Genetic control of neuronal activity enhances axonal growth only on permissive substrates

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Abstract

Neural tissue has limited regenerative ability, to cope with that, in the recent years a diverse set of novel tools have been used to tailor neurostimulation therapies and promote functional regeneration after axonal injuries. In this report, we explore cell-specific methods to modulate neuronal activity, including optogenetics and chemogenetics to assess the effect of specific neuronal stimulation in the promotion of axonal regeneration after injury. We found that opto- or chemogenetic modulations of neuronal activity on both dorsal root ganglia and corticospinal motor neurons increase their axonal growth capacity only on permissive substrates.

1 Introduction

Following injury, damaged axons from the central nervous system (CNS) degenerate and are unable to regenerate, while surviving fibres, have a limited capacity to sprout and to re-establish lost connections, leading to functional impairment (Fitch and Silver, 2008).

This failure of CNS axons to regenerate after injury is partly attributed to a hostile CNS environment for growth (Richardson et al., 1980; Huebner and Strittmatter, 2009; Cregg et al., 2014). The injury site is rich in growth inhibitory proteins including myelin-associated glycoproteins, and chondroitin sulfate proteoglycans (CSPG) while lacking trophic support for axon regeneration (Jones et al., 2003; Ohtake and Li, 2015; Sami et al., 2020). The limited intrinsic regenerative capacity of adult CNS axons also contributes to the failure of axon regeneration after injury (He and Jin, 2016; Curcio and Bradke, 2018; Mahar and Cavalli, 2018). Yet, in the last years there have been several studies approaching different aspects of the CNS physiology to both boost the intrinsic regrowth capacity and overcome the extrinsic inhibition (Wu et al., 2015; Anderson et al., 2018).

During development, when CNS neurons still retain their intrinsic ability to regenerate after axotomy (Arlotta et al., 2005), neuronal activity critically determines their connectivity, particularly onto spinal targets (Friel et al., 2014). In this direction, neuronal activation has also been explored to try to increase the intrinsic capacity for axon regeneration as well as overcome the extrinsic inhibition. Practically, electrical stimulation has been shown to enhance regeneration of sensory axons after peripheral nerve or dorsal columns injury (Udina et al., 2008a; Goganau et al., 2018a) and sprouting of cortical axons into contralateral spinal cord gray matter after pyramidotomy (Carmel et al., 2010a, 2014; Carmel and Martin, 2014a).

In order to gain insight on the actual mechanisms underlying these improvements, specifically and remotely activating neurons through opto- and chemogenetics, has become the state-of-the-art approach for tailored activity modulation (Sternson and Roth, 2014; Deisseroth, 2015).

In this direction, the current study explores the therapeutic potential of modulating neuronal activity of sensory ascending (dorsal root ganglia (DRG)) and motor descending (corticospinal motor neurons (CSMN)) neurons, via opto- and chemogenetic stimulation in regulating axonal growth after injury.

We found that specific neuronal stimulation induced increased axonal growth capacity both *in vitro* and *in vivo*, and both in sensory and motor neurons, but only when in presence of permissive and trophic environments. Opto- or chemogenetic stimulation did not overcome the inhibitory signalling induced by myelin-associated glycoproteins and CSPGs.

2 Materials and Methods

Mice

B6.Cg-Tg(Thy1-COP4/EYFP)18Gfng/J (Thy1-ChR2) (Jackson Laboratories) (Arenkiel et al., 2007) or wild type (WT) C57BL/6J mice (Charles River Laboratories) ranging from 6-10 weeks of age were used for the experiments and were randomly divided into the different experimental groups. Mice were anaesthetized with isoflurane (5% induction, 2% maintenance) during surgeries. For DRG neuronal cultures adult Thy1-ChR2 mice were used. Embryonic day 16.5 (E16.5) OF1 pregnant females were purchased from Charles River Laboratories, and the embryos were used for neuronal cortical cultures. All procedures were approved by the Ethics Committee on Animal Experimentation (CEEAA) of the University of Barcelona (CEEAA approval #276/16 and 141/15).

Dorsal root ganglia (DRG) neuronal culture

DRGs from adult Thy1-ChR2 mice were dissected, washed in cooled Hank's balanced salt solution (HBSS; ThermoFisher Scientific), and enzymatically digested (5mg/ml Dispase II (Merck) and 2.5mg/ml Collagenase Type II (ThermoFisher Scientific) in DMEM (ThermoFisher Scientific)) for 45min at 37°C. Next, the DRGs were resuspended in DMEM:F12 (ThermoFisher Scientific) media supplemented with 10% heat-inactivated Fetal Bovine Serum (FBS; ThermoFisher Scientific) and 1x B27 (ThermoFisher Scientific) and were mechanically dissociated by pipetting. After centrifugation, the resulting single cells were resuspended in culture media (DMEM:F12 media with 1x B27 and penicillin/streptomycin (P/S; ThermoFisher Scientific)) and plated in glass-coverslips (3000-4000 cells/well) pre-coated with 0.1mg/ml poly-D-lysine (2h, 37°C; Merck) and 2µg/ml laminin (over-night (O/N); ThermoFisher Scientific) at RT (room temperature)). An additional incubation with 20 µg/ml

CSPGs (Merck) was performed (2h at 37°C) in growth-inhibitory substrate experiments. Cells were allowed to grow for 24h at 37°C in a 5% CO₂ atmosphere.

Neuronal cortical culture

E16.5 WT mice embryos were used for neuronal cortical cultures. Brains were extracted, washed in cooled 0.1M phosphate-buffered saline (PBS; ThermoFisher Scientific) containing 6.5 mg/ml glucose (Sigma Aldrich) (PBS-glucose) and the meninges excised. Both cortical lobes were dissected, sliced in a McIlwain tissue chopper (Capdem Instruments) and trypsinized for 15 min at 37°C. The suspension was inactivated with normal horse serum (NHS; ThermoFisher Scientific), incubated with 0.025% DNase (Roche) PBS-glucose for 10 min at 37°C and mechanically dissociated. Single cells were spun down and resuspended in Neurobasal™ (ThermoFisher Scientific) medium supplemented with 2mM glutamine (ThermoFisher Scientific), P/S, 6.5mg/ml glucose, NaHCO₃ (Merck), 1x B27 and NHS 5% and plated in poly-D-lysine (0.1mg/ml) pre-coated microfluidic devices (see below). One day after seeding, culture media was changed, and NHS was excluded from the new media. Cells were maintained at 37°C in a 5% CO₂ atmosphere and culture media was changed every two days.

Microfluidic devices

The design used for microfluidic devices was a modification of a previously published device (Taylor et al., 2005; Sala-Jarque et al., 2020). The resulting device consists in 4 receptacles of 7mm diameter, connected in pairs by a longitudinal compartment (cell body and axonal compartments) which are in turn interconnected by 100 microchannels (10µm x 10µm x 900µm). The master was produced using standard photolithography techniques at IBEC Microfab Space. Poly(dimethylsiloxane) (PDMS; Dow) was used to prepare the devices by soft lithography, which were subsequently attached to glass bottom dishes using oxygen plasma treatment. Neurons were seeded in one of the larger compartments (cell body compartment) and allowed to extend their axons across the microchannels. Vacuum aspiration in the axonal compartment allowed complete axotomies (Taylor et al., 2005).

In vitro lentiviral production and infection

LV-EF1α-hChR2(H134R)-EYFP-WPRE (LV-ChR2) produced in our laboratory were used to introduce ChR2 expression in primary cortical neurons. For LV production, 293-FT (ATCC) cells were simultaneously co-transfected with three plasmids: pMD2.G (VSV-G envelope expressing plasmid), psPAX2 (lentiviral packaging plasmid), and pLenti-EF1α-hChR2(H134R)-EYFP-WPRE (transfer plasmid) in Opti-MEM (ThermoFisher Scientific) using Lipofectamine 2000 Transfection Reagent (). Transfection media was changed for culture media (Advanced DMEM (ThermoFisher Scientific) supplemented with 10% FBS, 1% P/S and 0.5% glutamine) 6h after. Media was recovered at 48h and 72h post-transfection, filtered and centrifuged at 1200g to remove debris. The supernatant was then centrifuged at 26.000rpm for 2h at 4°C in a Beckman conical tube containing a sucrose cushion (20%) for purification, and the viral pellet was finally resuspended in PBS-1% BSA. Cortical neurons were infected at 4 days *in vitro* (DIV) for 24h, and high levels of YFP fluorescence could be observed at 7DIV, indicating positive infection.

In vitro optogenetic stimulation

Thy1-ChR2 DRG neurons or cortical neurons infected with LV-ChR2 were used for *in vitro* optogenetic stimulation experiments. A 470nm emission LED array (LuxeonRebel™) under the control of a Driver LED (FemtoBuck, SparkFun) of 600mA and a pulse generator PulsePal (OpenEphys) (Siegle et al., 2017) was used to deliver blue light (470nm) to neuronal cultures. The

optogenetic stimulation protocol consisted in 1h of illumination at 20Hz of frequency with 5ms-45ms pulses, in 1s ON-1s OFF periods. In the case of DRG neurons, stimulation was applied 2h after seeding. For cortical neurons, two different stimulation time-points after the axotomy were assessed: 30 min after axotomy and 6h after axotomy.

Corticospinal tract and dorsal columns axotomy

A fine incision to the skin between the shoulder blades of anaesthetized mice and subsequent muscle removal allowed thoracic vertebral column exposure. A T9 laminectomy was performed, and the dura mater was excised. In corticospinal tract injuries, two thirds of the spinal cord approximately were severed laterally with a scalpel. In dorsal column axotomies (DCA), a dorsal hemisection was conducted with fine forceps.

Sciatic nerve crush (SNC)

The sciatic nerve was exposed after a small incision on the skin over the posterior hindlimb and blunt dissection of the gluteus maximus and the biceps femoralis. Fine forceps were used to carefully compress the nerve orthogonally 2x10s. Animals were allowed to recover for 24h, when were then sacrificed and the sciatic nerve and the sciatic DRGs (L4, L5, L6) were dissected and processed.

***In vivo* optogenetic stimulation**

An optic fiber cannula (1.25mm, 0.22 NA; Thorlabs) was implanted in the motor cortex (M1) of Thy1-ChR2 transgenic mice by stereotaxic surgery (-1mm antero-posterior (AP), 1.5mm lateral to bregma, 0.5mm deep, (hind limb innervation region)) prior to injury and was fixed to the cranium with screws and dental cement. At 7 days post-injury (DPI) animals started receiving daily optogenetic stimulations consisting in 1h of illumination with 470nm blue light, at 10Hz of frequency with 10ms pulses, in 1sON-1sOFF periods, for 5 consecutive days, 2 resting days followed by 5 more days. The illumination was delivered through the optic fiber cannula which was coupled to a 470nm wavelength LED source (M470F3, Thorlabs) controlled by a pulse generator (Pulse Pal, Open ephys) (Siegle et al., 2017). The control group went through the same procedures than the experimental group without receiving illumination.

***In vivo* chemogenetic stimulation**

The commercial AAV5-hSynhM3D(Gq)-mCherry or the control virus AAV5-hSyn-mCherry (Addgene) were injected into the sciatic nerve of WT C57BL/6J mice (1μL/sciatic nerve) 4-5 weeks before the experiment. For chemogenetic stimulation, the animals received two daily intraperitoneal (i.p.) injections of 5mg/kg CNO (Tocris). After DCA, injections were delivered starting from 3DPI and lasted 7 consecutive days. In the case of SNC, the injections were given prior to injury: from 3 days pre-injury to the same day of the injury (in total 4 days of chemogenetic stimulation).

Behavioral assessment of sensorimotor function

Sensorimotor deficits and recovery were evaluated using the gridwalk and the BMS (Basso Mouse Scale) tests at -1, 1, 7, 14, 21, 28 and 35 DPI for CST injuries and at -1, 1, 3, 7, 10, 21 and 28 DPI for DCA. During the gridwalk animals were recorded while walking three times on a grid of 1 x 1 cm squares (total longitude of the grid: 50cm). The number of missteps in relation to the total number of steps was blindly quantified. In BMS evaluations mice were allowed to freely move in an open field and a score was assigned to each of them according to the BMS scale(Basso et al., 2006). In brief, this

157 scale grades from 0 to 9 the locomotor capacity of the hind limbs depending on several aspects such as
158 ankle movement, paw placing and position or stepping.

159 **Tracing of injured spinal cords**

160 A fluorescent tracer (10% Dextran AlexaFluor 594, ThermoFisher Scientific) was injected with a
161 stereotaxic frame (KOPF) into the motor cortex of animals with injured CST at 35DPI using the fibre
162 optic cannula hole, in order to trace the stimulated CST. The tracer was injected at 0.2 µl/minute for a
163 total of 1 µl, adding 5 more minutes at the end to avoid liquid spillage. 5d were waited before sacrifice,
164 to allow the tracer to be reach the axonal terminations

165 **Immunocytochemistry (ICC)**

166 For immunocytochemistry (ICC), cells were fixed with 4% paraformaldehyde (PFA) 15 min on ice,
167 washed and incubated in blocking solution (1% Bovine serum albumin (BSA), 0,25% Triton X-100
168 (0,25% Tx) in PBS (0.1M)) for 1h at RT. Primary antibody (anti-βIII tubulin (Tuj1, 1:1000,
169 BioLegend), c-Fos (1:200, Cell Signaling), ChR2 (1:500, Progen), GFP (1:500, ThermoFisher
170 Scientific) was added to the cells in blocking solution and incubated O/N at 4°C. After washing, the
171 cells were incubated 1h at RT with Alexa-Fluor-conjugated secondary antibodies (568 goat and 488
172 goat and donkey).

173 **Tissue processing for immunohistochemistry**

174 Anaesthetized mice were transcardially perfused with ice-cold 4% paraformaldehyde (PFA) and the
175 spinal cord, the brain or the DRGs were dissected and allowed to post-fix in PFA 4% for 24h at 4°C.
176 In peripheral experiments the sciatic nerves and the DRGs were directly dissected, as perfusion was
177 not needed, and allowed to fix in 4% PFA for 2h on ice. Fixed tissues were cryoprotected in 30%
178 sucrose in PBS for 24h at 4°C, brains were then directly frozen and sliced at 30 µm with a freezing
179 microtome (2000R Leica), while spinal cords and DRGs were embedded in tissue freezing medium
180 (OCT, Sigma Aldrich), frozen and 18 µm or 10 µm slices respectively were obtained using a cryostat
181 (CM 1900 Leica) and directly mounted. For whole mount stainings cryprotection was not needed,
182 instead tissues were kept in PBS.

183 **Immunohistochemistry (IHC)**

184 Prior to IHC, brain slices were maintained in cryoprotection solution (30% glycerol, 30% ethylene
185 glycol, 40% PBS). Brain IHCs were performed directly on free floating slices. Slices were blocked for
186 1h at RT (10% FBS, 0.5% Tx, 0.2% gelatine, 0.2M glycine in 1xPBS) and incubated with primary
187 antibody O/N at 4°C (5% FBS, 0.5% Tx, 0.2% gelatine in PBS; GFP (1:500, ThermoFisher Scientific).
188 Slices were then repeatedly washed with PBS-0.5% Tx, incubated with secondary antibodies (Alexa
189 Fluor 488 donkey) and Hoechst.

190 IHCs were performed directly on spinal cord preparation. Slides were blocked for 1h at RT (Blocking
191 solution: 8% BSA, 0.3% Tx in TBS) was added and incubated for 2h at RT, followed by O/N incubation
192 of primary antibody (GFAP (1:500, Dako) in 2% BSA, 0.2% Tx in TBS at 4°C. Secondary antibodies
193 (Alexa Fluor 488 donkey) were added after washing with TTBS and incubated 1h at RT.

194 Whole-mount IHCs were performed for sciatic nerves. Blocking solution (8% BSA, 1% Tx and 1/150
195 mAbIgG (Fabs) in TBS) was added and incubated for 2h at RT, followed by 3 O/N of primary antibody
196 incubation (SCG-10 (1:1000, Novus Biologicals) in 2% BSA, 0.3% Tx in TBS at 4°C. Secondary

197 antibodies (Alexa Fluor 488 donkey) were added after washing with TTBS and incubated 1 O/N at
198 4°C.

199 DRG preparations were stained with Hoechst (Sigma Aldrich).

200 Each preparation was subsequently mounted in MowiolTM.

201 **Fluorescence intensity analysis**

202 To measure c-Fos intensity, DRG neurons were immunostained for c-Fos and ChR2 and imaged at 40x
203 magnification with an Olympus microscope IX71 using an Orca Flash 4. Only ChR2⁺ cells were
204 selected for this analysis. About 20 cells per well were analysed. The nuclei of the cells were selected
205 and its mean c-FOS intensity determined by subtracting the corresponding background value to the
206 integrated density (Corrected total cell fluorescence; CTCF), measured using ImageJTM.

207 **Neurite and axonal length analysis**

208 Images were taken at 10x magnification with an Olympus microscope IX71 using an Orca Flash 4 or
209 a CX50 Olympus camera. TUJ1 or GFP was used to immunolabel neurites and axons. For DRG neurite
210 analysis, three fields per well were analysed and the average neurite length per cell was obtained. Small
211 diameter (<35µm) neurons were excluded from the analysis. For cortical neuron cultures, 8-9 fields
212 per device were analyzed. Either average axon length or total area covered by axons was quantified.
213 Neurite-J plugin in ImageJTM software was used for neurite and axonal measurements (Meijering et
214 al., 2004). Area measurements were measured using ImageJTM software.

215 **Sprouting quantification**

216 To quantify the number of sprouting axons before and after the lesion, we measured the tracer
217 fluorescence intensity on sections at 0,5mm pre-injury and post-injury level. The spinal cord section
218 images were divided in two different ROIs corresponding to grey matter and white matter for analysis.

219 **Nerve regeneration analysis**

220 Regenerating sensory axons were immunolabeled with SCG10. Whole mount preparations were
221 imaged with a confocal microscope (LSM 800, AxioCam 503c; Zeiss) at 5x magnification. 6-8 tiles
222 and 10-15 slices were obtained per nerve and the ZenTM software and ImageJTM were used to
223 reconstruct the nerve and obtain a Maximum Intensity Z-projection. The number of regenerating axons
224 at several distances from the crush was determined.

225 **Statistical analysis**

226 Prism 6.0 (GraphPadTM Software) was used for statistics and graphical representation. Plotted data
227 shows mean±s.e.m (standard error of the mean). Normality was determined with Shapiro-Wilk test.
228 Significant differences are indicated by asterisks (* $p < 0.05$; ** $p < 0.01$; *** $p < 0.005$; **** $p < 0.001$).
229 ANOVA followed by Bonferroni post hoc test or Student's t-test were used in normal distributions as
230 opposed to Mann-Whitney or Kruskal-Wallis tests as non-parametric tests for samples that did not
231 meet normality.

232 **3 Results**

233 ***Optogenetic stimulation of DRG neurons increase their axonal growth in vitro***

234 We first sought to determine if optogenetic activation of adult DRG neurons could enhance axonal
235 growth *in vitro*. Adult dissociated Thy1-ChR2 DRG neurons were optically stimulated and allowed to
236 grow for 24h.

237 Optical stimulation (Fig 1A) effectively increased neuronal activity in DRG neurons, as displayed by
238 the increased levels of c-Fos staining after stimulation (49627 ± 4373 a.u. of nuclear CTCF intensity for
239 stimulated DRGs versus 33897 ± 3212 a.u. of nuclear CTCF intensity for non-stimulated; $p=0.0058$
240 Student's T test) (Fig 1B-C)

241 Optically stimulated neurons showed increased neurite outgrowth ($4053 \pm 433.1 \mu\text{m}$) after 24h of culture
242 when compared to non- stimulated ones ($2029 \pm 647.7 \mu\text{m}$; $p=0.0267$ Student's T test) (Fig 1D-E)

243 ***Chemogenetic stimulation of DRG neurons improve their regenerative capacity after sciatic nerve*** 244 ***crush***

245 We wanted to determine if the *in vitro* results in fact translated to enhanced axon regeneration *in vivo*
246 after PNS injury. Due to the difficulty to apply light for long periods of time in awake animals in the
247 DRGs, we switched the activity control tool to chemogenetics.

248 We first wanted to determine if DRGs were transduced *in vivo*. AAV5-hSyn-mCherry or AAV5-hSyn-
249 hM3Dq-mCherry were carefully injected into the sciatic nerve and mCherry expression was examined
250 4-5 weeks later. Both vectors transduced DRG neurons with similar efficiency (~35% of total DRG
251 neurons; ~65% of large diameter ($>35 \mu\text{m}$) neurons. mCherry expression was mainly localized in the
252 soma of large diameter DRG neurons (Fig 2A).

253 To assess the effects of neuronal activity on PNS regeneration, animals received 2 daily injections of
254 5mg/kg CNO 3 days before performing a bilateral SNC. 24h after injury hM3Dq stimulated animals
255 showed increased number of regenerating sensory axons (SCG10⁺) at further distances ($>750 \mu\text{m}$) when
256 compared to mCherry controls. The two-way ANOVA showed a significant effect of the distance ($p<$
257 0.0001) and stimulation ($p=0.0145$) as well as their interaction ($p=0.0101$). At $750 \mu\text{m}$ from the injury
258 stimulated animals showed 27.67 ± 4.19 axons versus 12.78 ± 3.34 axons on non-stimulated animals
259 ($p=0.0694$ Bonferroni post Hoc test), at $1000 \mu\text{m}$ from the injury stimulated animals showed
260 21.56 ± 3.72 axons versus 6.44 ± 1.43 axons on non-stimulated animals ($p=0.0163$ Bonferroni post Hoc
261 test), at $1250 \mu\text{m}$ from the injury stimulated animals showed 14.89 ± 3.12 axons versus 3.11 ± 1.14 axons
262 on non-stimulated animals ($p=0.0261$ Bonferroni post Hoc test), and at $1500 \mu\text{m}$ from the injury
263 stimulated animals showed 8.56 ± 1.81 axons versus 1.56 ± 1.06 axons on non-stimulated animals
264 ($p=0.0259$ Bonferroni post Hoc test) (Fig 2 D-E).

265 ***Optogenetic stimulation of cortical neurons does not improve the sensorimotor performance after*** 266 ***SCI***

267 We then wanted to test if this increased regeneration capacity also was present in the CNS neurons. To
268 this aim we first evaluated regenerative capacity of cortical neurons *in vitro* after optogenetic
269 stimulation. Embryonic cortical neurons were seeded in custom microfluidic devices as previously
270 described, and ChR2 was induced by infection with LV-ChR2 (Supp fig 1A), then axons were allowed
271 to grow for few days through the microchannels until they reached the other chamber, when a vacuum
272 assisted axotomy was performed. Optogenetic stimulation of cortical neurons decreased the axonal
273 regrowth when stimulated 30 minutes after the axotomy (Supp Fig 1B). In contrast, 6 hours after
274 axotomy, optogenetic stimulation resulted in a non- significant ($p=0.0759$; Student T-Test) increase in

axonal regrowth when compared to non-stimulated ones (Supp Fig 1 C-E). To control the intrinsic effect of blue light on neuronal growth, we applied the light stimulation pattern on non-stimulated (GFP controls) embryonic cortical neurons and observed no difference in axonal growth (Supp. Fig 1D).

To test if this results further translated to a better motor performance after SCI *in vivo*, we implanted optic fibre cannulas on Thy1-ChR2 animals, that have the cortical expression of ChR2 predominantly concentrated in the layer V, where corticospinal motor projecting neurons lay (Supp. Fig 2). After recovery, animals were subjected to a CST axotomy, and optical stimulation was performed daily from day 7 after injury (Fig 3A). Stimulated animals did not show any improvement in their motor performance on the Gridwalk (Fig 3B) or the BMS (Fig 3C) when compared to non-stimulated controls. For each test evaluated, the two-way ANOVA showed a significant effect of the time (BMS $p < 0.0001$; Gridwalk $p < 0.0001$) but neither from the stimulation (BMS $p = 0.7201$; Gridwalk $p = 0.1308$) nor from their interaction (BMS $p = 0.9905$; Gridwalk $p = 0.0707$). In all tests, non-significant changes were observed for each timepoint when compared stimulated versus non-stimulated mice (Bonferroni multiple comparisons).

These results indicate that most likely CNS neurons have the ability to increase their axonal growth capacity when stimulated, but there are inhibitory signals in the CNS injury that blocks this regeneration *in vivo*.

Chemogenetic stimulation of DRG neurons does not improve their sensorimotor performance after SCI

To further address this issue, we tested the effects of chemogenetic stimulation on DRG neurons after a dorsal hemisection *in vivo*. Similarly, to what was done for the PNS injury, AAV5-hSyn-mCherry or AAV5-hSyn-hM3Dq-mCherry were carefully injected into the sciatic nerve to induce its expression on the DRG. 4-5 weeks later animals were subjected to a dorsal hemisection, and 3 days after injury animals received 2 daily injections of CNO (5mg/kg b.w.) for 7 days (Fig 3 D). As observed for the cortical stimulation, stimulated animals did not show any improvement in their motor performance on the Gridwalk (Fig 3B) or the BMS (Fig 3C) when compared to mCherry-veh, or mCherry-CNO controls. For each test evaluated, the two-way ANOVA showed a significant effect of the time (BMS $p < 0.0001$; Gridwalk $p < 0.0001$) but neither from the stimulation (BMS $p = 0.2289$; Gridwalk $p = 0.0694$) nor from their interaction (BMS $p = 0.3614$; Gridwalk $p = 0.0785$). In all tests, non-significant changes were observed for each timepoint when compared stimulated versus non-stimulated mice (Bonferroni multiple comparisons).

Highlighting again the presence of an inhibitory environment in the CNS injury that blocks the regeneration induced by neuronal stimulation.

Optogenetic stimulation of dorsal root ganglia neurons increase their axonal growth in vitro, only on permissive substrates

To further assess the impact of CNS inhibitory signals in the activity-induced increased axonal growth, we cultured adult DRG neurons in both permissive (Laminin) and inhibitory (CSPG) substrates and subjected them to optogenetic stimulation.

As previously observed (Fig 1D-E), optogenetic stimulation increased the neurite outgrowth in permissive substrates ($3682 \pm 316.2 \mu\text{m}$ for stimulated DRGs versus $2721 \pm 183.1 \mu\text{m}$ for non-stimulated DRGs; $p=0.0128$ Student's T test), but not on neurons seeded on CSPGs ($86.4 \pm 13.4 \mu\text{m}$ for stimulated DRGs versus $73.04 \pm 10.27 \mu\text{m}$ for non-stimulated DRGs; $p=0.4381$ Student's T test), where optogenetic stimulation did not produce any change in neurite outgrowth (Fig4 A-B).

Optogenetic stimulation of corticospinal motor neurons increases prelesion sprouting but does not induce regeneration across the lesion.

Since activity dependant therapies have been shown to increase lateral sprouting and plasticity (Engesser-Cesar et al., 2007a; Goldshmit et al., 2008a; Carmel et al., 2010b, 2013a; Carmel and Martin, 2014b; Sánchez-Ventura et al., 2021), we sought to investigate if our stimulation paradigm increased the sprouting of injured axons before the lesion. To this aim, we analysed the spinal cords of stimulated Thy1-ChR2 animals after tracing their corticospinal tracts. Interestingly, we found that optogenetic stimulated animals did show more axonal tracing in the gray matter (33.60 ± 1.13 a.u. of CTCF intensity for stimulated animals versus 23.27 ± 0.89 a.u. of CTCF intensity on non-stimulated animals; $p=0.0015$ Student's T test) in segments before (-0.5mm) the injury, but not beyond (+0.5mm) the injury (14.61 ± 1.21 a.u. of CTCF intensity for stimulated animals versus 12.80 ± 0.47 a.u. of CTCF intensity on non-stimulated animals; $p=0.30328$ Student's T test) (Fig 4C-E). These findings support once again that the positive effects of modulation of the activity on the axonal regrowth are inhibited by CNS inhibitory signals present in the injury.

4 Discussion

Neuronal activity-triggered plasticity is commonly recognized as the main component of recovery in current activity-based therapies (Carulli et al., 2011; Hogan et al., 2020), however, this is mainly built on therapies that use exercise or electrical stimulation to induce neuronal activity (as in (Engesser-Cesar et al., 2007a; Goldshmit et al., 2008a; Carmel et al., 2010b, 2013a; Carmel and Martin, 2014b; Sánchez-Ventura et al., 2021)). Even so, even though these approaches increase neuronal activity, they do so without cellular specificity, hindering the identification of underlying cellular and molecular mechanisms induced by neuronal activity itself.

Taking advantage of the cellular specificity of optogenetics and chemogenetics (Sternson and Roth, 2014; Deisseroth, 2015) we performed different experiments to assess the specific cellular effects of neuronal activity on axonal growth using different *in vitro* and *in vivo* neuronal models.

Consistent with previous works (Park et al., 2015a; Ward et al., 2016; Jaiswal and English, 2017; Sala-Jarque et al., 2020), we showed that increasing neuronal activity on DRG sensory neurons enhanced axonal growth in non-inhibitory *in vitro* and *in vivo* peripheral injury models. Additionally, and still accordingly to previous studies (Ward et al., 2018), we also demonstrated these effects in embryonic cortical cells *in vitro*, highlighting the presence of similar mechanisms among different neuronal types.

It is important to mention that before the achievement of these results, a fine tuning of the stimulation protocol was needed, as axonal growth capacities showed to be highly dependent on the timing after injury and the pulse frequency during the stimulation (data not shown), highlighting the fact that neuronal activity needs, not only to be stimulated, but to be finely regulated in order to promote the desired outcomes. This has also been emphasized in previous studies indicating activity-dependant effects on gene expression (Lee et al., 2017; Tyssowski et al., 2018; Miyasaka and Yamamoto, 2021) and plasticity (reviewed in (Fauth and Tetzlaff, 2016)).

In accordance, little changes in the stimulation patterns might lead to opposed effects, therefore, we used optogenetics, whenever possible, as our method of choice to stimulate activity since it allowed us a higher temporal resolution (Rost et al., 2017). Nonetheless, the anatomical setting of the DRGs impeded the implantation of a permanent optic fiber canula, and therefore did not allow us to perform awake stimulations, as this would have compromised the wellbeing of the animals, therefore, we opted for a chemogenetic approach for *in vivo* DRG stimulation. Effectively, this method rendered similar results on neurons as those observed *in vitro* with optogenetics, resulting in enhanced regeneration. For this experiment we delivered the DREADD agonist before the injury (as a preconditioning), leading to similar effects as those observed in previous studies using enriched environment (Hutson et al., 2019) or electrical stimulation (Udina et al., 2008b; Goganau et al., 2018b; Senger et al., 2018) as a preconditioning.

Since our results have shown comparable regeneration outcomes than electrical stimulations (Al-Majed et al., 2000, 2004; Ahlborn et al., 2007), we believe that neuronal activity itself might be responsible, at least partially, of these effects on axonal growth.

However, when we tested this paradigm in an *in vivo* model of SCI, we found that optogenetic stimulation of spinal-projecting cortical neurons did not promote the expected functional recovery after SCI, contrarily to what other studies using electrical stimulation (Carmel et al., 2010a, 2013b; Carmel and Martin, 2014a), or exercise (Engesser-Cesar et al., 2007b; Goldshmit et al., 2008b; Loy et al., 2018a) showed.

We then focused on clarifying whether our model did not induce any kind of axonal regeneration, or it did, but was insufficient to promote recovery. In that sense, an *in vivo* CNS model of injury implies the presence of a number of factors absent in our previous experiments, including the presence of extrinsic inhibitory substrates for regeneration (Mahar and Cavalli, 2018; Bradbury and Burnside, 2019; Mesquida-Veny et al., 2021). Accordingly, when we performed further histological characterization together with complementing *in vitro* experiments, we found that stimulated neurons displayed increased axonal growth in permissive substrates (laminin), but not in growth inhibitory substrates (CSPGs), and stimulated neurons showed an increased amount of growth and sprouting in the rostral vicinity of the injury, especially in the grey matter of the SC, but not caudally across or beyond the injury. These data suggests that neuronal activity stimulates the axonal growth capacity of these neurons through a mechanism that cannot overcome the repressive signalling of growth-inhibitory molecules, such as CSPGs. However, *in vivo* injured neurons start growing through uninjured areas of the SC, similarly to what is described in other studies with classical stimulation approaches (Carmel et al., 2010a, 2013b; Carmel and Martin, 2014a), where injured and uninjured axons grow axonal processes searching for spared intraspinal circuitry, these new connections will eventually, through neuronal plasticity, restore the injured connections, bypassing the injury (Courtine and Sofroniew, 2019a). Interestingly, classical non-specific stimulations such as epidural electrical stimulation or rehabilitation, lead to stimulation of intraspinal circuits, involving spinal interneurons and motoneurons, this stimulation in turn promotes the reorganization and activation of this circuitry (Wagner et al., 2018a). Whether the lack of these functional effects with our paradigm was due to a shorter stimulation period, or that it lacked the direct stimulation of this intraspinal circuitry and thus not inducing the plasticity needed, remains still unanswered.

Parallely, a recent study also observed a blockage of activity-induced axonal growth in the presence of inhibitory-substrates after cellular specific stimulation (Wu et al., 2020a). In this study, functional recovery after SCI was only achieved after combining chemogenetic stimulation and ChABC administration (a CSPG degrading enzyme) (Wu et al., 2020b), surprisingly, functional recovery was

only achieved after 6 weeks of continuous combined treatment. In contrast to both this work and our results, both chemogenetic stimulation and visual stimulation induce functional regeneration after optic nerve crush, a CNS injury model (Lim et al., 2016). Importantly however, the expression of the different chemogenetic receptors in this study was induced by intravitreal injection of the vectors, meaning that retinal interneurons, including amacrine and bipolar cells, will most likely be expressing the channels and be subjected to stimulation upon agonist administration. This paradigm resembles to that of electrical stimulation, considering activation of this retinal interneurons might be inducing neuronal plasticity that facilitates functional recovery.

There is plenty of evidence that neuronal activity itself is key in promoting regeneration and recovery, however, the presence of growth inhibitory substrates in the injured CNS limits the success of activity-based therapies. In line with this, combining functional rehabilitation or exercise with CSPG degrading therapies (including ChABC (García-Alías et al., 2009; Wang et al., 2011a) or ADAMTS4 (Griffin et al., 2020) lead to synergistic effects, even when applied at chronic time-points (Wang et al., 2011b). These reinforce the concept that multifactorial therapies are the way to go in order to tackle the different aspects of the pathophysiology of SCIs (Griffin and Bradke, 2020).

Although our experiments did not demonstrate the presence of an activity-triggered transcriptional switch, more chronic stimulations could be leading to it, although there are no evidences about that yet. Parallely, more local and transient cellular mechanisms might explain the growth differences observed in our paradigm stimulating activity specifically at a cellular level. For example, neuronal activity changes the excitability of neurons through reorganization of several ionic channels (Misonou et al., 2004; Enes et al., 2010a; Romer et al., 2016). This can in turn influence growth, as a result of intracellular ionic adjustments and their downstream associated signalling (Shim and Ming, 2010; Tedeschi et al., 2016), or even translational changes (Enes et al., 2010b). Neuronal activity can also alter neuronal metabolism, increasing glycolysis, and lipid synthesis and integration to the membrane, a cellular process essential during axonal extension (Bas-Orth et al., 2017; Segarra-Mondejar et al., 2018). Another important cellular component altered by activity is the cytoskeleton and its dynamics, for instance, neuronal activity has been shown to increase dendritic spine microtubule polymerization (Hu et al., 2008). Accordingly, chemogenetic stimulation increases microtubule dynamics in the distal axonal portion, by reducing tubulin acetylation and increasing tyrosination in this region (Wu et al., 2020b). These mechanisms may not be exclusive, on the contrary, they are likely to take place all together facilitating activity-induced axonal growth.

Success of activity-based therapies does not depend solely on the cellular effects of activity modulations, instead, these therapies, that target activity in a more general and chronic way (as different neurons or even circuits are stimulated simultaneously), benefit from a raised general excitability that seems to be ultimate the responsible for the plasticity and reorganization that leads to recovery (Courtine and Sofroniew, 2019b). In that direction, rehabilitation and electrical stimulation after injury promote the formation of new synapses in the spinal cord (Loy et al., 2018b; Eisdorfer et al., 2020), respiratory function recovery after optogenetic stimulation is also attributed to synaptic plasticity (Alilain et al., 2008), and a recent study showing that rehabilitation together with electrochemical stimulation promotes recovery, is credited to cortico-reticulo-spinal circuit remodelling (Asboth et al., 2018). Additionally, sprouting of spared axons, instead of injured ones, is also responsible for recovery after exercise and/or electrical stimulation (Engesser-Cesar et al., 2007b; Goldshmit et al., 2008b; Carmel et al., 2010a, 2013b; Carmel and Martin, 2014a). In agreement to that, success of these approaches is only significant in incomplete injuries, and greater as more tissue is spared. Incomplete injuries might also help explaining why in our model we did not observe recovery after optogenetic stimulation of the motor cortex while others did (Deng et al., 2021), as different spinal injuries were

used. Compellingly, compression injury leaves more uninjured tissue and spinal tracts than our injury, that axotomizes all the dorsal tracts, including the main component of the CST in mice. Besides, complete injuries also present larger glial scars, and therefore greater accumulation of growth inhibitory molecules, which translates in larger distances and hurdles to overcome or bypass to achieve functional connections, together with a greater loss of intraspinal circuits.

These observations strengthen the view that current activity-based therapies stimulate plasticity on top of inducing regeneration in specifically stimulated neurons, but probably without inducing long-lasting cellular reprogramming. This plasticity results from activity-triggered local changes, therefore prolonged stimulations are more effective increasing growth or regeneration (Courtine and Sofroniew, 2019a). This is also evidenced by the presence of functional recovery after 12, but not after 4 weeks of chemogenetic stimulation (Wu et al., 2020b). Therapeutically, holistic more unspecific approaches, have a more potent effect than cellular specific stimulations alone, as evidenced by enriched environment compared to chemogenetic stimulation, which presents a lower rate of regeneration (Hutson et al., 2019). Studies have also shown that this activity-induced plasticity can be accelerated and improved by linking the activation of different relays topographically in a system (McPherson et al., 2015; Mishra et al., 2017; Wagner et al., 2018b). These systems are however more challenging to define underlying mechanisms and understand the true nature of the gains of these therapies.

Briefly, we found that specific cellular stimulation of neuronal activity induced axonal growth, but only in the absence of inhibitory substrates. Likewise, this approach seems to be therapeutically less efficient in enhancing recovery than other more chronic or general stimulations. This also suggests and strengthens the idea that activity-based therapies succeed because of local transient cellular changes coupled with neuronal plasticity, rather than resulting in a cellular reprogramming of growth capacities, and that longer stimulation periods elicit more robust responses.

5 Conflict of Interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

6 Author Contributions

FMV performed, designed experiments, performed data analysis, and wrote the manuscript; SMT performed and designed experiments and performed data analysis; JADR supervised experiments, provided experimental funds and edited the manuscript; AH performed, designed experiments, performed data analysis, provided experimental funds and wrote the manuscript.

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10 Figure legends

Figure 1. Increased neuronal activity promotes neurite outgrowth. A. Diagram of the optogenetic illumination protocol. B. Representative images of c-Fos (red) and ChR2 (green) immunostaining. Scale bar: 50µm. C. c-Fos expression in the nucleus is increased just after optogenetic stimulation of Thy1-ChR2 DRG neurons (n=20-25 cells). MFI: mean fluorescence intensity. a.u.: arbitrary units. Data are expressed as mean nuclear fluorescence intensity±s.e.m. $^{**}p < 0.01$ denotes significant differences in Student's t-test. D. Stimulated neurons presented significantly higher neurite lengths when compared to non-stimulated ones. Data are expressed as average neurite length per neuron ±s.e.m 24h *in vitro* (n=7 wells from 3 different experiments). $^{*}p < 0.05$ denotes significant differences in Student's t-test. E. Representative images of Tuj-1 staining used to analyze neurite length. Scale bar: 500µm.

Figure 2. Chemogenetic stimulation induced PNS regeneration. A-B. mCherry expression in AAV-mCherry and AAV-hM3Dq infected DRG neurons 4-5 weeks after viral injection. Large-diameter DRG neurons are preferentially infected. Data are expressed as mean % of large-diameter (diameter >35µm) infected neurons±s.e.m. (n=7-9 DRGs per group). Scale bar: 100µm. C. Schematic timeline of the experiment. D. The number of regenerating sensory axons (SCG-10⁺) in stimulated nerves (hM3Dq-CNO) is increased in all assessed distances compared to the non-stimulated (mCherry-CNO) reaching statistical significance in long distances. Data are expressed as mean±s.e.m. at each distance from the injury site (n=9 sciatic nerves per group). $^{*}p < 0.05$ denotes significant differences in ANOVA followed by Bonferroni post-hoc test. E. SCG-10 immunostaining of mCherry or hM3Dq infected sciatic nerves 24h after SNC. Dotted lines indicate the injury site. Scale bar: 200µm

Figure 3. Increasing neuronal activity does not induce recovery after CNS injury. A. Timeline of the CST injury and stimulation experiments. B-C. % of missteps in the gridwalk test (B) (n=8-9 mice per group for each timepoint) and BMS score (C) (n=10 mice per group for each timepoint) show no differences in sensorimotor recovery in stimulated Thy1-ChR2 (ChR2 Light) mice when compared to non-stimulated (ChR2 No light) after CST injury. D. Timeline of the DCA and stimulation experiments. E-F. Chemogenetically stimulated animals (hM3Dq-CNO) show similar sensorimotor function recovery to non-stimulated ones (mCherry-CNO, mCherry-veh) after DCA as seen by the gridwalk (E) (n=3-5 mice per group for each timepoint) and BMS (F) (n=5-6 mice per group for each timepoint) tests. Represented data correspond to the BMS score and % of missteps in the gridwalk. Data are expressed as mean±s.e.m. ANOVA followed by Bonferroni post-hoc test.

Figure 4. Neuronal activity induces growth but is not sufficient to overcome inhibitory environments. A. Neurite outgrowth was significantly increased in stimulated DRG neurons on growth-permissive substrates (LAM: laminin), but not over non-permissive substrates (CSPGs). Average neurite length per neuron was determined at 24h *in vitro*. Data are expressed as mean ±s.e.m (n=11-23 images from 3 different experiments). $^{*}p < 0.05$ denotes significant differences in Student's t-test. B. Representative images of Tuj-1 staining used to analyze neurite length. Scale bar: 200µm. C-D. Tracer intensity quantification of injured stimulated and non-stimulated Thy1-ChR2 mice at 0,5mm pre-injury and post-injury. The spinal cord was divided in grey matter and white matter for analysis. Optogenetic stimulation of the CST resulted in increased sprouting in the grey matter pre-injury, but not post-injury. a.u.: arbitrary units. Data are expressed as mean tracer intensity±s.e.m (n=5 spinal cords per group). $^{**}p < 0.01$ denotes significant differences in Student's t-test. E. Representative images of traced (Dextran⁺) spinal cords. Scale bar: 150µm

Supplementary Figure 1. Optogenetic stimulation of cortical neurons after axotomy. A. Cortical neurons (Tuj-1) express ChR2-eYFP after LV-ChR2 infection. White arrows in the high magnification image indicate neurite expression of ChR2. Scale bar: 250µm. B-C. Optogenetic stimulation of cortical neurons 30min after axotomy (B) resulted in reduced axon regeneration, while delivering the stimulation 6h after axotomy (C) increased axon regeneration without reaching statistical significance. Individual ChR2⁺ axon lengths were quantified (B: n=16 images; C: n=7-15 images). Data are expressed as mean±s.e.m. *** $p < 0.001$ denotes significant differences in Student's t-test. D. Optogenetic stimulation (ChR2 Light) of WT cortical neurons did not induce any changes in axonal regeneration. Total growth area/microchannel was computed (n=7-8 images corresponding to 4 different devices). Data are expressed as mean±s.e.m. E. Representative images of GFP staining used to analyze axon regeneration when stimulation is applied 6h after axotomy. Scale bar: 200µm.

Supplementary Figure 2. ChR2 expression in neocortex of Thy1-ChR2 mice. While some layer II-III neurons express ChR2, expression is concentrated in layer V neurons. Scale bar: 100µm

Figure 1

Figure 2

Figure 3