Hensch, 2005). BDNF has been shown
 283 to strongly promote neuronal activity, excitability (Figurov et al., 1995; Levine et al., 1995, 1998) and
 284 LTP (Kang & Schuman, 1994; Minichiello, 2009) in hippocampal and cortical excitatory neurons. In
 285 contrast, we found that optical activation of TrkB specifically in PV interneurons decreases
 286 excitability in PV cells and promotes field LTP in the visual cortex, indicating an increase in network-
 287 wide excitatory transmission and dynamics. These findings suggest that activation of TrkB produces
 288 differential effects on the cell excitability, increasing and decreasing it in pyramidal neurons and
 289 interneurons, respectively. Importantly, however, from the point of view of cortical networks, these
 290 differential effects on excitability cooperatively drive increased network activity, through enhanced
 291 excitability of pyramidal neurons and reduced activity of inhibitory neurons.

292

293 **Activation of TrkB mediates intrinsic changes in PV interneurons to enhance excitatory**

294 **transmission in the visual cortex**

295 The hallmark of PV interneurons is their high-frequency firing, which is enabled by the high
 296 expression of voltage-gated Kv3 channels (Chow et al., 1999; Hu et al., 2014). Kv3 channels are
 297 characterized by their fast deactivation during membrane repolarization, enabling sustained high-
 298 frequency firing. A decrease in Kv3 channel expression as well as their inhibition through PKC
 299 mediated phosphorylation is known to reduce cell excitability (Rudy et al., 1999; Song & Kaczmarek,
 300 2006). PKC is a downstream target of TrkB signaling (Minichiello, 2009) and our data show that TrkB
 301 activation in PV interneurons results in enhanced phosphorylation of Kv3.1 and decreased
 302 expression of Kv3.1 and Kv3.2 channels, demonstrating control of TrkB signaling over PV intrinsic
 303 excitability. The decreased firing frequency of PV cells in turn enhances excitatory transmission
 304 necessary for visual cortex plasticity. The maturation of PV cells throughout critical period is
 305 concurrent with an increase in the fast-spiking firing frequency and an increase in Kv3.1 expression
 306 (Du et al., 1996; Plotkin et al., 2005). Interestingly, the fast-spiking properties of PV interneurons
 307 fails to develop in BDNF null mice (Itami et al., 2007). Considering that Kv3.1 channels are specifically
 308 expressed in high-frequency firing neurons, this could account for the differential effects of TrkB
 309 actions in pyramidal cells and PV interneurons.

310

311 **OptoTrkB-induced changes in gene expression in PV interneurons**

312 Using snRNA-seq, we found genes that were differentially regulated after optoTrkB activation in PV
 313 interneurons. Particularly the expression of genes related to excitatory transmission, synaptic
 314 plasticity and excitability was affected. *Grin1* and *Grik3* are subunits of the NMDA and kainate
 315 receptor, respectively, and these essential components of excitatory synapses were reduced by

316 optoTRKB activation. Knocking out *Grin1* in PV interneurons enhances network oscillations,
 317 particularly theta and gamma frequencies (Korotkova et al., 2010). The specific function of the *Grik3*
 318 subunit is not fully characterized, but kainate receptors have been shown to regulate both the
 319 maturation and excitability of GABAergic interneurons (Jack et al., 2019; Segerstråle et al., 2010), as
 320 well as the generation of synchronized activity at the level of neuronal networks (Bartos et al., 2007).
 321 As both NMDA and kainate receptors are known to mediate and modulate excitatory synaptic
 322 transmission (Zhang et al., 2013), the downregulation of *Grin1* and *Grik3* might therefore contribute
 323 to the changes in intrinsic properties of PV interneurons. In addition, we observed a decrease of
 324 *Cux2*, deletion of which reduces the amplitude and lowers the frequency of mEPSC (Cubelos et al.,
 325 2010). Furthermore, *Lingo-2* that is broadly expressed in neurons was reduced after optoTrkB
 326 activation, and the *Lingo* family members act as negative regulators of TrkB signalling (Meabon et
 327 al., 2016). Finally, we found an upregulation of the *Igf1r* expression. IGF-1 is another neurotrophic
 328 factor and shares components with the intracellular pathway of BDNF (Zheng & Quirion, 2004).
 329 Studies from the visual cortex have shown that IGF-1 reactivates ocular dominance plasticity and
 330 reduces GABAergic transmission in the adult visual cortex (Maya-Vetencourt et al., 2012).
 331 Upregulation of the IGF-1 receptor could render PV interneurons more responsive to IGF-1, thereby
 332 further contributing to a decrease in GABAergic transmission.

333

334 **Activation of optoTrkB regulates PV plasticity**

335 Exciting research suggests that PV interneurons show intrinsic plasticity and can change between
 336 high and low plasticity states when exposed to different environmental conditions (Donato et al.,
 337 2013). For example, environmental enrichment that is known to promote neuronal plasticity,
 338 induces a low PV-expressing network state. In contrast, contextual fear conditioning that is

339 associated with network consolidation, shifts the PV network into a high-PV expressing state (Donato
340 et al., 2013). Additionally, PV expression progressively increases throughout development and could
341 therefore also account for the reduction in brain plasticity during adulthood (Donato et al., 2013;
342 Umemori et al., 2015). Interestingly, TrkB activation in PV interneurons can regulate PV plasticity
343 states, and this effect may be mediated through direct regulation of PV mRNA expression as
344 confirmed by single nuclei sequencing and qPCR. As PV is a Ca^{2+} buffer, a low PV expressing state
345 could also directly regulate PV interneuron firing (Eggermann & Jonas, 2012; Hu et al., 2014)

346 In the hippocampus, PV intensity is generally higher in PV cells enwrapped by PNNs and PV cells with
347 weak staining intensity often lack PNNs (Yamada et al., 2015). The maturation of the PV network is
348 associated with the formation of PNNs (Pizzorusso et al., 2002). Enzymatic digestion of PNNs
349 restores plasticity in adulthood (Lensjø et al., 2017; Pizzorusso et al., 2002) and results in a decrease
350 in PV intensity and PV mRNA levels, suggesting a correlation between PNN expression and PV
351 configuration states (Yamada et al., 2015). In fact, a recent study by Devienne et al. demonstrates
352 that transient electrical silencing of visual cortical PV interneurons induces a regression of PNN
353 (Devienne et al., 2019), indicating a causal relationship between PV activity and the accumulation of
354 PNNs. Our data suggest that TrkB activation within PV interneurons directly regulates PV expression
355 and leads to the reduction in PNN levels, thereby contributing to the plasticity state of PV neurons.

356 In conclusion, we show that TrkB receptor activation within PV interneurons is sufficient and
357 necessary to rapidly reduce their excitability and plasticity state, thereby changing cortical network
358 dynamics. Hence, TrkB activity in PV cells dynamically controls network rewiring, learning and
359 consolidation. Although we used the visual cortex as a model network, these mechanisms might be
360 extrapolated to other brain areas, such as the hippocampus and amygdala, and therefore aids in the
361 understanding of the pathophysiology of neuropsychiatric disorders and the rational design of

362 clinical interventions.

363

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Declaration of competing interests

EC has received a lecturer fee from Jansen Cilac.

Author contributions

F.W., J.U. and E.C. conceived and designed the project. F.W. performed the experiments. M.V. analyzed the in vivo electrophysiology recordings under the supervision of S.P.. G.D. helped with the Western Blot experiments. M.L. cut the brains on the vibratome and helped with

386 immunohistochemistry. E.J., S.M., and J.U. helped with single nuclei sequencing experiments and
 387 data analysis. A.S. helped with the behavioral experiment using hPV-TrkB CKO mice combined with
 388 fluoxetine treatment. J.H. analyzed the PV/PNN intensities. S.K., C.R., J.E., M.R., S.K., T.T. and S.L.
 389 provided supervision and support during the electrophysiological experiments. F.W., J.U. and E.C.
 390 wrote the manuscript.

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Figures

Fig. 1. TrkB activation in PV interneurons induces visual cortex plasticity.

(A) Structure of optoTrkB conjugated to a light reactive PHR domain, which dimerizes optoTrkB upon blue light exposure. (B) Experimental timeline of the shift in ocular dominance paradigm with optoTrkB. OptoTrkB was stimulated with blue light twice daily (30 seconds) for 7 days during MD. (C)

MD of the contralateral eye induces a shift in ocular dominance when combined with stimulation of optoTrkB (IOS I-II) (n = 6-8 animals/group). Two-way ANOVA with Sidak's post-hoc test comparing the ODI of IOSI and II (control, p = 0.9094; light, p = 0.001). The shift is preserved if the eyes are not deprived (IOS II-III). Two-way ANOVA with Sidak's post-hoc comparing the ODI of IOS II with III (control, p = 0.4985; light, p = 0.6915). Stimulation of optoTrkB reverses shift in ocular dominance when combined with MD of the ipsilateral eye (IOS III-IV) (n = 6-7 animals/group; one mouse excluded due to corneal infection). Two-way ANOVA with Sidak's post-hoc test comparing ODI of IOS III and IV (control, p = 0.9094; light, p = 0.001). (D) LTP recordings from layer II/III in the V1. Theta-burst stimulation (TBS) results in LTP in slices where optoTrkB has been activated for 30 seconds 30 minutes prior to LTP induction, but not in control slices. Two-way ANOVA (p < 0.0001). n = 6-7 recordings/group randomly recorded from 5 animals. (E) The broadband (4–112 Hz) LFP power (blue: optoTrkB activated at 470 nm, black: control wavelength at 780 nm) averaged over animals (light = 5, control = 4) for each 20 minute recording session as a function of time after light stimulation. LFP power shows a significant increase over time (Regression analysis; Control: $y = 0.04 [\%/min] \cdot x [min] + 100.25$, $R^2 = 0.7573$, $p < 0.0001$; Light: $y = 0.09 [\%/min] \cdot x [min] + 101.01$, $R^2 = 0.8240$, $p < 0.0001$). (F) Broadband LFP power increase from baseline period (left plot of each half) to 100-minute after blue light (470nm) stimulation (right plot of each half) was approximately twice as strong after optoTrkB activation (blue, difference in medians: d = 10.72) than after stimulation with control wavelength (780nm) (black, difference in medians: d = 4.90). A Welch's t-test confirmed that the LFP power in the 100-minute condition bins after optoTrkB activation is higher than after control stimulation (Welch's t-test, t = 21.60, p < 0.0001). (G) Changes in frequency bands 100 min after light stimulation compared to baseline. (H) Wavelet spectra of control (left) and light stimulated (right) mice. Color scale. Flx, fluoxetine; ODI, ocular dominance index; MD, monocular deprivation; V1, primary visual cortex; PHR, photolyase homology region; EXD, extracellular domain; TMD, transmembrane domain; TK, tyrosine kinase domain. Bars represent means $\pm$ SEM. ** p < 0.01 *** p < 0.001; **** p < 0.0001; Bars represent means $\pm$ SEM, **** p < 0.00001.

Fig. 2. TrkB regulates plasticity-related genes in PV interneurons.

(A) Two-dimensional Uniform Manifold Approximation and Projection (UMAP) plots based on 3,000 differentially expressed (DE) genes for 10,706 cells. After PCA analysis followed by clustering, the cells were grouped into 18 subpopulations including interneurons. (B) UMAP plots based on 3,000 DE genes for selected 586 cells clustered to PV interneurons. After PCA analysis, the cluster was further divided into 4 groups (PV_0, 1, 2, 3). (C) Markers differently expressed in each cluster. PV_3 cluster include both Chandelier (CHC) and basket (BS) cells. (D) Gene ontology (GO) analysis on differentially expressed (DE) genes between non-light-exposed and light-exposed samples in cluster 3_3. (E) Selected DE genes of immediate early genes (IEGs) and relevant genes to decreased excitation in cluster PV_3.

Fig. 3. TrkB activation in PV interneurons reduces cell excitability.

(A) Whole cell patch clamp recordings of intrinsic excitability of optoTrkB-transfected PV cells. Activation of optoTrkB decreases intrinsic excitability of PV cells at 30-60 minutes after light stimulation. Two-way ANOVA with Tukey's post-hoc test comparing control with light stimulation (control vs. 30-60 min, $p = 0.0354$). $n = 6-8$ recordings/group randomly recorded from 14 animals. (B) Trends towards larger AP half-width 30-60 min after light stimulation. One-way ANOVA with Bonferroni's post-hoc test (30-60 min, $p = 0.1622$). (C) sEPSC frequency in PV interneurons is lower at 10-30 minutes after light stimulation as compared to controls. One-way ANOVA with Bonferroni's post-hoc test (10-30 min, $p = 0.0233$; 30-60 min, $p = 0.0261$). $n = 6-8$ recordings/group randomly recorded from 14 animals. (D) Light stimulation of optoTrkB results in increased intensity of phospho-Kv3.1 staining in PV interneurons in layer II/III of the V1. Unpaired t-test. $n = 6$ animals/group; 7 days twice daily light stimulation. (E) Representative images of PV, phospho-Kv3.1 and merged immunohistochemistry staining. (F) Representative traces used to estimate intrinsic excitability (top row, 150 pA), AP half-width (middle row) and sEPSC frequency (lower row). (G-H) qPCR quantification of Kv3.1 and Kv3.2 mRNA. Kv3.1 ($p = 0.0923$) (G) and Kv3.2 (H) mRNA levels are

489 decreased 60 min after light stimulation measured by qPCR. qPCR measurements of control, 15
 490 minutes, 30 minutes and 60 minutes after light stimulation. AP, action potential; spontaneous
 491 excitatory postsynaptic current, sEPSC; Bars represent means $\pm$ SEM. * $p < 0.05$; *** $p < 0.0001$; ****
 492 $p < 0.00001$

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Fig. 4. TrkB activation regulates PV-plasticity.

(A) Representative immunohistochemical images of PV and PNN expression in layer II/III of the V1. (B-D) Number of cells expressing PV and/or PNN after optoTrkB activation (n = 6 animals/group). (B) Chronic light stimulation of optoTrkB significantly reduced numbers of PNN positive cells (Unpaired t-test; p 0.0012), (C) PV/PNN double-positive cells (Unpaired t-test; p = 0.0021), and (D) PV cells expressing PNN (Unpaired t-test; p = 0.0156). (E) Representative images of low, intermediate-low, intermediate-high and high PV-expressing cells, where PV interneurons are categorized according to PV intensity. (F) Image analysis on PV intensities. Chronic stimulation of optoTrkB significantly reduces the PV staining intensities of high PV expressing cells. Two-way ANOVA with Bonferroni's post-hoc test comparing control vs. light stimulated samples (p = 0.0034). (G) Image analysis on PNN intensities. Activation of optoTrkB decreases PNN intensities in high PV-expressing cells. Two-way ANOVA with Bonferroni's post-hoc test comparing control vs. light stimulated samples (p = 0.0003). Bars represent means $\pm$ SEM. * p < 0.05; *** p < 0.0001; **** p < 0.00001

Fig. 5. TrkB activation in PV interneurons is necessary for visual cortex plasticity

(A) ODI after chronic Flx treatment in WT and hPV-TrkB CKO mice. Flx permits MD to shift the ODI towards the non-deprived eye in the V1 of WT mice, but fails to do so in hPV-TrkB CKO mice. Two-way ANOVA with Sidak's post-hoc test comparing the ODI of IOS I and II. WT water, $p = 0.7311$; WT Flx $p < 0.0001$; CKO water, $p = 0.9621$; CKO Flx, $p = 0.5048$. $n = 5 - 10$ animals/group. (B) LTP recordings from layer II/III in the V1 in WT and CKO mice after Flx treatment. TBS induces LTP only in WT mice treated with Flx. Two-way ANOVA with Tukey's post-hoc test comparing treatment in WT and treatment in CKO mice (WT water vs. WT Flx, $p = < 0.0001$; CKO water vs. CKO Flx, $p = 0.8871$). $n = 7-12$ recordings/group from 3 animals.

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Fig. 6. TrkB expression is necessary for fluoxetine-induced changes in PV intrinsic properties

(A) Whole cell patch clamp analysis of WT and hPV-TrkB CKO mice after chronic Flx treatment. The intrinsic excitability is reduced after Flx treatment in WT mice but not in hPV-TrkB CKO mice. Two-way ANOVA with Tukey's post-hoc test comparing Flx treatment in WT and CKO mice (WT water vs. WT Flx, $p < 0.0001$; CKO water vs. CKO Flx, $p = 0.7068$). $n = 7-12$ recordings/group from 3 animals. (B) Flx treatment increases AP half-width in WT mice but not in hPV-TrkB CKO mice. Two-way ANOVA with Tukey's post-hoc test comparing Flx treatment in WT and CKO mice (WT water vs. WT Flx, $p = 0.03$; CKO water vs. CKO Flx, $p = 0.6672$). (C) Flx treatment induces a trend towards reduced frequency of sEPSC in WT ($p = 0.5397$) but not in hPV-TrkB CKO mice ($p = 0.9866$). Two-way ANOVA with Tukey's post-hoc test. (D) Flx treatment decreases PV intensities in high PV expressing cells in WT but this effect is not seen in hPV-TrkB CKO mice. Two-way ANOVA with Tukey's post-hoc test comparing Flx treatment in WT and hPV-TrkB CKO mice (WT water vs. WT Flx, $p = 0.0325$; CKO water vs. CKO Flx, $p = 0.9838$). (E) Flx treatment reduces PNN intensities in intermediate-high and high PV expressing cells only in WT mice but fails to do so in hPV-TrkB CKO mice. Two-way ANOVA with Tukey's post-hoc test comparing Flx treatment in WT and hPV-TrkB CKO mice (WT water vs. WT Flx, intermediate-high, $p = 0.0014$; high PV, $p = 0.0325$; CKO water vs. CKO Flx, intermediate-high, $p = 0.3217$; high PV, $p = 0.6112$). $n = 6-10$ animals/group. (F) Flx-treated WT mice show a trend towards lower numbers of PNN ($p = 0.07$). (G) Flx-treated WT mice have significantly lower percentages of PV interneurons also expressing PNNs, but this effect is abolished in hPV-TrkB CKO mice. Two-way ANOVA with Tukey's post-hoc test comparing Flx treatment in WT and CKO mice (WT water vs. WT Flx, $p = 0.0001$; CKO water vs. CKO Flx, $p = 0.6462$). (H) Representative immunohistochemical images of PV and PNN expression in layer II/III of Flx-treated WT and hPV-TrkB CKO mice. Number (B, C) and intensity (D, E) of cells expressing PV and/or PNN by imaging analysis after Flx treatment. Bars represent means $\pm$ SEM. ** $p < 0.01$; *** $p < 0.001$; **** $p < 0.0001$

604 **Material and Methods**

605 EXPERIMENTAL MODEL AND SUBJECT DETAILS

606 All animal procedures were done according to the guidelines of the National Institutes of Health
607 Guide for the Care and Use of Laboratory Animals and were approved by the experimental Animal
608 Ethical Committee of Southern Finland. Female mice (PV-cre, WT or hPV-TrkB CKO) older than 110
609 days were group housed with *ad libitum* access to food and water. For the shift in ocular dominance
610 paradigm, the mice underwent transparent skull surgery and were monocularly deprived for 7 days.

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613 **Mice**

614 Heterozygous mice with TrkB deletion specifically in PV⁺ interneurons (PV-hTrkB mice; PV^{pvr/wt},
615 TrkB^{flx/wt}) were produced by mating females of heterozygous PV specific Cre line (PV^{pvr/wt}) (Pvalb-
616 IRES-Cre, JAX: 008069, Jackson laboratory) (Hippenmeyer et al., 2005) and male of homozygous
617 floxed TrkB mice (TrkB^{flx/flx}) (Minichiello et al., 1999). Single floxed mice (TrkB^{flx/wt}) served as control.
618 For patch-clamp electrophysiology, females of the homozygous PV Cre line (PV^{pvr/pvr}) were crossed
619 with males harboring a homozygous TdTomato indicator allele (Rosa26^{TdT/TdT}) (Ai14, JAX: 007914,
620 Jackson laboratory, Bar Harbor, ME) (Madisen et al., 2010) and heterozygous floxed TrkB allele
621 (TrkB^{flx/wt})(Minichiello et al., 1999) to reproduce compound transgenic mice with TrkB deletion and
622 TdT expression specifically in PV interneurons (PV^{pvr/wt}, TrkB^{flx/wt}, Rosa26^{TdT/wt}) (hPV-TrkB CKO) and
623 wild-type allele in TrkB (PV^{pvr/wt}, TrkB^{wt/wt}, Rosa26^{TdT/wt}) (WT). All of the parental strains were back-
624 crossed with C57BL/6J for more than six generations. To assure the complete closure of the critical
625 periods the mice were older than 110 days at the start of the experiments. The mice were kept
626 under 12h light/dark cycle with light on at 6 am and *ad libitum* access to food and water.

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628 Construction of optoTrkB

629 Photolyase homology region (PHR) domain of optoTrkB (Chang et al., 2014) was optimized in codon
630 usage for mice and connected with a flexible tag (Kennedy et al., 2010) to the C-terminus of a full-
631 length mouse TRKB, as previously described (Umemori et al, submitted). For cre-dependent
632 expression of optoTrkB, a double floxed inverted open-reading frame (DIO) structure (Fenno et al.,
633 2014) of optoTrkB (DIO-optoTrkB) was constructed (Figure S1A). We artificially synthesized DNA
634 including human Synapsin promoter (Kügler et al., 2001), lox2272 (Lee & Saito, 1998), loxP, inverted
635 optoTrkB sequence [PHR, flexible tag (Kennedy et al., 2010), mouse full-length TrkB
636 (NM_001025074)], lox2272, loxP, and cloned into pFCK(0.4)GW (a gift from Pavel Osten) (Addgene
637 plasmid # 27229 ; <http://n2t.net/addgene:27229>; RRID:Addgene_27229) (Dittgen et al., 2004) using
638 PacI and EcoRI cloning sites (Figure S1a). Furthermore, for electrophysiological studies, we
639 constructed DIO-optoTrkB expressing TdTomato (Shaner et al., 2004) (DIO-optoTrkB-IRES-TdT)
640 (Figure S1C). A fragment of Internal ribosome entry site (IRES) with TdTomato was amplified by PCR
641 with primers (5'- ggcgcgCCCCCTCTCCCTCCCCCCC -3' and 5'-
642 ggcgcgccTTACTTGTACAGCTCGTCCATGCCGTACAG -3') using Hot start Q5® polymerase (NEB,
643 Frankfurt, Germany) from LeGO-iT (a gift from Boris Fehse) (Addgene plasmid # 27361 ;
644 <http://n2t.net/addgene:27361> ; RRID:Addgene_27361) (Weber et al., 2010), and cloned into pCR™
645 Blunt II-TOPO® vector (ThermoFisher Scientific, Hvidovre, Denmark). Then the sequence was
646 confirmed by sequencing and sub-cloning into Ascl cloning sites of DIO-optoTrkB (Figure S1A) to
647 construct DIO-optoTrkB-IRES-TdT. Cre-dependent Inversions of DIO-optoTrkB and DIO-optoTrkB-
648 IRES-TdT were confirmed in HEK293 cells co-transfected with the DIO-vectors and plasmids
649 expressing Cre recombinase followed by PCR (Set1 (Figure S1B), and immunocytology (Figure S1D).
650 The following primers were used to confirm the inversion of DIO-optoTrkB:
651 Set1 (5'- AGTCGTGTCGTGCCTGAGA -3' and 5'- GAAATTTATGTGCCGCAGGT -3'); Set2 (5'-

652 CTGCTGGCAAAGGCTATTTTC -3' and 5'- GGGCCACAACCTCCTCATAAA -3').

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654 Virus generation

655 DIO-optoTrkB/DIO-optoTrkB-IRES-TdT, pLP1, pLP2, and pVSVG were co-transfected into HEL293FT
656 cells by jetPEI® (Polyplus Transfection, Illkirch, France) according to the manufacturer's instruction,
657 for producing lentivirus through a previously reported method (Hioki et al., 2007). Briefly, 3µg of
658 the plasmids was transfected into HEL293FT cells and cultured on a poly-D-Lysine coated dish
659 containing pre-warmed Opti-MEM® medium (ThermoFisher Scientific) with 2% FBS for 72 hrs. Then
660 the culture supernatant was collected and exchanged with a new Opti-MEM® medium with 2% FBS
661 followed by incubation for 72 hrs. The culture supernatant was collected again and centrifuged at 4
662 °C at 2000 g for 10 min. The supernatant was then concentrated with Amicon® Centrifugal Filter
663 Units Ultra -15 (Merck Millipore, Darmstadt, Germany) into less than 10ml. The concentrated
664 solution was purified using the sucrose gradient method (Tiscornia et al., 2006) and aliquots were
665 stored at -80 °C until use. From one aliquot the p24 capsid protein concentration was measured to
666 estimate the infection unit (IU).

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668 Virus injection

669 PV-cre mice were anaesthetized with 2.5% isoflurane and DIO-optoTrkB (p24 concentration/titer
670 4.04×10^7) was stereotactically injected into the binocular area of the V1, previously identified
671 through optical imaging (see below) and using blood vessels as landmarks. DIO-optoTrkB was
672 stimulated with blue LED light (470nm) through the transparent skull during monocular deprivation
673 twice a day for 30 seconds at 8-10 am and 4-6 pm. Blue LED light was provided by a BLS Super High
674 Power Fiber-Coupled LED Light Sources (BLS-FCS-0470-100, Mightex, Pleasanton, CA) connected to
675 a BioLED Light Source Control Modules (BLS-3000-2, Mightex). To avoid light exposure through the

676 skull, the transparent skull was covered by black nail polish between light stimulation sessions.ht

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678 Surgery

679 For chronic imaging of intrinsic signals, the animals underwent transparent skull surgery as
680 described previously (Steinzeig et al., 2017).

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682 Fluoxetine treatment

683 Fluoxetine (Bosche Scientific, New Brunswick, NJ) was administered via drinking water (0.08 mg/ml),
684 corresponding to a dose of approximately 8-12 mg/kg/day, and kept in light-protected bottles. The
685 drinking water of both the control and fluoxetine group contained 0.1 % saccharin, was changed
686 twice a week and the consumption was monitored. Drug treatment started after transparent skull
687 surgery and continued throughout the whole experiment.

688

689 Monocular deprivation and virus stimulation

690 The animals were anaesthetized with intraperitoneal injection as described above. The eye lashes
691 were trimmed and the eye lid margins were sutured shut with 3 mattress sutures. To prevent
692 postoperative infections, an eye ointment containing dexamethasone was applied. The eyes were
693 checked daily until reopening, re-sutured if needed, and mice with signs of corneal injury were
694 excluded from the experiments.

695

696 Optical imaging of intrinsic signals

697 We determined the strength of neuronal responses to stimulation of either eye in the binocular
698 region of the V1 using imaging of intrinsic signals (IOS) (Kalatsky & Stryker, 2003) before (IOS I; IOS

699 III) and after (IOS II; IOS IV) monocular deprivation. The animals were chamber anaesthetized with
700 1.8% isoflurane in a 1:2 mixture of O₂/air and then intubated and ventilated with 1.2% isoflurane in
701 the same mixture.

702 Intrinsic optical signal responses were recorded from the V1 of the right hemisphere according to a
703 previously described protocol (Kalatsky & Stryker, 2003), which was modified for the measurement
704 of OD plasticity (Cang et al., 2005).

705

706 Data analysis of optical imaging

707 Cortical maps were computed based on the acquired frames using Fourier decomposition to extract
708 the signal from biological noise using an analysis software package for continuous recording of
709 optical intrinsic signals (VK Imaging, USA) (Kalatsky & Stryker, 2003). The ODI was then calculated
710 for every pixel within the binocularly responding region based on the formula $(C-I)/(C+I)$, where “C”
711 refers to the response magnitude of the contralateral eye and “I” to that of the ipsilateral eye. For
712 each animal, several ODIs were collected and then averaged. Positive ODI values represent
713 contralateral dominance, negative represent ipsilateral dominance, while ODI values of 0
714 correspond to equally strong contralateral and ipsilateral eyes.

715

716 In vivo electrophysiology

717 To measure oscillations in the binocular region of the optoTrkB transfected area, PV-cre mice
718 underwent transparent skull surgery and optoTrkB was injected as described above. Under 15%
719 urethane anesthesia (in PBS), a hole for a 16 channel optrode (A1x16-10mm-100-177-OA16LP,
720 NeuroNexus, Ann Arbor, MI) was drilled next to the injection site and a hole for a ground electrode
721 next to lambda. The animals were head-fixed and the optrode was covered with Dil and slowly
722 inserted into the V1 to a depth of about 2200 μ m. The data was recorded using a Smartbox

723 (NeuroNexus) at a sampling rate of 20 kHz. After collecting a 20 minute baseline, optoTrkB was
724 stimulated for 30 seconds with blue LED light and recorded for 2h, separated into 20-minute blocks.
725 Finally, the animals were transcardially perfused and the brains fixed with 4% PFA. To confirm the
726 position of the electrodes and co-localization with optoTrkB, 250 μ m coronal slices were cut on a
727 vibratome.

728 The digitized data were first offline notch filtered at 50 Hz by the Python RHD2000 interface provided
729 by Intan. Afterwards, the data were band-pass filtered between 4 and 150 Hz, using a 2nd order
730 Butterworth filter with zero-phase shift, and then downsampled to a sampling rate of 1 kHz to
731 extract the local field potential (LFP) component. The LFP for each electrode contact was normalized
732 to zero and the underlying current source density (CSD) was calculated as the second spatial
733 derivative, using previously published methods (Voigt & Kral, 2019).

734 Time-frequency analysis of the CSD data was performed by wavelet filtering using 36 Morlet
735 wavelets with a width parameter $m = 6$ and with frequencies ranging from 4 to 112 Hz with log-
736 constant scaling. For further analyses, the signal from all electrodes that were identified to reside in
737 areas with optoTrkB expression by the histological analysis was averaged, and the data were binned
738 into non-overlapping 20 seconds bins. Morlet power spectra were estimated by computing the
739 averaged magnitude over all bins of the 20-minute recording period per condition. For statistical
740 analysis, broadband power in the 20-minute baseline period and the 20-minute period beginning
741 100 minutes after LED onset was compared using permutation testing. Surrogate data was generated
742 by 1000 random permutations of condition labels and the effect size was measured using Cohen's
743 d. All data analyses and statistics were performed in Python (ver. 2.7) using SciPy (ver. 0.19.0), the
744 NeuroDSP toolbox (ver. 2.0.1-dev)(Cole et al., 2019), and custom written scripts.

745

746 Electrophysiology in acute slices

747 The brains of optoTrkB-infected PV-cre mice were dissected in darkness using red light illumination,
 748 and kept in the dark throughout the whole experiment. The brains, including brains of fluoxetine-
 749 treated WT and hPV-TrkB cKO mice, were dissected and immersed in ice-cold dissection solution
 750 containing (in mM): 124 NaCl, 3 KCl, 1.25 NaH₂PO₄, 1 MgSO₄, 26 NaHCO₃, 15 D-glucose, 9 MgSO₄
 751 and 0.5 CaCl₂. The cerebellum and anterior part of the brain were removed and coronal 350µm brain
 752 slices of the V1 were cut on a vibratome (Leica Biosystems, Wetzlar, Germany). Slices were divided
 753 into two groups and allowed to recover for 30 min at 31-32°C in artificial cerebrospinal fluid (ACSF)
 754 containing (in mM): 124 NaCl, 3 KCl, 1.25 NaH₂PO₄, 1 MgSO₄, 26 NaHCO₃, 15 D-glucose, and 2 CaCl₂
 755 and bubbled with 5% CO₂/95% O₂.

756 OptoTrkB-transfected slices in one of the groups were acutely stimulated for 30 seconds with blue
 757 light after transferring them to the recording chamber, whereas the transfected slices in the other
 758 group were kept in darkness.

759 Field excitatory postsynaptic currents (fEPSPs) were recorded in an interface chamber (32°C) with
 760 ACSF-filled glass microelectrodes (2-4 MΩ) positioned within layer II/III of the V1 using an Axopatch
 761 200B amplifier (Molecular devices, San Jose, CA). Electric stimulation (100 µsec duration) was
 762 delivered with a bipolar stimulation electrode placed at the border of the white matter (WM) and
 763 layer VI. Baseline synaptic responses were evoked every 20 seconds with a stimulation intensity that
 764 yielded a half-maximum response. After obtaining a 15 minute stable baseline, θ burst stimulation
 765 (TBS) (4 sweeps at 0.1 Hz, each sweep with 10 trains of 4 pulses at 100 Hz at 200 ms intervals) was
 766 delivered and field potentials in response to 0.05 Hz stimulation were recorded for additional 45
 767 minute. WinLTP (0.95b or 0.96, www.winltp.com) was used for data acquisition and analysis.

768

769 To measure intrinsic excitability, the brains of PV-cre mice transfected with the DIO-optoTrkB-IRES-

770 TdT lentivirus were dissected as described above but cut in a protective NMDG ACSF (Ting et al.,
771 2014) containing (in mM): 92 NMDG, 2.5 KCl, 1.25 NaH₂PO₄, 30 NaHCO₃, 20 HEPES, 25 glucose, 2
772 thiourea, 5 Na-ascorbate, 3 Na-pyruvate, 0.5 CaCl₂·4H₂O and 10 MgSO₄·7H₂O, pH 7.3–7.4 The slices
773 were allowed to recover at 32°C for 10 min, after which they were transferred to modified ACSF
774 containing (mM): 92 NaCl, 2.5 KCl, 1.25 NaH₂PO₄, 30 NaHCO₃, 20 HEPES, 25 glucose, 2 thiourea, 5
775 Na-ascorbate, 3 Na-pyruvate, 2 CaCl₂·4H₂O and 2 MgSO₄·7H₂O , pH 7.3-7.4 for storage. Recordings
776 were done in submerged chamber perfused with normal ACSF (32°C).

777 Whole cell patch clamp recordings from the TdT expressing PV cells were obtained under visual
778 guidance under ambient light with glass microelectrodes (3-5 MΩ) filled with a low Cl⁻-filling solution
779 containing (in mM): 135 K-gluconate, 10 HEPES, 2 KCl, 2, Ca(OH)₂, 5 EGTA, 4 Mg-ATP, 0.5 Na-GTP
780 using a Multiclamp 700A amplifier (Axon Instruments, Foster City, CA). Uncompensated series
781 resistance (Rs) was monitored by measuring the peak amplitude of the current response to a 5 mV
782 step. Only experiments where Rs < 30 MΩ, and with < 20 % change in Rs during the experiment,
783 were included in analysis.

784 Intrinsic excitability was measured in current clamp mode by injecting currents ranging from -50 to
785 600 pA for 600 ms in 50 pA steps from the resting membrane potential of -60/-70 mV. The recordings
786 were analyzed in Clampfit (Molecular Devices, San Jose, CA) programs. A minimum of three action
787 potentials (AP) were averaged based on their AP half-width (10th AP at 200pA injected from
788 rheobase). sEPSCs were recorded under voltage clamp at -70 mV and analyzed in Clampfit
789 (Molecular Devices). The threshold for detection of inward sEPSC events was three times the
790 baseline noise level, and all detected events were verified visually.

791

792 Sample collection

793 For collecting tissue samples for qPCR and Western Blot experiments, the brains of optoTrkB infected
 794 PV-cre mice were dissected and immersed in ice-cold dissection solution containing (in mM): 124
 795 NaCl, 3 KCl, 1.25 NaH₂PO₄, 1 MgSO₄, 26 NaHCO₃, 15 D-glucose, 9 MgSO₄ and 0.5 CaCl₂. The V1 was
 796 dissected and incubated at 31-32°C in artificial cerebrospinal fluid (ACSF) containing (in mM): 124
 797 NaCl, 3 KCl, 1.25 NaH₂PO₄, 1 MgSO₄, 26 NaHCO₃, 15 D-glucose, and 2 CaCl₂ and bubbled with 5%
 798 CO₂/95% O₂. The tissue samples were either immediately collected in NP lysis buffer and
 799 homogenized (control), or stimulated with blue LED light for 30 seconds and collected and
 800 homogenized after 15 minutes, 30 minutes or 60 minutes. All of the procedures were done in dark
 801 conditions. The samples were further used for qPCR and Western Blot analysis.

802

803 Western Blot

804 The samples were centrifuged (16000 g, 15 min at +4°C) and the supernatant was used to measure
 805 the protein concentrations using the Lowry method (Biorad DC protein assay, Bio-Rad, Richmond,
 806 CA) (Lowry et al, 1951). The samples were separated in a SDS-PAGE (2-4% gradient gel, NuPage™;
 807 ThermoFisher Scientific) and blotted to a PVDF membrane (300 mA, 1 h, + 41°C). The membranes
 808 were washed in Tris Buffer Solution with 0,001% Tween[®] 20 (TBST), blocked in TBST with 3% BSA for
 809 1 hour at room temperature and incubated in primary antibody solutions (in TBST with 3% BSA)
 810 directed against: phosphorylated (pY816, #4168, 1:1000; pY515, #9141, 1:1000; pY706-7, #4621S,
 811 1:1000, Cell signaling technology (CST), Danvers, MA) and non-phosphorylated forms of TrkB and
 812 CREB (TrkB, 1:2000, BD Transduction Laboratories, San Jose, CA; CREB, #4820S, CST, Danvers, MA) at
 813 +4°C for overnight. After washing in TBST, the membranes were further incubated in secondary
 814 antibody solutions (TBST with 5% Non-Fat Dry Skinned Milk and Horseradish Peroxidase conjugated

815 secondary antibodies Goat Anti-Rabbit/Mouse, 1:10000, Bio-Rad) for 2 hours at room temperature.
816 After washing with TBST and rinsing with PBS, secondary antibodies were visualized by an
817 electrochemiluminescence kit (ECL plus, ThermoFisher Scientific) according to the manufactures
818 instruction, and detected using a FUJIFILM LAS-3000 dark box.

819

820 qPCR

821 RNA was purified from the lysate following the manufacturer's protocol using a combined protocol
822 of QIAzol® (Qiagen, Hilden, Germany) and NucleoSpin® (Macherey-Nagel, Düren, Germany). Briefly,
823 the aqua layer was isolated after Qiazol and chloroform extraction. The RNA was washed in 100%
824 ethanol and the DNA was digested in the spin columns. The purified RNA was then reverse
825 transcribed to cDNA using Maxima First Strand cDNA Synthesis Kit (ThermoFisher Scientific). The
826 amount of cDNA synthesized from the target mRNA was quantified by real-time PCR (qPCR). The
827 following primers were used to amplify specific cDNA regions of the transcripts of interest:

828 Kv3.1 (5' -AGAGATTGGCACTCAGTGACT-3' and 5' -TTGTTACGATGGGGTTGAAG-3'), Kv3.2 (5'
829 -AGGCTATGGGGATATGTACCC-3' and 5' -TGCAAAATGTAGGCGAGCTTG-3'), PV (5'-
830 TGTCGATGACAGACGTGCTC-3' and 5'-TTCTTCAACCCCAATCTTGC-3').

831

832 Immunohistochemistry

833 Animals were transcardially perfused with PBS followed by chilled 4% paraformaldehyde (PFA) in
834 PBS. Brains were removed under ambient light and left for fixation in 4% PFA overnight at +4 °C. For
835 cutting, the brains were embedded in 3% agar and cut into 40 µm coronal visual cortical sections
836 using a vibratome (Leica Biosystems, Wetzlar, Germany). After washing with PBST (1x PBS and 0.2%
837 TritonX100), the sections were incubated in 10% donkey serum (Vector Laboratories, Peterborough,

UK) and 3% Bovine Serum Albumin (BSA) (Sigma-Aldrich, Steinheim, Germany) in PBST for 30 minutes at room temperature. Next, the sections were incubated with the following antibodies: 1) guinea pig anti-parvalbumin (1:1000) (#195004, Synaptic Systems, Göttingen, Germany), 2) biotinylated lectin from Wisteria floribunda (WFA; 1:200) (#L1516-2MG, Sigma-Aldrich), and 3) rabbit anti-phospho Kv3.1 (1:100) (#75-041, Phosphosolutions, Aurora, CO) overnight at +4° C. After washing in PBST, the samples were further incubated in: 1) goat anti-guinea pig secondary antibody conjugated with Alexa Fluor647/546 (1:1000) (Abcam, Cambridge, UK), 2) streptavidin conjugated with Alexa Fluor488 (1:1000) (Thermofisher Science), or 3) Goat anti-rabbit conjugated with Alexa Fluor 647 (1:1000) (Life technologies, Carlsbad, CA) for 2 hours at room temperature protected from light. After final washing in PBS, the sections were transferred to 0.1M PB with gelatin, mounted on glass slides and covered with DAKO mounting medium (Sigma Aldrich).

849

850 Image acquisition and analysis

851 Quantitative analysis of immunostainings was performed blind. Images were taken from the V1
852 according to the mouse brain atlas.

853 Laser scanning confocal microscopy was used to detect PV positive (PV⁺), PNN positive (PNN⁺),
854 double positive (PV⁺PNN⁺) and pKv3.1-positive cells. Images were obtained using a confocal
855 microscope LSM 700 (Carl Zeiss, Vantaa, Finland) equipped with a 10× objective lens (10x Plan-
856 Apochromat 10x/0.45, Carl Zeiss) and imaging Software ZEN 2012 lite (Carl Zeiss). From each section,
857 a z-stack containing at least 10 consecutive images was obtained. A minimum number of 3 sections
858 per animal were imaged using the same microscope and the same camera settings for all samples.
859 Image processing was done using ImageJ software version 1 (Schneider et al., 2012). All images in
860 each z-stack were analyzed and the number of cells was averaged per z-stack.

861 To determine the PV cell populations, frequency distribution analyses were performed on PV

intensities taken from non-stimulated optoTrkB samples or control WT samples to serve as reference group. The PV cell populations were defined as low PV (0-8000 a.u.), intermediate-low PV (int-low PV, 8000-16000 a.u.), intermediate-high PV (int-high, 16000-24000 a.u.) and high PV (24000-36000 a.u.) expressing cells and these criteria were applied to the light stimulated samples.

Nuclei preparation

Single nuclei RNA sequencing (snRNA-seq) was conducted by following previously reported methods with some modifications (Bakken et al., 2018; Krishnaswami et al., 2016) (Figure S3A). Considering that optoTrkB is sensitive to light stimulation, we used snRNA-seq instead of single cell RNA sequencing (scRNA-seq) after FACS selected by TdTomato fluorescence, to avoid laser-induced optoTrkB activation during FACS. There is a good correlation of gene expression and gene detection sensitivity in each cell between SnRNA-seq and scRNA-seq (Bakken et al., 2018), and snRNA-seq has several advantages, such as reduced dissociation bias and dissociation-induced transcriptional stress responses (Wu et al., 2019). Tissue samples were obtained bilaterally from the visual cortex of adult female PV-cre mice infected with DIO-optoTrkB-TdT. After deep anesthesia with pentobarbital, the mice were perfused with ice-cold NMDG ACSF (Ting et al., 2014) consisted of 0.5 mM CaCl_2 , 25mM D-Glucose, 98 mM HCl, 20 mM HEPES, 10 mM MgSO_4 , 1.25 mM NaH_2PO_4 , 3 mM Myo-inositol, 12 mM N-acetylcysteine, 96 mM N-methyl-D-glucamine, 2.5 mM KCl, 25 mM NaHCO_3 , 5 mM sodium L-Ascorbate, 3 mM sodium pyruvate, 0.01 mM Taurine, and 2 mM Thiourea, bubbled with a carbogen gas (95% O_2 and 5% CO_2), and the brains were isolated and sliced on a vibratome (MICROM, HM 650V, Thermofisher Science) in NMDG ACSF (Ting et al., 2014). Then, the slices were exposed to blue LED light (30 seconds), and incubated in a modified HEPES ACSF including 92 mM NaCl, 2.5 mM KCl, 1.25 mM NaH_2PO_4 , 30 mM NaHCO_3 , 20 mM HEPES, 25 mM glucose, 2 mM thiourea, 5 mM Na-ascorbate, 3 mM Na-pyruvate, 2 mM CaCl_2 , 2 mM MgSO_4 , 3 mM Myo-inositol, and 0.01 mM Taurine

886 for one hour. The visual cortex was bilaterally isolated from the slices under a microscope using red
887 light illumination, and tissues were transferred to pre-cooled Dounce homogenizers filled with cold
888 homogenization buffer (250 mM sucrose, 25 mM KCl, 5 mM MgCl₂, 10 mM Tris buffer, pH 8.0, 1 μM
889 DTT, 1× protease inhibitor (cOmplete, Sigma-aldrich), 0.4 U/μl RNase Inhibitor (Promega), 0.2 U/μl
890 Superscript (ThermoFisher Science, Waltham, MA), 0.1% (v/v) Triton X-100 (Sigma-Aldrich), 10 μg/ml
891 Cyclohexamide (Sigma-Aldrich), 10 μg/ml Actinomycin D (Sigma-Aldrich), 10 ng/ml Hoechst 33342
892 (ThermoFisher Science)). The tissues were homogenized with five strokes of the loose pestle,
893 followed by 10 strokes of the tight pestle. Finally, nuclei of the tissues were obtained by filtration
894 through a BD Falcon™ cell strainers (BD Falcon, San Jose, CA). All procedures were performed with
895 RNase free in the dark condition.

896

897 Immunostaining and sorting of nuclei

898 The nuclei were centrifuged (1,000 x g, 8 min, 4 °C) and resuspended in staining buffer (PBS, pH 7.4,
899 with 0.5% RNase-free BSA (Promega, Heidelberg, Germany) and 0.2 U/μl of RNasin Plus RNase
900 inhibitor (Promega)). Samples were incubated in blocking/washing buffer (PBS with 0.5% BSA). The
901 nuclei were incubated with mouse monoclonal anti-NeuN antibody (1:5000) (MAB377, Sigma-
902 Aldrich) or purified mouse IgG1k (1:5000) (554121, BD Pharmingen, San Diego, CA) as a negative
903 control for staining and sorting for 30 min with rotation at 4 °C. Then the samples were washed in
904 staining buffer by inverting the tubes several times, and were centrifuged for 5 min at 400g at 4 °C.
905 The pellet nuclei were re-suspended in 500 μl of staining buffer containing goat anti-mouse Alexa
906 Fluor 488 secondary antibody (ThermoFisher Science) with a final dilution of 1:5000, and were
907 incubated with rotation for 30 min at 4 °C followed by washing with blocking/washing buffer.

908

909 FACS of nuclei

910 Hoechst 33342 and NeuN positive nuclei stained with Alexa488 were sorted and collected by BD
911 Influx cell sorter (BD Biosciences, Heidelberg, Germany) (Figure S3B). After checking the quality,
912 collected nuclei were loaded on Chromium Controller (10x Genomics).

913

914 Single nuclei RNA sequencing (snRNA-seq)

915 Preparation of cDNA Library was done using Chromium Single Cell 3' Gene Expression v3 library
916 preparation kit (10x Genomics), and sequenced with NovaSeq 6000 (Illumina, San Diego, CA) with
917 read lengths: 28bp (Read 1), 8bp (i7 Index), 0 bp (i5 Index) and 89bp (Read 2). Data was pre-
918 processed using CellRanger 3.0. The reads are mapped to reference genome that included introns,
919 and we obtained 3,789 and 6,923 cells with approximately 10k median reads per cell (Figure S4).

920

921 Data analysis for snRNA-seq

922 Data analysis was carried out using Chipster version 3.16 (Kallio et al., 2011), which utilizes Seurat
923 v3 (Butler et al., 2018). Cells with more than 7,000 and less than 100 reads were filtered out, and
924 reads for the remaining cells were scaled logarithmically. Based on results for plotting variance
925 against expression, 3000 most highly varied genes in 10,706 cells were used for clustering and
926 sample integration. By Seurat's integrated analysis (Butler et al., 2018), 20 canonical components
927 were used for finding anchors and 20 principal components to integrate the samples together.
928 Clusters were obtained using UMAP (McInnes et al., 2018) with 20 principal components and
929 resolution 0.5 (Figure S5A, and S5B). This resulted in 18 clusters and we identified the clusters of
930 pyramidal neurons and interneurons with previously reported markers (Tasic et al., 2018), such as
931 *Glutamate decarboxylase 1 (Gad1)*, and those of interneurons with *Solute Carrier Family 6 Member*
932 *1 (Slc6a1)*, *Parvalbumin (Pvalb)*, *Paired box protein (Pax6)*, *Vasoactive intestinal peptide (VIP)*, and

933 *Somatostatin* (*Sst*) (Fig. 2A) (Figure S5C). Then a subset of data containing only parvalbumin-positive
 934 interneurons was selected and clustered again by UMAP with the same parameters resulting in four
 935 different clusters (Fig. 2B) (Figure S6A and S6B). Markers and differentially expressed genes between
 936 non- and light- exposed samples were detected in each cluster (Figure S6D) (Table S3-6). Pathway
 937 and gene ontology analysis was carried out with DAVID (Huang et al., 2009).

938

939 Statistical analysis

940 All statistical graphs were generated using Graphpad Prism v.6.07 (GraphPad Software, San Diego,
 941 CA). Unpaired t-Test, Two-way and one-way ANOVA followed by Tukey's or Bonferroni post hoc tests
 942 were also performed using Graphpad Prism v.6.07. The confidence level was set to 0.05 (P value)
 943 and all results were presented as means $\pm$ s.e.m. All the individual data points are shown in the
 944 histograms. The data distribution was assumed to be normal but this was not formally tested. The
 945 sample size was determined based on our previous experience.

946

947 DATA AND SOFTWARE AVAILABILITY

948 snRNA-seq data have been deposited in NCBI's Gene Expression Omnibus (Edgar et al., 2002) and
 949 are accessible through GEO Series accession number GSE 142797
 950 (<https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE142797>). Other raw data is available in
 951 Mendeley Data, V1, doi: 10.17632/992cg2vrj5.1
 952 ([https://data.mendeley.com/datasets/992cg2vrj5/draft?a=ab1dd775-63b5-4eca-aa01-](https://data.mendeley.com/datasets/992cg2vrj5/draft?a=ab1dd775-63b5-4eca-aa01-0a565fb6fa0d)
 953 [0a565fb6fa0d](https://data.mendeley.com/datasets/992cg2vrj5/draft?a=ab1dd775-63b5-4eca-aa01-0a565fb6fa0d)).

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Figure S2. OptoTroB activation mediates plasticity in the visual cortex.

(A-E) Western Blot analysis of V1 of PV-cre mice infected with DIO-optoTrkB without light stimulation (Control) and 15 minutes, 30 minutes and 60 minutes after light stimulation. Acute stimulation of optoTrkB with light results in increased phosphorylation of tyrosine sites: (A) Y816, (B) Y515, (C) Y706 and (D) phosphorylation of CREB. (E) Representative Western Blots of Y816, Y515, Y706 and CREB. (F) PV-cre mice without optoTrkB expression stimulated with blue light twice daily during 7 days of MD show no shift in ocular dominance. (G) PV mRNA levels are decreased 60 min after light stimulation measured by qPCR. qPCR measurements of control, 15 minutes, 30 minutes and 60 minutes after light stimulation. OptoTrkB reduces the expression of PV 60 minutes after stimulation. One-way ANOVA ($F(3, 11) = 5.112$; $p = 0.0186$) with Bonferroni's post-hoc test comparing control vs. 15 min, 30 min and 60 min (control vs. 15 min, $p = > 0.9999$; control vs. 30 min, $p = 0.1456$; control vs. 60 min, $p = 0.0125$). $n = 3-4$ animals/group. Bars represent means $\pm$ SEM. * $p < 0.05$

Figure S3. snRNA-seq after activation of optoTrkB

(A) Scheme of snRNA-seq. The primary visual cortices were bilaterally isolated from PV mice transfected by DIO-optoTrkB-TdT. Nuclei were extracted from the tissue, and Hoechst+ and NeuN+ cells were sorted by FACS. A cDNA library was constructed with Poly (A)-RNA in droplets. The fragmented cDNA, were then sequenced on a Next generation sequencer (NGS). (B) Sorting nuclei of neurons by FACS. Cells were stained with Hoechst 33342, NeuN antibody, and mouse IgG antibody (negative control) followed by labelling with anti-mouse IgG conjugated with Alexa488, and then sorted by BD Influx cell sorter (BD Biosciences). Hoechst 33342 positive nuclei were efficiently sorted in all samples. While small portion of nuclei (0.15%) labelled with IgG bound to Alexa488 were sorted in negative control, large portion of nuclei (~72%) stained with NeuN bound to Alexa488 were efficiently sorted and collected.

Figure S4. Number of genes, reads, and mitochondrial genes in nuclei.

The plots represent genes and transcripts per cell, where the horizontal axis is cell id and vertical axis is the number of genes or transcripts, respectively. Each dot is a unique cell and orange areas represent cell density at vertical axis. In the right panels, the horizontal axis is cell id and the vertical axis is the percentage of transcripts originating from mitochondrial DNA. Each cell should be represented with a dot and the black line in 0.000 % shows absence of mitochondrial genes. Since we used only nuclei, mitochondrial genes are absent. Based on genes and transcripts per cell, we decided to remove cells that had fewer than 100 or greater than 7000 reads.

Figure S5. Clustering single nuclei

(A) For 20 first principal components, heatmaps are drawn with cells as columns and the 15 most important genes as rows. The heatmaps show major variation captured in 20 first principal components. (B) Elbow plot is generated by calculating how much of the variation is explained by the first 20 principal components. Horizontal axis represents ordinal number of principal component and vertical axis the standard deviation of explained percentage in variation. Most of the variation in the data is included in 20 first principal components, which were used for later analysis. (C) In the violin plot, the vertical axis indicates the expression level of a gene in a cluster (number below), and the horizontal axis indicates cell density at a given expression level. Pyramidal neurons were identified by using the markers Nrn1, Sv2b and Satb2 and while interneurons were done by Gad1 and Slc6a1. The cluster 5, 11, and 13 contained parvalbumin-, somatostatin-, and VIP-positive interneurons, respectively.

Figure S6. Clustering PV interneuron

(A) Genes in representative PCAs explain four clusters in PV neurons. Heatmaps of unique genes in each clusters show that the PV cluster can be further divided into four clusters (0-3). (B) Number and ratio of PV cells used for clustering and DE analysis. (C) Distribution of expression of genes in cluster specific genes (*Pvalb*, *Il1rapl2*, *Synpr*, *Nkain3*), representative markers for chandelier cells (*Hs3st4*, *Errfi1*), and basket cells (*Reln*, *Tac1*) as reported previously (Tasic et al., 2018). (D) Distribution of cells expressing each marker in the four clusters of PV interneurons. These results demonstrate that each cluster has unique expression of marker genes, and the cluster 3 includes both chandelier and basket cells.

Figure S7. Representative traces of electrophysiological recordings
Representative traces used to estimate intrinsic excitability (150 pA, top row), AP half-width (2nd row) and sEPSC (bottom row) in WT mice treated with water (first column), WT mice treated with Flx (2nd column), CKO mice treated with water (3rd column) and CKO mice treated with Flx (last column).

Number of cells in each cluster

Cluster	Neuronal type ^a	Non-activated		Activated	
		N ^b	% ^c	N ^b	% ^c
0	Pyramidal	696	18.4	1200	17.3
1	Pyramidal	349	9.2	874	12.6
2	Pyramidal	368	9.7	827	11.9
3	Pyramidal	390	10.3	746	10.8
4	Pyramidal	222	5.9	408	5.9
5	Internuron	173	4.6	413	6.0
6	Pyramidal	218	5.8	360	5.2
7	Pyramidal	172	4.5	334	4.8
8	Pyramidal	261	6.9	235	3.4
9	Pyramidal	192	5.1	293	4.2
10	Pyramidal	189	5.0	266	3.8
11	Internuron	146	3.9	237	3.4
12	Internuron	82	2.2	169	2.4
13	Internuron	82	2.2	164	2.4
14	Pyramidal	87	2.3	143	2.1
15	Pyramidal	68	1.8	130	1.9
16	Pyramidal	76	2.0	102	1.5
17	Internuron	14	0.4	20	0.3

Table S1. Number of nuclei in each cluster.

(a) All clusters were identified by expression of makers previously reported (Tasic et al., 2018).

(b) Number of cells in each cluster were provided by Chipster/Seurat version 3.

(c) Percentage was calculated by dividing the number of nuclei in each cluster with that of total nuclei in non-activated or activated sample

A

Input resistance	Control	10-30 min	30-60 min
Mean	237.5	233.8	272.6
Std. Deviation	63.99	58.7	77.3
Std. Error of Mean	26.12	22.19	31.56

B

Input resistance	WT water	WT Flx	hPV-TrkB CKO water	hPV-TrkB CKO Flx
Mean	171.2	251.2	189.4	204.3
Std. Deviation	46.84	69.45	55.23	71.69
Std. Error of Mean	15.61	24.55	20.87	20.69

Table S6. Input resistance is after optoTrkB activation and fluoxetine treatment in WT and hPV-TrkB CKO mice.

(A) Input resistance is unchanged after optoTrkB activation. (B) No effect of fluoxetine treatment on input resistance in WT and hPV-TrkB CKO mice.

Supplemental Table 2. Markers in all clusters.

Supplemental Table 3. DE genes after optoTrkB activation in all clusters

DE genes were identified by Chipster/Seurat version 3. Gene ontology (GO) in each gene was annotated by DAVID program

Supplemental Table 4. Marker genes in clusters of Parvalbumin interneurons

Supplemental Table 5. DE genes after optoTrkB activation in clusters of Parvalbumin interneurons.

DE genes were identified by Chipster/Seurat version 3. GO in each gene was annotated by DAVID program

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