Reversible and reusable compartmentalized thermoplastic chip for coculture of dorsal root ganglion neurons

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Abstract

Compartmentalized microfluidic chips play an important role in understanding the cellular mechanisms involved in neurodegenerative disorders. Dorsal root ganglia are a well-established model for modelling the peripheral nervous system (PNS), but their development on a chip remains limited. Furthermore, it would be beneficial for the devices to be openable in order to access the biological material inside for analyses. Easy to prototype and biocompatible, styrenic block copolymers (SBC) are an alternative to polydimethylsiloxane (PDMS) that offer both reversible and permanent bonding properties. This paper presents a fast and straightforward method to produce compartmentalized SBC chips. The study validates the culture of murine dorsal root ganglia explants, comparing it to the standard methods, to obtain a model of the PNS. Moreover, the reversible bonding properties of the SBC permit the reuse of the chip with a quick and easy cleaning protocol. It provides direct access to the cells, opening the way for imaging and molecular biology analysis. The comparison of the resources required to produce PDMS and SBC chips highlights the importance of moving to reusable devices. These detachable, easy-to-manufacture and sustainable all-thermoplastic platforms provide an alternative way of prototyping compartmentalized devices *for in vitro* PNS modeling.

Keywords: Microfluidic chip, Dorsal root ganglia, sensory neurons, compartmentalized culture, styrenic bloc copolymers, soft thermoplastic elastomers

1. Introduction

The reduction of animal numbers and their replacement by alternative experimental methods are two of the 4R principles (reduction, replacement, refinement, and responsibility) and constitute a major strategic ethical challenge, prompting the development of *in vitro* models that can be used instead of *in vivo* studies involving animal models. Despite the rise of microphysiological systems, in neuropharmacology, the complexity of the physiological processes and the long time scales might explain why only few neurological conditions have been modeled to date.[1] While industrial sectors have played a role in democratizing these tools, reducing their demand on laboratory resources and broadening their applications in neurobiology could further advance this effort.[2]

Microfluidic devices for compartmentalized culture have offered the possibility to study the neuronal functions and structures in a highly controlled way.[3,4] Robert Campenot was the first to develop a device that enabled local control of axons and somas.[5] The aim was to physically separate the cell bodies from the axons to mimic the diverse microenvironments that a neuron might encounter. Subsequently, a variety of microfluidic devices have been developed for neuronal studies. Taylor initiated a two-compartment microfluidic device connected by straight microchannels [6] for the study of axonal injury, regeneration and transport. Samson *et al.* employed a five-parallel culture chamber configuration to circumvent cross-contamination and to establish an environmentally isolated neuronal population with synaptically interconnected cells.[7] Courte *et al*. designed a microchip with arc-shaped microchannels to control the orientation and filtering of axonal growth.[8] Sharma *et al.* proposed a keyhole design to model human nerves,[9] while Park *et al.* developed central circular chamber designs to promote axon growth in microchannels and the formation of a denser axonal network layer.[10,11] Electrospun aligned nanofiber-integrated compartmentalized microfluidic neuron culture systems have been developed by Luo *et al.* to improve axon alignment, length and stability.[12] Two-dimensional (2D) and three-dimensional (3D) compartmentalized co-culture systems on microdevices have been developed to analyze the interactions between neurons and glial cells in both the central nervous system (CNS)[11,13] and in the peripheral nervous system (PNS).[9,11,14–16]

Most of the PNS models focus on motor neurons, while there are only a few studies on sensory neurons on chip.[17–20] However, these studies focus on co-cultures of sensory neurons and oligodendrocytes, a glial cell that is specific to the central nervous system. Dorsal root ganglia (DRGs) represent a well-established model for PNS modeling, but their development on a chip remains limited. They mainly focus on pain signaling[21] and therapeutics[22] as well as on "nerve-on-chip".[23,24] In 2023, Mutschler *et al*.[25] proposed the first co-culture of neurons from dissociated DRGs and Schwann cells on a compartmentalized microfluidic device. They highlighted the challenge of culturing DRGs as

explants in microfluidic chips. Notably, Park *et al.* were the only group to successfully culture a chick DRG explant in a compartmentalized microfluidic device.[26] To facilitate the seeding of the explant, they designed a device with an open-top chamber.

Compartmentalized systems and organ-on-chip devices are becoming more widespread in the field of Biology, requiring a re-evaluation of their sustainability and of the materials used in their manufacture for specific applications. Currently, polydimethylsiloxane (PDMS) is predominantly used due to its high oxygen permeability, while polystyrene (PS) is favored for large-scale production and alignment with biological standards.[27] However, there are no neurobiological applications that uses materials combining the advantages of PDMS and PS, such as styrenic block copolymers (SBCs).[28] Reversible bonding of SBCs would be an advantage for accessing tissues during culture and for reusing devices.[29–31] Moreover, as the organ-on-chip industry is expanding rapidly and access and production costs are becoming critical for laboratory resources and carbon footprint,[32] there is no known comprehensive assessment or methodology to evaluate the resources involved.

Here, we describe a fast and straightforward method for the fabrication of compartmentalized SBC chips and demonstrate their reversible bonding and reusability for the culture of compartmentalized peripheral nervous system model. We first propose a simple process for fabricating SBC chips with an axisymmetric design adapted to anisotropic compartmentalized culture, particularly suitable for primary DRG explants. We confirm the theoretical compatibility of the design with neuron culture using analytical and finite element models. The viability and axonal growth of $E13.5$ mouse embryonic DRGs within SBC and equivalent PDMS chips were then compared. We present a rapid and reliable cleaning procedure for SBC chips and evaluate their reusability. We also demonstrate the potential of reversible bonding chips for direct imaging using electron microscopy, as well as for biological analysis techniques (immunostaining, Western blot, RT-qPCR). Finally, we discuss the resources (energy, time, cost and CO_2 equivalent emission) involved in the production of a PDMS compared to an SBC chip.

2. Results & discussion

2.1. Development of a compartmentalized SBC chip for coculture of DRG neurons

We developed a design and a workflow to produce axisymmetric compartmentalized 6-plexed SBC devices. With the exception of the photolithography step, which is necessary for wide channels, the fabrication process of SBC microfluidic chips **(Figure 1A)** is an out-of-the-cleanroom process. The first phase was to replicate the SU-8 mold in an epoxy mold, which is more compatible with hot embossing. It was used for 2 years, embossing more than 200 chips without any visible damage. The replication of the molds was confirmed by profilometry *(***Figure SI.1***)*. These open-top devices are made of two layers that are simple to assemble thanks to the adhesive properties of SBC. A first $1\ mm$-thick

110 layer containing the structures and a $2\ mm$-thick reservoir layer to increase the volume capacity and
111 prevent rapid evaporation of the culture media (**Movie SI.1**) as well as shear stress during pipetting.
112 The central chamber without the reservoir can only hold 7 µL, whereas it reaches a volume of $63\ \mu L$
113 when the second layer is added. Due to the radial growth of axons in the DRGs, we designed a circular
114 central chamber with a diameter of $4\ mm$ to facilitate the penetration of axons into the microchannels
115 **(Figure 1B)**. The dimensions of the microchannels have been chosen to enable compartmentalization
116 of somas in the central chamber and axons in the peripheral chambers.[6,33] Treatments can be added
117 in the outer chambers without flowing back to the central chamber. The high fluidic resistance of the
118 microchannels creates a small but sustained flow between the chambers, thereby counteracting the
119 diffusion (**Figure SI.3***)*. This fluidic isolation allowed 6 independent peripheral chambers to be created.
120 The sTPE devices bond easily to polystyrene (PS) **(Figure 1C)**, making them adaptable to already existing
121 basic culture plates. The Young's modulus of the SBC polymer is $1.15\ MPa$[28] which is in the same range
122 as PDMS ($0.5 - 3\ MPa$). In addition, this material is biocompatible, gas permeable, optically
123 transparent and exhibits reversible bonding that can withstand $1\ bar$ even after 10 sealing/unsealing
124 cycles,[31] making it a suitable candidate for cell culture. The air plasma hydrophilization remains stable
125 in time for at least one week for the SBC chips. The culture of peripheral neurons was demonstrated
126 using either explants or dissociated embryonic DRGs harvested from $E13.5$ mouse embryos.

127 The device was also compatible with dissociated DRGs. It has been described that the radial
128 flow pressure resulting from the circular fluidic design facilitates the positioning of cells in close
129 proximity to the microchannels inlet.[10,11,34] Moreover, the circular design allows for axon coverage that
130 is more than 3.5 times higher than that of square designs.[10] It allows the cultivation of both dissociated
131 DRGs and explants while most of the DRGs cultured on chips are dissociated. Overall, this model
132 provides an experimental framework to study different aspects of Schwann cell development such as
133 proliferation, migration, differentiation, and myelination of axons.[35] These co-cultures of neurons,
134 Schwann cells and fibroblasts were placed directly into the central chambers and cultured within the
135 SBC chips bonded to 6-well PS plates. The cultures could be maintained for one month within the
136 devices, and no leakage or contamination was observed.

137 **2.2. A shear stress compatible with DRG culture on a chip**

138 In order to assess the shear stress experienced by the DRG in the chip, we performed a Finite
139 Elements Method (FEM) analysis and particle velocimetry. As demonstrated by Liu *et al.*, it is essential
140 to quantify the shear stress forces generated within the microchannels in order to evaluate the
141 suitability of the device for biological studies. We used the FEM to visualize the speed v and the shear
142 rate σ in a microchannel cross section **(Figure 2A)**, the two physical quantities being intrinsically linked
143 by $\sigma = \mu \nabla v$. The following values were obtained for a pressure gradient of $30\ \mathrm{Pa}$, which represents

the theoretical maximum pressure that can be applied within the microfluidic device under study, given the finite volume of the reservoirs. The maximum shear stress inside the cross section is $0.8\ \mathrm{dyn.cm^{-2}}$. A pressure gradient of $30\ \mathrm{Pa}$ is achieved when the central chamber is full and the peripheral chambers are empty. However, this extreme case never occurs in practice when cells are in the chips. During the media change, the pressure reached a maximum of $25\ \mathrm{Pa}$ for a few seconds, due to the presence of a small volume of media within the chambers. Experimental validation was performed by measuring the time required for $0.2\ \mu\mathrm{m}$ microbeads to cross the microchannels **(Figure 2B)**. The experiment was started with a pressure of $25\ \mathrm{Pa}$ and stopped after $9\ \mathrm{h}$, when the pressure had reached $0\ \mathrm{Pa}$. The flow rate (Q) inside the microchannels was calculated from the velocity of the microbeads and the area of the cross section of a microchannel (A): $Q = v \times A$. An exponential regression was fitted to the experimental results, with $v_{max} = 132.5\ \mu m.s^{-1}$ and a time constant of $\tau = 3.18\ h$. It was then possible to derive the shear stress inside the microchannels based on the following equation: $\sigma = \frac{6\eta Q}{h^2 w}$, where η was fixed as the viscosity of water in Pa.s. For a maximum pressure of 25 Pa, the mean shear stress obtained was 2.73 dyn.cm^{-2}.

2.3. Viability and axonal growth of DRG culture on a chip

In order to validate the DRG culture on our devices, we assessed the viability and the axonal growth over time and according to the material. For viability, the dissociated cells formed a rather uniform distribution of cells, thereby facilitating the Live/Dead staining. For axonal growth, the axisymmetric growth of the explant is directional, thereby facilitating the segmentation and the quantification of the network. DRGs at E13.5 are at a stage at which we can obtain a high yield of sensory neurons and Schwann cells. Both exhibit high viability and plasticity.[35] After the DRGs were picked from the extracted spinal cord, dissociated and seeded in PDMS and SBC chips, or in well plates as a control. Their viability was assessed at day *in vitro* (DIV) $DIV3$ and $DIV7$. Comprehensive tiles of each culture enable exhaustive quantification and better visualization **(Figure 3A**). All the cells were segmented and counted exhaustively (**Figure SI.6**). At $DIV3$, the cell density was below the expected seeding cell density due to the low survival rate of the primary cells during the seeding step. The number of live cells was significantly higher in the control than in the chips (**Figure 3B**), with more than 1800 cells.mm^{-2} in the 96-well plate against 600 cells.mm^{-2} in the PDMS chips, 880 cells.mm^{-2} in the SBC chips and 930 cells.mm^{-2} in the reused SBC chips at DIV7. However, there was no significant difference in the number of dead cells between all the conditions. The overall number of cells increased by $13\ \%$ in the control group between $DIV3$ and $DIV7$, compared to 64% in the PDMS chips. This rate was found to be similar between SBC and reused SBC chips, at $31\ \%$ and $35\ \%$ respectively. Similarly, DRG explants were fixed at $DIV3$, before the axons reached the microchannels, and at $DIV7$, after the axons had crossed the microchannels **(Figure 3C)**. The growth speed was deduced from the axonal

178 length (**Figure SI.7**). Due to the fluidic isolation, the antibodies were unable to diffuse from the
179 peripheral chambers into the microchannels. As the axons started to cross the microchannels around
180 $DIV4$, it was necessary to detach the SBC chips at $DIV7$ before staining the cells to fully observe them
181 in the peripheral chambers (**Movie SI.2***)*. However, due to the irreversible bonding of the PDMS, it was
182 not possible to obtain the length of the axon at $DIV7$ inside them. No significant difference was
183 observed between the axonal length in 4-well control and the axonal length in the chips **(Figure 3D)**.
184 From $\mathrm{DIV0}$ to $\mathrm{DIV3}$, the axonal growth speed was $443 \pm 33\ \mu\mathrm{m.day}^{-1}$ and decreased to a speed of
185 $324 \pm 36\ \mu\mathrm{m.day}^{-1}$ from $\mathrm{DIV3}$ to $\mathrm{DIV7}$.

186 This design enables a uniform treatment in the central chamber and a gradient treatment in
187 the peripheral chambers, due to the high resistance of the microchannels. This feature can be useful
188 for drug treatments; however it prevents antibodies to reach the microchannels during
189 immunostaining constraining us to open the chip to observe the entirety of the axons. In addition, the
190 open-top device facilitates media changes and the CO2 exchange, while minimizing mechanical stress
191 and shear stress compared to a closed-compartment device.[20] The stable hydrophilization was
192 especially convenient for transport and long-term use compared to PDMS. We compared the
193 maximum shear stress value obtained in our device to the maximum shear stress value that is
194 sustainable for cortical neurons. We considered the value of $5\ \mathrm{dyn.cm}^{-2}$ which was measured for a
195 $24\ \mathrm{h}$ exposure[36] as a reference, as to our knowledge there are no data on the effect of shear stress on
196 sensory neurons. The maximum experimental shear stress obtained in our device is 1.25 times higher
197 than the maximum theoretical shear stress. However, it is almost 2 times lower than the cell viability
198 limit for cortical neurons. The maximum shear stress in our device is only reached for a few seconds,
199 when the cells are inside, during the change of medium. An experimental study of the effect of the
200 shear stress on the cell viability would help identify the limit in the context of peripheral nervous
201 system (i.e. sensory neurons) but it would require a controlled flow rate. We numerically confirmed
202 that the axonal growth rate is not impacted by the microfluidic chip, assuming a comparable threshold
203 in our biological context. We made tiled images of the central chambers of the chips and of the wells
204 of the controls to confirm the viability of the cells and ensure that our microfluidic architecture does
205 not affect them. The tiles enabled us to monitor viability over time and confirm the homogeneous
206 spatial distribution of dead cells in the chips. In contrast, in a standard well plate, cells tended to
207 accumulate on the walls.

208 We had to adjust the number of cells seeded for each material to ensure the targeted seeding
209 density remained consistent. Even when using the same 3 mm puncher to open the central chamber
210 of the SBC and PDMS chips, the diameters of the holes obtained are not the same (**Figure SI.2**). This is
211 a consequence of the elongation at break of the SBC ($720\ \%$) compared to the PDMS ($76\ \%$) one.[27,31]

212 As the diameters are different, the seeding area is different for both materials. We have a surface area
213 of $7.1\ \mathrm{mm}^2$ for SBC devices and $3.14\ \mathrm{mm}^2$ for PDMS ones. However, only the open top parts are
214 different in size, the central chamber is still $4\ \mathrm{mm}$ in diameter for both materials, which explains why
215 there are fewer cells per mm^2 in the PDMS devices than in the SBC devices.

216 The consistently high viability (>80%) across conditions indicates that the cells generally survive
217 well under all tested conditions, which is critical for biological relevance and assay reliability. The
218 axonal growth rate in the chips is comparable to that of the control, indicating that the presence of
219 microchannels has no effect on axonal growth. It is also similar to that observed in the literature, with
220 a rate of around $400\mu\mathrm{m/day}$.[37] The lower cell density in the chips compared to well plates may reflect
221 differences in culture environment, such as nutrient availability or surface interactions, which can
222 influence cell proliferation behavior. The SBC material exhibits denser cell growth at DIV3 and DIV7 in
223 comparison to the PDMS, suggesting that it may offer a better support for cell proliferation. The fact
224 that the reuse of SBC material does not affect viability is encouraging for reproducibility and material
225 stability. In case of in vitro preclinical study, the viability remains relatively high for all three conditions
226 and networks neuron networks develop normally. Further phenotyping would be required to assess
227 advanced structural features such as myelination (Ki67 proliferation marker, c-fos stress marker, MBP
228 and MPZ myelin markers).

229

230 **2.4. SBC chips can be cleaned and re-used up to 3 times**

231 A rapid, simple and reproducible protocol was developed to clean and reuse the chip **(Figure**
232 **4A)**. We validated the regeneration of the chip using immunofluorescence, profilometry and electron
233 microscopy. We validated the compatibility of re-used chips using axonal growth assays. The process
234 was completed in less than $3\ min$ per device and required only milliQ water, soap, ethanol $70\ \%$ and
235 a toothbrush. Chips were sterilized again by UV-sterilization prior to culture. DRGs were grown up to
236 one month in reused chips. It was confirmed that there was no noticeable difference in the surface of
237 the SBC device before and after cleaning. Epifluorescence and SEM images were taken before use,
238 after use and after the cleaning **(Figure 4B)**. Due to the sensitivity of the sTPE to electron beam, cracks
239 appeared during SEM imaging. Using these two microscopy techniques, we confirmed that all the
240 cellular debris present after culture had been removed by the cleaning protocol. After the third
241 cleaning, it was observed that the axons were unable to enter the microchannels **(Figure 4C)**,
242 suggesting that they were obstructed at the entrance. This finding was confirmed by optical
243 profilometry measurement **(Figure 4D)**, which revealed that the microchannels were clogged. The
244 clogging can be readily observed with a brightfield microscope (**Figure SI.9**) and is therefore easily
245 detected. Consequently, each chip can be used up to 3 times before exhibiting irreversible deformation
246 and becoming inoperable. The SBC material also has intrinsic advantages for cell culture compared to

247 the PDMS in our context: the hydrophilization of SBC remains steady, allowing longer storage and
248 reduced risk of cells detachment. The tendency of PDMS to leach and absorb other material can also
249 result in issues when used in conjunction with other materials, particularly 3D printed materials or
250 hydrogels.[38] The use of SBC materials would facilitate the production of composite devices for more
251 complex applications.

252 **2.5. The reversible bonding of the SBC provides direct access to the cells**

253 The reversible bonding properties and detachment of the SBC microfluidic chips also allowed
254 direct access to the cells (**Movie SI.2**). Immunostaining of the cells was performed after the
255 detachment of the device without damaging the cells **(Figure 5A)**. We can clearly observe axons and
256 nuclei of Schwann cells and fibroblasts that had grown and migrated inside the microchannels. NFH is
257 an anti-neurofilaments-heavy chain while SMI312 is staining the neurofilaments-medium chains (NFM)
258 and NucRed the nuclei. Similar images were obtained on fixed cells by SEM after opening of the chips
259 **(Figure 5B)**. The cells were not damaged by the detachment or the drying process required prior to
260 metallization. The reversible bonding properties also allowed direct access to the cells, which could
261 then be scraped in order to realize WB and RT-qPCR (see **Figure 5C-D**). WB was performed after 11
262 days of culture by scraping DRGs either from a 4-well plate or from a 6-well plate after detaching the
263 SBC chips. Four DRGs were pooled together, and the results showed no significant difference in the
264 expression of SOX10, a protein expressed in Schwann cells, between the DRGs extracted from the
265 control and the SBC devices (**Figure 5C**). RT-qPCR was conducted on a single DRG explant scraped from
266 a well or a chip after 7 days of culture. The comparison of the amount of RNA recovered in control
267 wells and that recovered in the reversibly bonded chips revealed that the SBC chips provide more
268 material under equivalent culture conditions – see **Figure SI.14**. The expression levels of *Krox20*, *Mbp*
269 and *Dhh*, which are markers of the Schwann cell lineage, were found to be comparable in the control
270 and SBC chips (**Figure 5D**). Dhh and Mbp are expressed at the precursor level of Schwann whereas
271 *Krox20* specifically confirms the transition to immature Schwan Cells (prior to myelinating Schwann
272 Cells).[35]

273 Many reversible bonding strategies have recently been reported, for applications beyond cell
274 culture.[30] Most of them require the intervention of an external help, a change in the design or the
275 addition of an interlayer between the chip and the substrate. PDMS can be reversibly bonded in the
276 absence of plasma treatment, but it is not easily reproducible and leakage occurs rapidly. In this case,
277 axons can grow under the microchannels rather than inside them and the hydrophobicity of the PDMS,
278 which is usually made hydrophilic by plasma bonding, prevents liquid from easily entering the device.[29]
279 The self-adhesive properties of thermoplastic elastomers have also been used for reversible bonding[39]
280 and their reported maximum working pressure without plasma bonding is 3.4 times higher than that

281 of PDMS one.[40] Recent studies have shown that PDMS can exhibit reversible bonding properties after
282 plasma activation of the substrate, 0,1% APTES treatment and overnight bonding at 75°C[41]. However,
283 the SBC we use has self-adhesive properties that do not require any chemical treatments or design
284 changes. We have shown that it can be reused without impacting the cell culture and no leakage was
285 observed. Axons grew inside the microchannels and the SBC chips were hydrophilized by using plasma
286 treatment without impacting their reversible bonding. Cleaning the chips after use has been suggested
287 before,[42] but to our knowledge, has never been described. We proposed a fast and simple detailed
288 protocol for cleaning the devices after use, thus enabling them to be reused. However, the reusability
289 of the chips is limited in this particular context. After 3 uses, the microchannels were repeatedly
290 clogged, completely blocking the flow between compartments. We hypothesize that this could be due
291 to the uniform repetition of the different thermal cycles and plasma treatments, as the microchannels
292 clogging was identical and regularly distributed. For chips with thicker channels (in this case the
293 clogged channels were $2.8\ \mu\mathrm{m}$ thick), it is possible to exceed the $\mathrm{n} = 3$ limit.
294 Axons are not only growing inside the central chamber, but also in the reservoirs (**Figure SI.12**).
295 When the chip is detached, these axons are torn at the same time. However, most of the axons present
296 in the microchannels remain attached. The samples can be immunostained after fixation. The resulting
297 images were more contrasted, with a more uniform background than with the devices, due to the
298 contribution of the autofluorescence of the polymer chip. This direct access also allowed to achieve
299 direct collection for biological analysis of proteins (western blot), which has already been performed
300 using a detachable substrate device,[14] or nucleic acids (RT-qPCR), which has been carried out directly
301 on the chip.[43] The detachment of the chips allowed easy scraping of the cells to collect most of the
302 biological material. WB was performed on 4 DRGs pooled, experiments have been conducted on 1, 4
303 and 8 DRGs in controls to assess the number of DRGs necessary to extract sufficient proteins (**Figure**
304 **SI.13**). The results showed that there was no significant difference by using 4 or 8 DRGs. However,
305 there is almost 40% of signal intensity difference between the 1 DRG and 4 DRGs experiments,
306 supporting our choice to pull 4 explants together. RT-qPCR was performed by pulling 1 or 2 DRGs
307 explants and similar results were obtained, confirming that enough RNA was collected by scrapping 1
308 DRG to perform an experiment. Moreover, around $20\ ng/\mu L$ of RNA was extracted from 1 DRG which
309 would be sufficient to test approximatively 30 genes.
310

311 **2.6. Comparison of carbon footprint of SBC and PDMS chips fabrication process**

312 The energy and time needed to assemble an SBC and a PDMS device were calculated at each
313 stage of the fabrication process **(Figure 6A-B)**. The total energy required to fabricate an SBC chip was
314 $289.3\ kJ$, almost 2 times less than the molding step of a PDMS chip alone, which required $556.8\ kJ$.

PDMS chip fabrication took more time and consumed more supplies, resulting in more waste than an SBC chip. Moreover, an SBC device could be cleaned and reused 2 times before disposal. The cleaning step required no energy and took only 3 min, thereby reducing the time and energy needed for the fabrication process, as well as the cost and kg of CO2 generated **(Figure 6C** and **Table SI.1**). Compared to a PDMS chip, a reused SBC device took 4 times less energy and time to manufacture. The overall price for a single PDMS chip is €2.83, while the price of an SBC chip is €3.10. However, the price of the SBC device decreased to €1.72 when it was reused. To determine the carbon footprint generated by the SBC raw material production, we assumed that it was the average of the carbon footprint of polystyrene, polybutadiene and polyethylene production. The resulting value for one SBC chip is $0.013\ kg$ CO_2e while for a PDMS chip it is $0.025\ kg$ CO_2e. When an SBC device is reused, as there is no need for new SBC there is no more carbon footprint generated by the material fabrication. The overall carbon footprint has been calculated by considering the mass of CO_2 equivalent generated per chip for electricity consumption, polymer for chip and 6-well plate production, washing of the chip and disposal of the wastes by incineration. Most of the carbon footprint was attributed to the disposal and washing protocols and was similar for the three types of chips. However, the reused SBC devices produced $0.05\ kg$ CO_2e less than the PDMS ones because they consumed less electricity and didn't require the production of polymer.

Time, energy and carbon footprint generated for SBC and re-used SBC chips fabrication were lower than for PDMS ones fabrication. PDMS required more material and thus generated more waste and cost. Thermoplastics were cheaper to produce[44] but because PDMS is produced in larger quantities, its final price is lower. It should be noted that the calculations are based on low carbon electricity, coming mostly from nuclear power plants. Depending on the country of reference, electricity can affect both the price and the carbon footprint. Costs and carbon footprint could be lowered for both SBC and PDMS devices fabrication by reducing the amount of waste generated as they cause half of the mass of CO_2e produced. The quantities of ethanol and milliQ water could be minimized, and increasing the density of chips per mold would also reduce the amount of energy and time necessary to produce them. However, the amount of wasted SBC is easier to reduce than for PDMS because the SBC can be cut to the desired size whereas PDMS must be poured over the entire surface of the mold. When comparing all the environmental impacts from cradle to grave, normalized to those of PDMS, SBC and reused SBC chips appeared to be lower for all impacts, in particular for marine eutrophication and mineral resource scarcity (**Table SI.2**), confirming the interest of using SBC as an alternative to PDMS. In our work, we choose to focus only on the manufacturing process for comparison purposes, we did not take into account the labor cost and the equipment maintenance as they are similar, in our case, for both materials. However, the total energy required to produce one

349 chip can be considered negligible when compared to the energy consumed in the lab or the energy
350 required to produce and transport the material.[45–47] If we considered the fabrication of 300 chips over
351 a year, the carbon footprint associated with chip fabrication corresponded to 1/60th of the mean
352 carbon footprint of one researcher in one year ($6t_{eq\ CO2}$).[45] In summary, the SBC already shows a very
353 interesting reduction in time and cost for the prototyping phase and could have a significant impact
354 on the environmental impact of a product on an industrial scale. This analysis underlines the
355 importance of quantifying and optimizing the environmental impact of microfluidic devices, especially
356 in a field where there is an ongoing industrial scale-up that should intensify with the growing
357 rationalization of animal models.

358 ## 3. Conclusion

359 We introduced a detachable, reusable, easy to produce and equivalent in cell culture
360 performance SBC device for an original application to DRGs culture in the context of peripheral nervous
361 system study. We proposed a new design adapted to DRGs explants and their radial growth with 6
362 peripheral compartments. We estimated the shear stress in the chip using simulations and particle
363 velicometry and experimentally confirmed its ability to perform axonal growth and viability assays of
364 a murine peripheral nervous system model. Based on the above biological assays, the PDMS devices,
365 used as a control, showed equivalent performance to the SBC devices. The demonstration of growth
366 and maintenance of a primary co-culture of nerves, which are particularly sensitive to stress, in an SBC
367 chip paves the way for tissue diversification and species growth in SBCs for the neurofluidic
368 community, but more generally for the microphysiological systems community. The reversible bonding
369 of the chips has been exploited both for direct access to different contact imaging microscopy and
370 biological analysis techniques such as SEM, WB and RT-qPCR. The possibility to recycle our chips led
371 us to develop an original workflow to evaluate the resources (time, energy, financial costs and CO2)
372 involved in the production and recycling of the chip, which could be used by the microfluidic
373 community in general to initiate a deeper rethinking. Switching to a reusable SBC chip allowed a
374 significant reduction in all these resources, more than 4 times in time and energy compared to PDMS,
375 contributing to more sustainable research. The carbon footprint assessment methodology can be
376 reused for other devices.
377 In the field of the microphysiological systems, the paradigm to move away from PDMS toward soft
378 thermoplastics makes it possible to reduce the amount of resources used and to overcome the usual
379 limitations for cell culture. This could help minimize the environmental impact of a rapidly growing
380 industry. It also contributes directly to efforts in microfluidics to accelerate the clinical translation of
381 therapeutics for neuropathologies.[48,49] From an animal ethics perspective, the development of
382 alternative *in vitro* models has the potential to significantly reduce the animal use, in alignment with

383 the 4Rs commitment. In the case of DRGs, the number of embryos in a pregnant mouse is typically
384 between five and seven, with an average of 40 DRGs per embryo. This yields a potential 250
385 experiments to be conducted from a single mouse, depositing one dorsal root ganglion per chip. This
386 approach allows for the testing of up to 1,500 conditions using the proposed design, rendering it highly
387 valuable for drug screening and mechanistic investigation studies. Finally, the demonstration of viable
388 DRG culture on our chip paves the way toward myelinating coculture in the PNS, which would provide
389 2D myelin models for neuropharmacological explorations. This could be used to study of the processes
390 of myelination, demyelination, and remyelination, but also to select *in vitro* drugs that could be used
391 for regenerative treatments by stimulating the production of new myelin.

392

393 # 4. Experimental Section

394 **4.1. Chip design and fabrication**

395 *a) Molds fabrication*

396 The photomask designs were drawn using CleWin 5.2.4 version and printed on a SodaLime substrate
397 with a resolution of 128k dpi (JD Photo Data, UK). A first layer of 2.8 µm thick microchannels was
398 structured with SU-8 2002 (Chimietech, France) on a 4-inch diameter silicon wafer $1\ mm$ thick using
399 soft lithography. This was followed by the addition of a second layer of $100\ \mu m$ thick chambers made
400 with SU-8 2050 (Chimietech, France). The Epoxym Kit (EdenTech, France) was used to create a negative
401 replica in $1{:}10$ Sylgard 184 PDMS (Neyco, France), followed by an epoxy positive replica of the SU-8
402 mold with Gallon KIT Conapoxy FR-1080 (Ellsworth Adhesives, USA).

403 *b) Thermomolding and assembly of SBC chips*

404 The microfluidic chips were embossed from styrene block copolymer (SBC) foils (EdenTech, France) at
405 $160\ °C$ for $3\ min$ $(0.07 - 0.08\ MPa)$ using the Sublym hot press machine (EdenTech, France) and the
406 previously fabricated epoxy mold. The central chamber was punched with a $3\ mm$ punch to obtain an
407 open top device. A second layer of SBC embossed on a flat epoxy mold was added to each chip with a
408 $6\ mm$ diameter reservoir for the central chamber. The outer chambers were punched simultaneously
409 in the patterned layer and the reservoir layer using a $4\ mm$ puncher. Finally, each chip was die-cut
410 with a $26\ mm$ puncher. The two-layer devices were then placed in a 6-well plate (VWR, USA) for
411 hydrophilization by plasma treatment for $10\ min$ in a plasma cleaner/sterilizer (Harrick Scientific
412 Products, USA). They were baked at $80\ °C$ for $2\ h$ to increase the bonding strength.

413 *c) Fabrication and assembly of PDMS chips*

414 Polydimethylsiloxane (PDMS) chips and reservoirs were respectively molded with 1:10 Sylgard 184
415 from the SU-8 mold and a basic silicon wafer, and baked at $80\ °C$ for $2\ hours$. The patterned layer and

416 the reservoir layer were punched with the same punchers used for the SBC chips and both were plasma
417 treated for $10\,min$ prior to assembly. 6-well plates were pre-treated with $1\,\%$ (3-
418 aminopropyl)triethoxysilane (APTES, Sigma-Aldrich, Germany) in milliQ water. Finally, the devices were
419 treated with plasma again for $10\,min$ and were bonded in the 6-well plates.

420 *d) Sterilization and coating of the chip*

421 The same protocol was used on the SBC and PDMS chips. The plates containing the devices were placed
422 in a UV/Ozone ProCleaner™ Plus (BioForce Nanosciences, USA) for $10\,min$ for sterilization. The chips
423 were then rinsed with EtOH $70\,\%$ and three times with Dulbecco's phosphate buffered saline 1X (D-
424 PBS 1X, Gibco, USA) before being incubated with $5\,\%$ Poly-L-lysine (Sigma-Aldrich, Germany) in a
425 solution of boric acid (Sigma-Aldrich, Germany) and sodium hydroxide for $30\,min.$ at 37°C. After
426 washes with Neurobasal medium (Gibco, USA), they were coated with a solution of $130\,\mu g.ml-1$ rat
427 tail collagen I (Clinisciences, France) in $17.5mM$ acetic acid for 1 h at room temperature. Finally, the
428 chips were rinsed again and placed at $37\,°C$ and $10\,\%$ CO_2 overnight before cell seeding. The same
429 coating procedure was followed for the 4-well and 96-well plates, which were used as controls.

430 *e) Detachment and cleaning of the chip*

431 The wells were filled with $1\,mL$ of D-PBS to immerge the chips before to progressively detach them
432 with a tweezer. After $10\,min$ of incubation at room temperature, the chips were carefully removed
433 from the wells. If cells observation was not needed, the devices were detached directly without the
434 incubation step. To clean them, they were first bathed in a solution of milliQ water and soap solution
435 (disinfectant Dermanios Scrub CG, Laboratoire Anios, France) and gently brushed with a toothbrush,
436 then rinsed successively with milliQ water, EtOH $70\,\%$ and with milliQ water again. Finally, they were
437 either dried in a laminar flow hood for $1\,h$ before storage or dried with compressed air and directly
438 reused following the usual protocol of sterilization and coating. Optical profilometry was performed
439 with a WykoNT9100 optical profilometer (Veeco, USA) on a new chip and a chip that had been used
440 up to 4 times and cleaned each time.
441

442 **4.2 Dorsal Root Ganglia (DRG) Cell Culture**

443 *a) Cells harvesting, seeding and maintenance*

444 Dorsal root ganglia (DRGs) were harvested from embryos at $E13.5$ surgically removed from pregnant
445 C57BL/6 J mice (Janvier, France) following a previously described protocol.[50] For explant culture, the
446 devices were filled with DRG plating media (DMEM high glucose (Gibco, USA), Glutamax 1X (Gibco,
447 USA), 10% Heat Inactivated Horse Serum (Gibco, USA), 2.5S NGF (R&D Systems, USA),
448 Antibiotic/Antimycotic 1X (Gibco, USA)). One DRG was placed in the central chamber of each

449 microfluidic chip or in the center of a well of a 4-well plate. For dissociated cell culture, the DRGs were
450 trypsinized with a solution of 0.25 Trypsin in Hanks' Balanced Salt Solution 1X (HBSS 1X, Gibco, USA)
451 for = at $37\ °C$ and $5\ \%$ CO_2. Trypsinization was stopped with L-15 media (Gibco, USA) containing 10%
452 horse serum. DRGs were then spun down at $1{,}500\ rpm$ for $5\ min$. The cells were resuspended in DRG
453 plating medium and seeded at a targeted density of $275\ 000\ cells.cm^{-2}$ inside the central chambers
454 or in a well from a 96-well plate, as a control. The dissociated cells were seeded in a volume of $7\ \mu L$ in
455 the central chamber and incubated $2\ h$ prior to fill the reservoir, to allow them to adhere to the surface
456 of the culture plate. DRGs were incubated at $37\ °C$ and $10\ \%$ CO_2. 18 to $20\ h$ after the seeding, the
457 medium was replaced with Neurobasal medium supplemented with Glutamax 1X, 2.5S NGF, B27
458 Supplement (Gibco, USA) and Antibiotic/Antimycotic 1X. The medium was changed every $2\ days$. To
459 prevent evaporation, $3\ mL$ of D-PBS was added in between the wells of the plates. All aspects of animal
460 care and animal experimentation were performed in accordance with the relevant guidelines and
461 regulations of INSERM and Université Paris Cité (authorization APAFIS#7405-2016092216181520).

462 *b) Live and dead cell assays*

463 Viability was assessed at day *in vitro* $DIV3$ and $DIV7$ using a LIVE/DEAD Viability/Cytotoxicity Assay Kit
464 (Invitrogen, USA). Cells were incubated for $30\ min$ at $37\ °C$ and $10\ \%$ CO_2 and rinsed with L15 medium
465 without red phenol (Gibco, USA). Cells were observed using the Cy5 and EGFP channels on a ZEISS
466 Axio Observer Z1 microscope (Zeiss, Germany) incubated at $37\ °C$ and $10\ \%$ CO_2 and the images were
467 processed using ImageJ 1.54j.[51] The data obtained were analyzed using R and tidyverse and the graphs
468 were plotted on GraphPad Prism 8 (GraphPad Software, USA). The full R markdown script is available
469 as supplementary information (*Script SI.1*). Sidak's multiple comparison tests were performed. The
470 alpha value threshold was set at $5\ \%$ and the *P*-values are presented as follows: $*\ P < 0.05$, $**\ P <$
471 0.01, $***\ P < 0.001$.

472 *c) Immunostaining*

473 Cell cultures at $DIV3$ and $DIV7$ were fixed with 4% paraformaldehyde (PFA, Electron Microscopy
474 Sciences, USA) for $10\ min$ at room temperature. At $DIV7$, the SBC chips were detached to facilitate
475 the staining. The cells were then permeabilized ($0.25\ \%$ Triton X100 (Sigma-Aldrich, Germany) and
476 $0.1\ \%$ Tween20 (Thermo Fischer Scientific, USA) in D-PBS) for $10\ min$. After washes with D-PBS and
477 incubation for $45\ min$ in a blocking solution ($0.5\ \%$ Triton X100, $0.1\ \%$ Tween20, $2\ \%$ bovine serum
478 albumin (BSA, Sigma-Aldrich, Germany), $3\ \%$ Mouse on Mouse blocking reagent (Bio-Techne, USA) and
479 $5\ \%$ Normal Donkey Serum (Sigma-Aldrich, Germany)), the cells were incubated overnight at $4\ °C$ with
480 the primary antibodies: rabbit anti-NFH ($1{:}\ 1000$, Merck Millipore Ref. AB1989, Germany) and mouse
481 anti-SMI312 ($1{:}\ 250$, BioLegend Ref. AB_2566782, USA). The cultures were again rinsed with D-PBS
482 and incubated for 1h at room temperature with the secondary antibodies: donkey anti-rabbit Alexa

483 Fluor™ 488 ($1:1000$, Invitrogen Ref. A-21206, USA) and donkey anti-mouse Cyanine3 ($1:165$, Jackson
484 Ref. AB_2340813, USA). For nuclear staining, the devices were incubated $30\,min$ in a solution of
485 NucRed (Invitrogen, USA) in HBSS. To reduce the amount of antibody and staining required for chip
486 detachment, rings were 3D printed in polylactic acid and placed in each well to reduce the surface area
487 (**Figure SI.1-2**). Images were captured on a ZEISS Axio Observer Z1 microscope and processed on
488 ImageJ using the *NeuriteJ* plug-in. The resulting data were processed in RStudio (Posit, USA) and graphs
489 plotted on GraphPad Prism 8. The full R markdown script is available as supplementary information
490 (**Script SI.2**) Sidak's multiple comparison tests were performed.

491 *d) Western Blot*

492 Western blots (WB) were performed on DRGs explant cultures at $DIV11$. The chips were detached,
493 and the proteins were extracted by scraping the DRGs in RIPA buffer ($50\,mM$ Tris Base, $150\,mM$ NaCl,
494 $1\,mM$ $MgCl_2$, $1\,\%$ NP-40 (Sigma-Aldrich, Germany), $0.1\,\%$ sodium dodecyl sulfate SDS (EU0660,
495 Euromedex, France), $0.5\,\%$ sodium deoxycholate (D6750, Sigma-Aldrich, Germany), Protease Inhibitor
496 Complete mini EDTA free (11836170001, Roche, Switzerland), Pyrophosphate ($10\,mM$, S9515, Sigma-
497 Aldrich, Germany), Phosphate Inhibitor Cocktail 2 ($10\,\mu L/ml$, P5726, Sigma-Aldrich, Germany),
498 Phosphate Inhibitor Cocktail 3 ($10\,\mu L/ml$, P0044, Sigma-Aldrich, Germany) and sodium fluoride
499 ($10\,mM$, S7920, Sigma-Aldrich, Germany)). The protein quantification was determined using a Bio-Rad
500 DC Protein Assay (Bio-Rad, USA) with BSA as the standard. A total of $15\,\mu g$ of protein was loaded into
501 a $4-20\,\%$ TGX 10-wells gel (Bio-Rad, USA) and the transfer step was performed using a Trans-Blot
502 Turbo transfer system (Bio-Rad, USA) on polyvinylidene difluoride membranes (Bio-Rad, USA). Non-
503 specific binding sites were blocked with blocking solution ($5\,\%$ BSA in Tris Buffered Saline (TBS,
504 Euromedex, France) and then incubated with primary antibodies rabbit anti-Sox10 ($1:1000$,
505 ab155279, Abcam, UK) and rabbit anti-Actin ($1:2\,000$, ab8227, Abcam, UK), followed by a goat anti-
506 rabbit Horseradish Peroxidase (HRP) coupled secondary antibody ($1:20\,000$, 111-035-144, Jackson IR,
507 UK). Finally, specific bands were detected with the Pierce®ECL 2 Western Blotting substrate kit
508 (RPN2209, Thermo Fischer Scientific, USA), using the ImageQuant LAS 4000 ECL imaging system (GE
509 Healthcare, USA). The images were analyzed using the gel tool analyzer of ImageJ (1.54g). The
510 integrated densities of Sox10 were normalized to β-actin expression and plotted using GraphPad Prism
511 10. A t-test was performed for statistical analysis.

512 *e) RT-qPCR*

513 RT-qPCR was performed on explants at $DIV7$. The chips were detached and the DRGs were scraped in
514 Trizol (15596018, Invitrogen, USA). The RNA was precipitated and resuspended in water prior to
515 quantification with a Nanodrop (Thermo Fischer Scientific, USA). The SuperScript III Reverse
516 Transcriptase (18080085, Invitrogen, USA) was used following the manufacturer's instructions to

517 realize the reverse transcriptase of $90\ ng$ of RNA. qPCR was performed on a CFX384 biorad qPCR
518 machine ($15\ s$ of denaturation at $95\ °C$, $30\ s$ of elongation at $60\ °C$ – 40 times) by using Sybr Green
519 Rox kit (AB-1162/B, Thermo Fischer Scientific, USA) containing $300\ nM$ of selected primers (**Table 1**).
520 The quality of the qPCR was tested by checking the dissociation curve. Each qPCR reaction was
521 performed in triplicates. Relative expression of target genes was normalized to Ppia and Rpl13a mRNA
522 levels and calculated using the ΔΔCt. Results are expressed as fold changes with respect to the controls
523 and the data were plotted using GraphPad Prism 10.
524

Gene	Sequence 3'-> 5'	Sequence 5'->3'
Ppia	GGCAAATGCTGGACCAAAC	CATTCCTGGACCCAAAACG
Rpl13a	TACGCTGTGAAGGCATCAAC	TTGGTATTCATCCGCTTCCG
Krox20	TGCACCTAGAAACCAGACCTTC	TGCCCGCACTCACAATATTG
Mbp	ACTCACACACGAGAACTACCC	GGTGTTCGAGGTGTCACAATG
Dhh	ATCCACGTATCGGTCAAAGC	AGTCACCACGATGTAGTTCCC
B-actin	ATGGTGGGAATGGGTCAGAAG	TGCCATGTTCAATGGGGTAC
Pgk1	ACGGTGTTGCCAAAATGTCG	TTGGAACAGCAGCCTTGATC
Gapdh	GTGGACCTCATGGCCTACAT	TGTGAGGGAGATGCTCAGTG

525 **Table 1:** qPCR primers used in this study
526
527 *f) Scanning Electron Microscopy*
528 After $21\ days$ of culture, the cells were rinsed with D-PBS and fixed in $4\ \%$ PFA. The bottom of each
529 well was then laser-cut using a Universal Laser Systems VLS 4.60 (Universal Laser Systems, Inc., USA)
530 to obtain circular slides suitable for scanning electron microscope. The chips were detached according
531 to the previous protocol and the biological samples were dried using an automated critical point dryer
532 Leica EM CPD300 (Leica Microsystems, France). The samples were then metallized with platinum in a
533 Cressington 208HR metallizer (Cressington Scientific Instruments, UK) and observed in a Zeiss Gemini
534 SEM 360 scanning electron microscope (Zeiss, Germany).

535 *g) Finite Element Method (FEM)*
536 COMSOL Multiphysics software was used to perform diffusion-convection simulations on the
537 microfluidic device. The *creeping flow* and *transport of diluted species* modules were implemented for
538 this study. In the diffusion-convection experiment, a pressure gradient of 5 and $20\ Pa$ between the
539 inlet and the outlet was used for the convective flow, and an inflow of 0.5 mM of water (diffusion
540 coefficient $D = 10^{-9}\ m^2.s^{-1}$) was used for diffusion, by keeping the inlet and outlet the same

541 (transport) or opposite (isolation). The system comprises two identical blocks of size $1 \times 1 \times$
542 $0.1\ mm^3$, merged with a smaller block of size $0.5 \times 1 \times 0.028\ mm^3$. This geometry represents the
543 junction between the central and peripheral chambers, which is of particular relevance for the study.
544 Approximate solutions of the following equations were numerically calculated by the FEM software:

545 $\rho \frac{\partial u}{\partial t} = \nabla \cdot [-pI + \mu(\nabla u + (\nabla u)^T)] + F$ (1)

546 $\rho \nabla \cdot u = 0$ (2)

547 $\frac{\partial c_i}{\partial t} + \nabla \cdot J_i + u \cdot \nabla c_i = R_i$ (3)

548 $J_i = -D_i \nabla c_i$ (4)

549 Equation (1) is the Navier-Stokes equation, (2) is the incompressibility of the flow, (3) is the diffusion-
550 convection equation, and equation (4) is Fick's law of diffusion with ρ the fluid density, u the fluid
551 velocity, p the fluid pressure, I the identity matrix, T the temperature, μ the fluid dynamic viscosity, F
552 the external forces applied to the fluid, and where, for species i, c_i is the concentration, J_i is the molar
553 flux, R_i is a net volumetric source for *c* and D_i is the diffusion coefficient.

554 *h) Particle image velocimetry*

555 The devices were first coated with 5% BSA in D-PBS1X for 10 min to reduce collision impact with the
556 walls. FluoSpheres™ amine ($0.2\ \mu m$ diameter) (Invitrogen, USA) were diluted $1/40000$ in the same
557 solution. To obtain a flow of $25\ Pa$, $60\ \mu L$ of the microbead solution was injected into the central
558 chamber and $5\ \mu L$ of the solvent solution was injected into the peripheral chambers. D-PBS was added
559 in between the wells to prevent evaporation and the plate was sealed with parafilm. Microbeads were
560 observed using the Cy5 channel on a ZEISS Axio Observer Z1 microscope. Images were analyzed directly
561 using the Zeiss Zen software and data were plotted using GraphPad Prism 8.

562 *i) Time, cost, carbon footprint and energy consumption calculations*

563 The estimated times, costs, carbon footprint and energy consumption were calculated for one chip.
564 Power consumption was measured using a wattmeter Primera-Line PM 231 E (Lextronic, France). The
565 measurements were converted to energy by multiplying the power by the time of use of each piece of
566 equipment. Whenever an equipment was used for several devices at the same time, the amount of
567 energy and time were divided by the number of chips to estimate the amount required for one device.
568 Material prices were based on supplier prices. Electricity prices were based on the French electricity
569 price in June 2024 ($€0.2516/kWh$). The carbon footprint was calculated based on the direct emissions
570 of products manufactured in Europe, using the Ecoinvent 3.5 database, required for the fabrication of
571 one chip. PDMS production was assimilated to silicone using the Ecoinvent process 'product

production | silicone product | APOS, S RER', while SBC manufacture was modeled as an average of 25 % polyethylene, 25 % polybutadiene and 50 % polystyrene using the processes 'polyethylene production, high density, granulate | polyethylene, high density, granulate | APOS, S, RER', 'polybutadiene production | polybutadiene | APOS, S, RER' and 'polystyrene production, general purpose | polystyrene, general purpose | APOS, S, RER'. In addition, the impact calculations included electricity consumption, 6-well plate production, washing protocol and waste disposal by incineration. Impacts were calculated using the ReCiPe 2016 Midpoint (H) method.

Data availability

The data for this article plots can be found on https://github.com/HugoSalmon/25Moreau.

Supporting Information

Supporting Information is available from the Wiley Online Library.

Author Contributions

H.S., S.B., S.M., A.S. and A.E-T conceptualized the study. H.S., A.S. and C.D-B. provided the resources. S.M., H.S, A.S. and A.E-T made data curation. S.M., A.S. and A.E-T realized the formal analysis. H.S. and S.B. supervised the study. H.S. acquired the fundings. H.S., T.EJ., A.S. and S.B. validated the study. Investigation was conducted by H.S., S.M., R.F-B., A.S. and C.D-B.. S.M. wrote the original draft and H.S., T.EJ., A.S., C.D-B. and S.B. review and edited the paper. H.S., S.B., S.M., T.EJ., R.F-B., A.S. and C.D-B. designed the methodology of the study. H.S. and S.B. handled the project administration.

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Conflict of Interest

The authors declare no conflict of interest.

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List of abbreviations

4R: Reduction, Replacement, Refinement and Responsibility

CNS: Central Nervous System

DIV: Days *in vitro*

DRG: Dorsal Root Ganglia

PDMS: PolyDiMethylSiloxane

PNS: Peripheral Nervous System

RT-qPCR: Reverse Transcription quantitative Polymerase Chain Reaction

SBC: Styrenic Block Copolymer

SEM: Scanning Electron Microscope

sTPE: soft Thermoplastic Elastomer

WB: Western Blot

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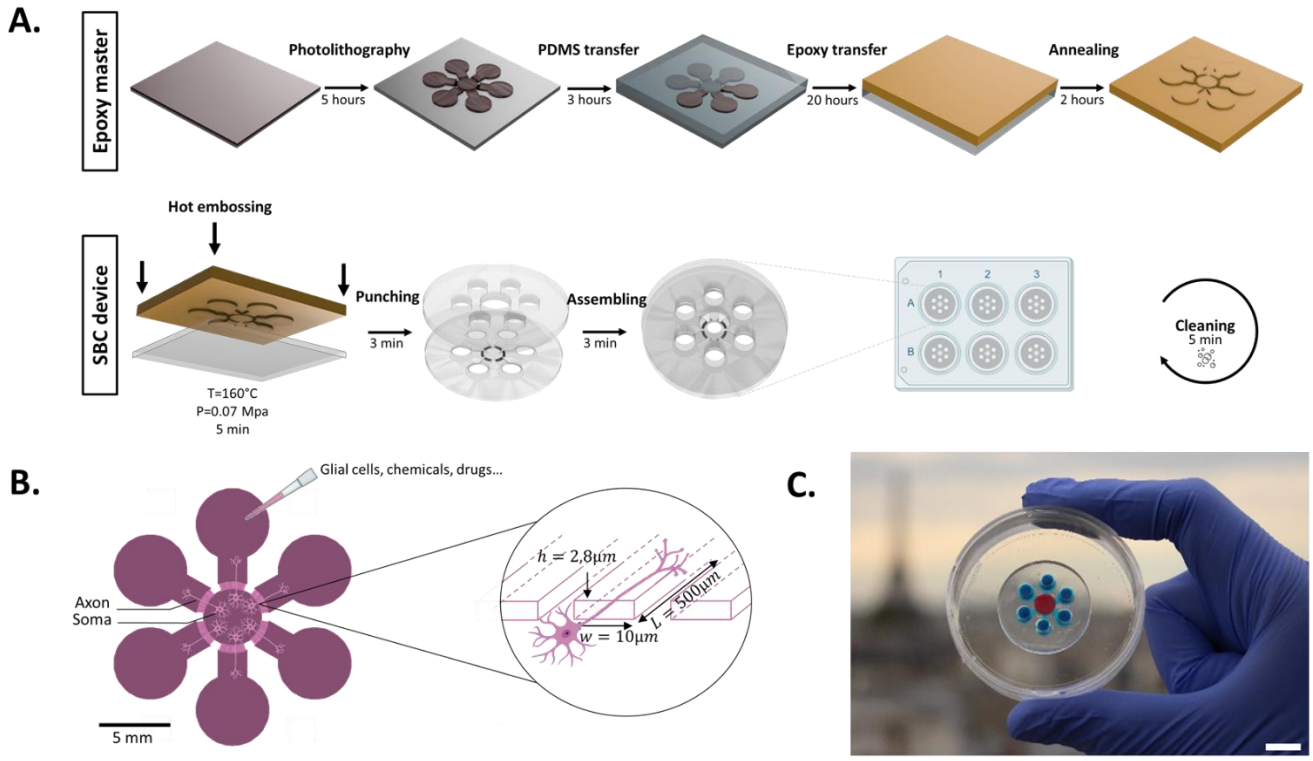


Figure 1: Fabrication and design of the thermoplastic chip. A. Fabrication process of the two-layer microfluidic chip in styrenic bloc copolymers (SBC). **B.** Scheme of the design of the chip. Somas are compartmentalized in the central chamber, while axons grow through the 292 microchannels. **C.** Image of the device filled with food dyes. Scale bar = 1 cm.

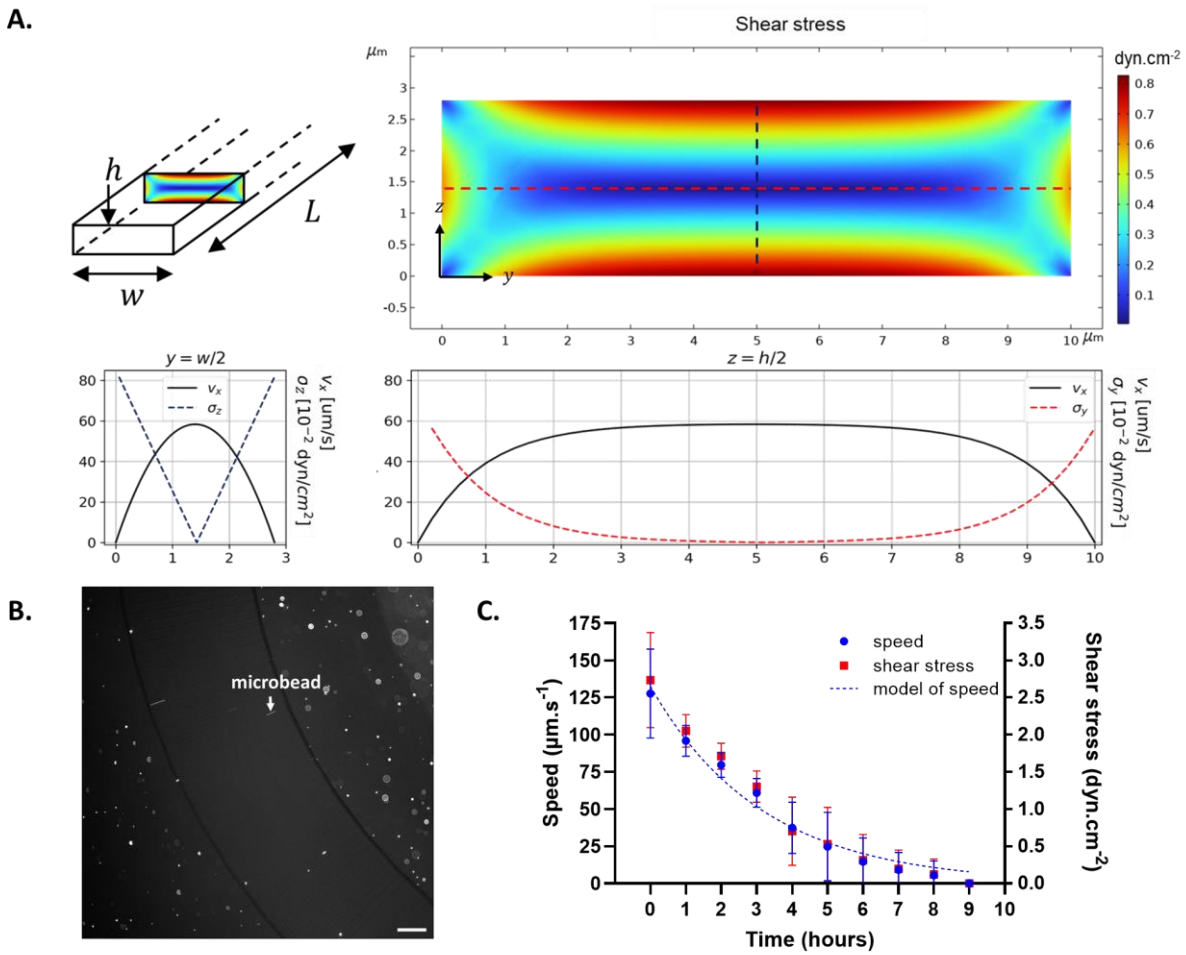


Figure 2: Estimation of the shear stress inside microchannels. A. Shear stress field induced by a 30Pa flow over a microchannel cross-section calculated by Finite Element Methods (FEM), with associated analytical plots of the velocity (in black) and shear stress along the y-axis (dashed red line) and along the z-axis (dashed blue line). **B.** Particle image velocimetry in the styrenic bloc copolymers (SBC) chip of 200 nm fluorescent polystyrene particles. Scale bar $= 100\ \mu m$ **C.** Evolution of the flow average speed inside the microchannels (blue curve), fitted exponential model for the average flow speed (dotted blue line), and associated shear stress (red curve) over time for initial pressure conditions of $25\ Pa$.

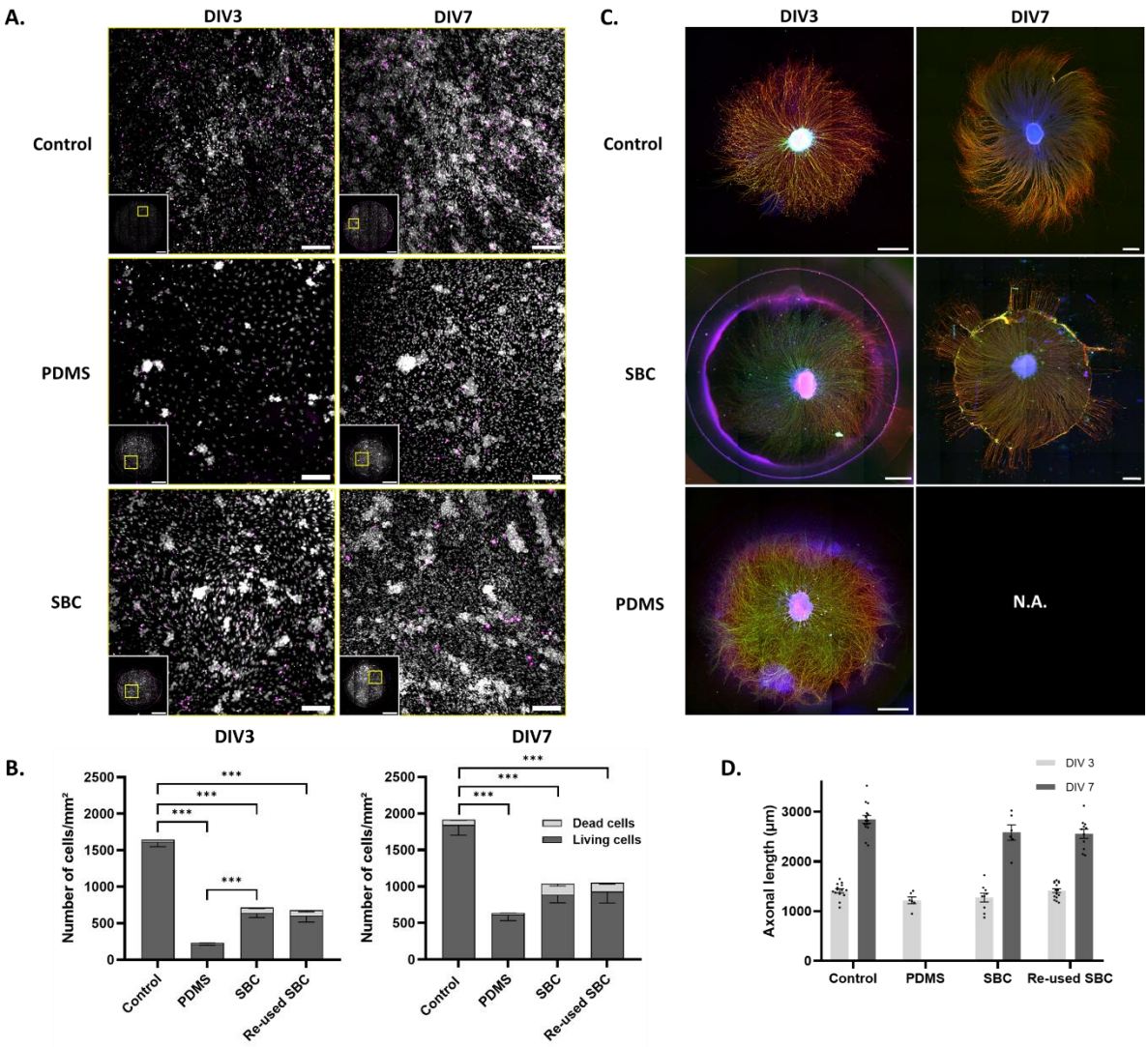


Figure 3: Comparison of viability and axonal growth of Dorsal Root Ganglia (DRG) explants within the chips A. Viability test for a control well from a 96-well plate and a central chamber of a Styrenic Block Copolymer (SBC) and a PolyDiMethylSiloxane (PDMS) chip at $DIV3$ and $DIV7$. The images show all cells (white) and dead cells (magenta). Miniature tiles (bottom left of each image), scale bars $= 1.2\,mm$, and its representative frame corresponding to the region in the yellow square, scale bars $= 150\,\mu m$. **B.** The number of living and dead cells per mm² inside the control wells, PDMS chips, SBC chips and reused SBC chips at $DIV3$ and $DIV7$. DRG $n = 6 - 24$, embryos $e = 6 - 12$. Results are represented as the mean ± SEM. Two-way ANOVA were performed. The alpha value threshold was set at 5% and the *P*-values are represented as follows: $* \, p < 0.05$, $** \, p < 0.01$, $*** \, p < 0.001$. **C.** Tiles of DRG explants inside the controls wells, PDMS chips, SBC chips and reused SBC chips at $DIV3$ and $DIV7$. Nuclei (blue), neurofilaments H (green), SMI312 (red), scale bars $= 500\,\mu m$ **D.** Axonal length inside the control wells, PDMS chips, SBC chips and reused SBC chips at $DIV3$ and $DIV7$. No data is available for PDMS chips at DIV7, as it was not possible to measure the axonal length. $n = 6 - 15$, $e = 6 - 12$. Results are represented as the mean ± SEM.

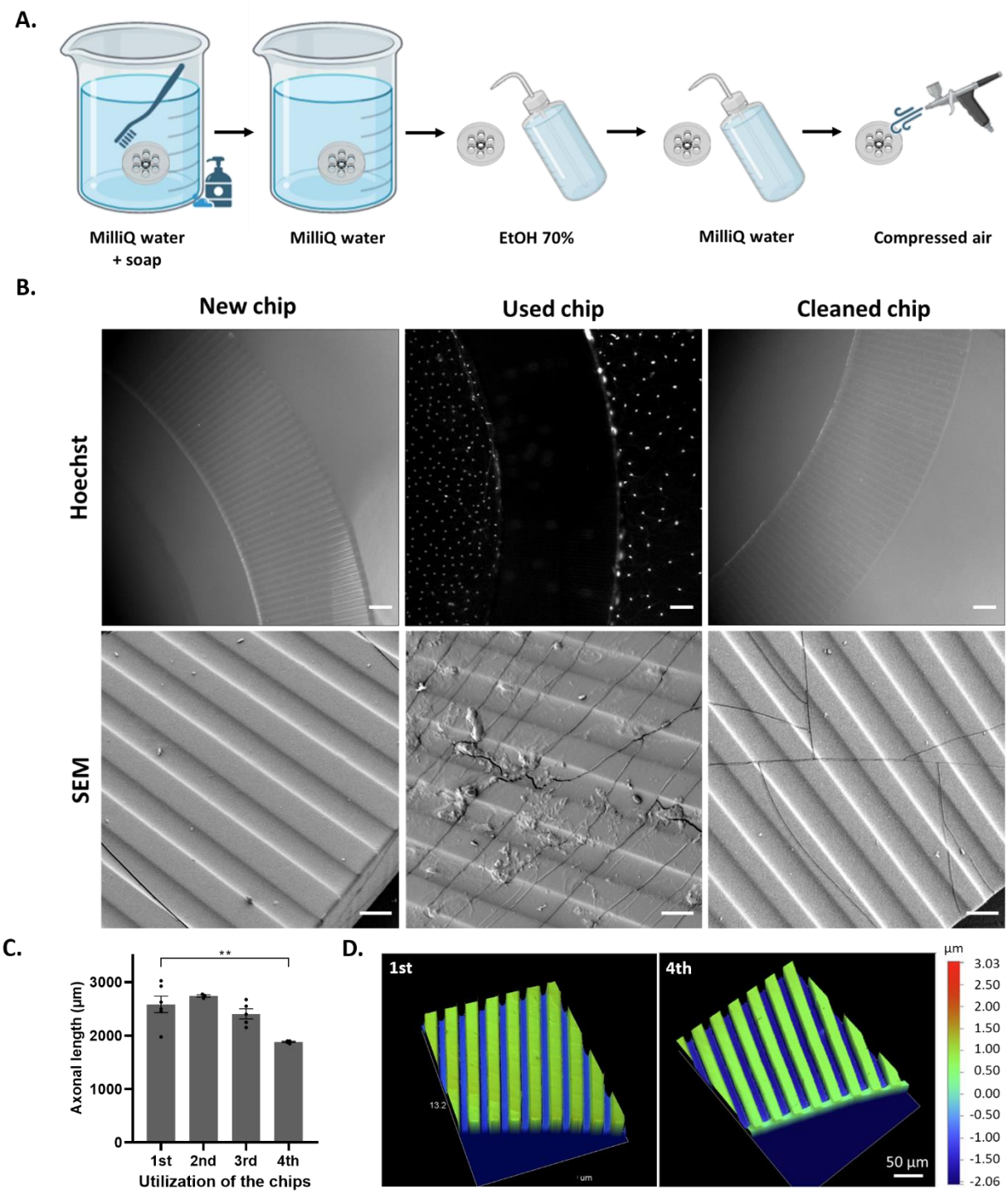


Figure 4: Reusability of Styrenic Block Copolymer (SBC) chips A. Cleaning process of the two-layer microfluidic chip in SBC. **B.** Hoechst staining (scale bar $= 100\ \mu m$) and SEM (scale bar $= 10\ \mu m$) imaging on a new, a used and a cleaned SBC chip. **C.** Axonal length of Dorsal Root Ganglia (DRGs) at DIV7 depending on the number of uses of the SBC chip. DRG $n = 3 - 6$, embryos $e = 3 - 6$. The results are presented as the mean±SEM. A Tukey's multiple comparison test was conducted. The alpha value threshold was set at $5\ \%$, and the *P*-values are represented as follows: $*\ p < 0.05, **\ p < 0.01, ***\ p < 0.001$. **D.** Optical profilometry of the microchannels of a new SBC chip and a SBC chip used four times.

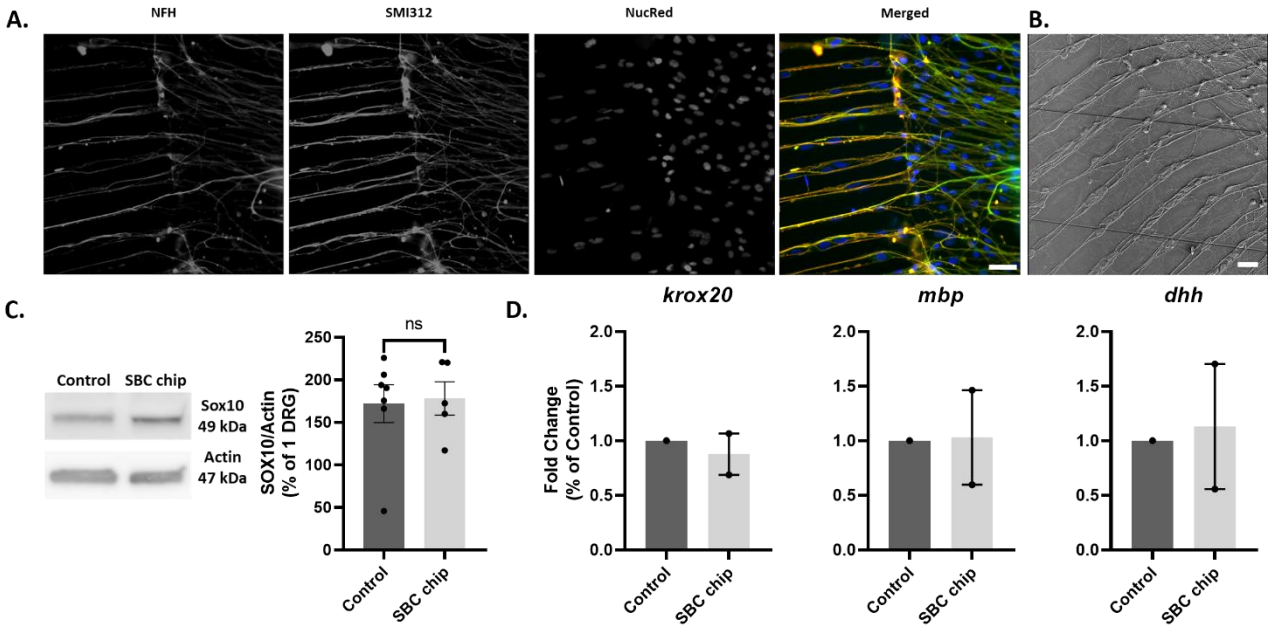
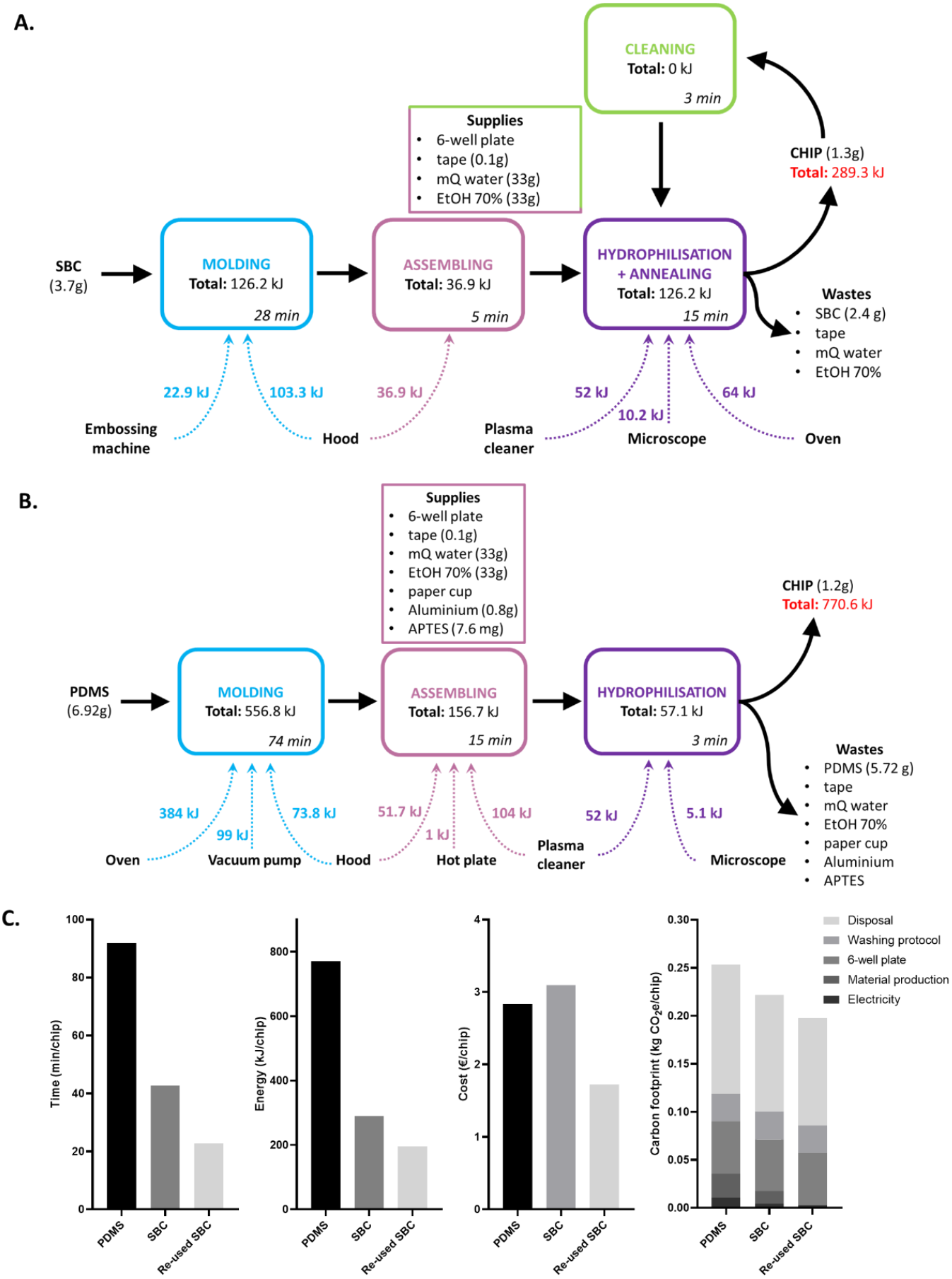


Figure 5: Direct accessibility to cells A. Immunostaining of the neurofilaments of a Dorsal Root Ganglia **(**DRG) at $DIV7$ after opening of the chip. Nuclei (blue), neurofilaments H (green), SMI312 (red). **B.** SEM image of the neurofilaments of a DRG at $DIV21$. Scale bars represent $20\ \mu m$ **C.** Western blot analysis of 4 explants DRGs at DIV11 in control wells from a 96-well plate and in Styrenic Block Copolymer (SBC) opened chips and corresponding graph representing Sox10 expression. Actin was used as a loading control. Results represent the mean ± SEM of three independent experiments ($n = 5 - 8$, embryo $e = 3$). A t-test was performed for statistical analysis (ns=non-significant). **D.** RT-qPCR analysis of *Krox20*, *Mbp* and *Dhh* genes of one DRG at $DIV7$ in controls and in SBC chips. Results represent the mean ± SEM of three independent experiments ($n = 6 - 10$, embryo $e \geq 5$). A Mann-Whitney-test was performed for statistical analysis (ns=non-significant).



770 **Figure 6: Assessment of the resources used and CO2 emissions generated to produce a Styrenic Block**

Copolymer (SBC) chip and a PolyDiMethylSiloxane **(PDMS) chip.** Energy consumption of the different steps of a fabrication process for **A.** a SBC-based chip and **B.** PDMS-based chip. **C.** Comparison of the time, energy, cost and carbon footprint associated to the fabrication of a PDMS chip, a SBC chip and a reused SBC chip. Note that we did not take into account the labor cost and the equipment maintenance, as these are similar in our context for both materials.