Medial plantar nerve ligation as a novel model of neuropathic pain in mice: pharmacological and molecular characterization

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Peripheral neuropathic pain is a consequence of an injury/disease of the peripheral nerves. The mechanisms involved in its pathophysiology are not entirely understood. To better understand the mechanisms involved in the development of peripheral nerve injury-induced neuropathic pain, more experimental models are required. Here, we developed a novel peripheral neuropathic pain model in mice by using a minimally invasive surgery and medial plantar nerve ligation (MPNL). After MPNL, mechanical allodynia was established, and mice quickly recovered from the surgery without any significant motor impairment. MPNL causes an increased expression of ATF-3 in the sensory neurons. At 14 days after surgery, gabapentin was capable of reversing the mechanical allodynia, whereas anti-inflammatory drugs and opioids were ineffective. MPNL-induced neuropathic pain was mediated by glial cells activation and the production of TNF-\(\alpha\) and IL-6 in the spinal cord. These results indicate MPNL as a reasonable animal model for the study of peripheral neuropathic pain, presenting analgesic pharmacological predictivity to clinically used drugs. The results also showed molecular phenotypic changes similar to other peripheral neuropathic pain models, with the advantage of a lack of motor impairment. These features indicate that MPNL might be more appropriate for the study of neuropathic pain than classical models.

Neuropathic pain is widely recognized as one of the most difficult pain syndromes to treat and represents a significant challenge to researchers and clinicians. While it often does not respond to conventional analgesic therapies, non-conventional painkillers including anti-epileptics (such as carbamazepine and gabapentin) or antidepressants (such as amitriptyline) can be effective\(^1\). Nevertheless, patients continue to suffer with moderate severity pain despite taking prescribed medications for their condition\(^2\).

Neuropathic pain results from lesions in the central or peripheral nervous system caused by mechanical trauma, metabolic diseases, neurotoxic chemicals, infection, some medications or tumour invasion\(^3\). The identification of a specific pathophysiological change involved in the induction and maintenance of neuropathic pain may provide the basis for the development of novel analgesic therapies for this disease. In this context, the understanding of the neurobiology of neuropathic pain has been greatly accelerated by the development of animal models that reflect some elements of human pain syndromes\(^4\). Mechanical trauma in the peripheral nerves are the most used experimental models to induce neuropathic pain\(^5\). For instance, chronic constriction injury (CCI) caused by ligation of the sciatic nerve has been extensively used to elucidate the pathophysiology of peripheral neuropathic pain\(^6-10\). Importantly, these models require vast invasive surgeries, and in some cases, they result in damaged muscles or osteotomy. Consequently, animals require a long period to recover after surgery, which complicates the study of neuropathic pain in the initial period (induction phase). During this recovery phase, which lasts about three to five days, animals exhibit a considerable loss of functional gait, which makes the evaluation of pain behaviours difficult. Therefore, to better understand the mechanisms involved in the induction and maintenance of peripheral nerve injury-induced neuropathic pain, more experimental models are required.
In the present study, we propose a novel model to study peripheral neuropathic pain that is based on a minimal invasive surgery and medial plantar nerve ligation (MPNL) in mice (Fig. 1). The pharmacological predictivity and molecular mechanisms involved in MPNL-induced peripheral neuropathic pain were also addressed.

Results

MPNL induced mechanical hypersensitivity and ATF-3 expression in the dorsal root ganglia.

On the first day following the surgical procedure, the MPNL group showed a marked mechanical allodynia ipsilateral to the nerve injury, which was marginally different from sham-operated animals (Fig. 2A). However, MPNL-induced mechanical allodynia persisted, at least, twenty-five days after surgery, whereas the nociceptive threshold of sham-operated animals recovered to control levels (P < 0.001, n = 5; Fig. 2). In another set of experiments, we performed the MPNL model parallel to another well-characterized model of peripheral neuropathic pain in mice, the chronic constriction injury model (CCI). Both MPNL and CCI induced mechanical allodynia (Fig. 2B). However, the intensity of mechanical allodynia was higher in the CCI group compared with that in the MPNL group (Fig. 2B). Interestingly, in our conditions, it was only possible to assess the mechanical allodynia 7 days after the CCI surgery because some animals dragged their hind paw, which makes the mechanical threshold measurement difficult (Fig. 2B). Importantly, in the CCI model, 100% of the mice responded to the two last filaments (0.02 and 0.008 g) of the von Frey series, whereas none on the MPNL model mice reached this level of mechanical allodynia.

To evaluate whether the MPNL model would cause injury to peripheral sensory neurons, DRGs (L3–L6 pooled) were collected, and protein levels were assessed. There was a significant time-dependent increase in ATF-3 expression after MPNL, which peaked seven days post-injury, returning to control levels 21 days after MPNL (Fig. 3A). In another set of experiments, DRGs were harvested separately (L3 to L6; a pool of 3) at seven days following surgery to evaluate which DRG was more affected after MPNL by measuring ATF-3 expression. The increased expression was more prominent in L4 ganglia, although L3 and L5 also showed significantly higher ATF-3 expression compared with sham-control ganglia (Fig. 3B). Importantly, the up-regulation of ATF-3 was higher in the DRGs (L3–L6 pooled) of the CCI injured mice compared with the MPNL mice (Fig. 3C).

Figure 1. Representative image of the steps involved in the induction of MPNL. (A) Mice under anaesthesia and medial paw region, site of procedure; (B) incisions of approximately 0.5 cm; (C) MPN exposure with a glass probe; (D–G) MPN ligation (H,I) suture of the incision.
Corroborating the western blotting data, the immunofluorescence of L4 ganglia revealed an increase in ATF-3 immunoreactivity in the nuclei of sensory neurons after MPNL compared with sham-operated DRGs, which was even higher in the CCI-operated mice (Fig. 4).

The motor functional capacity of animals undergoing MPNL surgery was evaluated using rota-rod test. No significant difference was observed between the sham and MPNL groups on the third (remaining time means of 117.6 and 119 min, respectively), seventh (means of 120 min in both groups) and fourteenth (means of 120 min in both groups) days after surgery. Corroborating the hypothesis that MPNL does not cause significant motor impairment, we found no cells expressing ATF-3 in the ventral horn of the spinal cord of the MPNL mice (Fig. 5). On the other hand, there was an increase in the ATF3-expressing cells in the ventral horn of the spinal cord of CCI mice, suggesting some degree of damage in the motor neurons (Fig. 5).

**Analgesic pharmacological predictive value of the MPNL model.** The pharmacological predictive value of MPNL was determined by treating mice on the third and fourteenth days following MPNL with different
classes of clinically used analgesic drugs, including steroids (dexamethasone; Fig. 6A,B), NSAIDs (indomethacin, Fig. 6C,D), opioids (morphine; Fig. 6E,F) and gabapentinoids (gabapentin, Fig. 6G,H). On the third day following MPNL, all drugs were significantly reversed mechanical allodynia (Fig. 6A,C,E,G). On the other hand, on the fourteenth day after surgery, dexamethasone, indomethacin and morphine produced only a slight reduction in mechanical allodynia after MPNL (Fig. 6B,D and F). Nevertheless, gabapentin reversed MPNL-induced mechanical allodynia by the 14th day after surgery (Fig. 6H).

Glial cell activation in the spinal cord contributed to the MPNL-induced pain-like behaviours. Glial cell activation in the spinal cord (mainly microglia and astrocytes) occurs after peripheral nerve injury and accounts for chronic pain induction and maintenance11–14. In this context, the participation of spinal glial cells in MPNL-induced pain-like behaviours was investigated. Firstly, microglial cell activation/proliferation in the spinal cord was assessed after MPNL induction through the analysis of Iba1 expression. There was a significant increase in the expression of Aif1 (Iba1 gene) mRNA from the third to the tenth day after surgery (Fig. 7A). There was also a significant increase in Iba1 protein expression three days after MPNL (P < 0.01) (Fig. 7B). These results were confirmed by immunofluorescence of spinal cord slices. The naive group showed the constitutive expression of Iba1 in the spinal cord (Fig. 7C), whereas a massive increase in Iba1-expressing cells was observed on the ipsilateral side (dorsal horn) of the spinal cord 3 days after MPNL (Fig. 7D). In only 10% of the slices analysed, there was an increase in the expression of Iba1 in the ventral horn of the spinal cord (data not shown).

Further supporting the hypothesis that MPNL triggers microglia activation in the spinal cord, we found that the morphology of microglia changed from a resting state (ramified) to an activated state (hypertrophied and amoeboid) after MPNL (Fig. 7C–F). Functionally, the inhibition of spinal microglia activity with intrathecal (i.t.)
injections of minocycline (preoperatively, and once a day for up to 3 days following MPNL), significantly inhibited MPNL-induced mechanical allodynia in a dose- and time-dependent manner (Fig. 7G). The higher dose of minocycline did not change the mechanical threshold of sham-operated mice (Fig. 7G). Remarkably, when minocycline treatment was interrupted, the MPNL-induced mechanical allodynia was restored (Fig. 7G).

Regarding astrocytes, there was an increase in the expression of GFAP (astrocyte activation marker) in the spinal cord 7 days after MPNL (Fig. 8A). Because the increased expression of GFAP protein was significant on day 7 after surgery, this time point was used to assess the participation of astrocytes in MPNL-induced mechanical allodynia. Then, (i.t.) treatment with fluorocitrate (an astrocyte function inhibitor) reduced MPNL-induced mechanical allodynia in a dose- and time-dependent manner (Fig. 8B). Interestingly, the higher dose (3 nmol/site) of fluorocitrate reduced the mechanical allodynia until 24 h after MPNL, but the allodynia returned to the control levels 48 h after treatment (Fig. 8B). The higher dose of fluorocitrate did not change the mechanical threshold of sham-operated mice (Fig. 8B).

**Role of spinal pro-inflammatory cytokines in the genesis of MPNL-induced pain-like behaviours.** In the last part of this study, the involvement of pro-inflammatory cytokines (TNF and IL-6) in MPNL-induced mechanical allodynia was investigated. These two cytokines were chosen because they have been implicated in the genesis of neuropathic pain in other models3,15. Firstly, the mRNA expression of these cytokines was evaluated in the spinal cord following MPNL based on real-time PCR. There was a time-dependent increase in Tnf and Il-6 gene expression after surgery (Fig. 9A,B). Whereas Tnf expression peaked 1 and 3 days after MPNL and returned to control levels 5 days after surgery, Il-6 expression was increased from day 3 until day 7, decreasing thereafter (Fig. 9A,B). Supporting these data, TNF and IL-6 protein expression was also up regulated in the spinal cord of MPNL mice at 3 days after surgery (Fig. 9C,D). To investigate whether these cytokines are involved in the genesis of MPNL-induced pain-like behaviours, MPNL was performed in C57BL/6 (WT), TNFR1/R2−/− and IL-6−/− mice. Mechanical allodynia triggered by MPNL was completely abrogated in TNFR1/R2−/− mice (Fig. 9E). As further support, the i.t. injection of infliximab (anti-TNF antibody) also reduced MPNL-induced mechanical allodynia (Fig. 9F). In addition, mechanical allodynia induced by MPNL was only slightly reduced in IL-6−/− compared with WT mice (Fig. 9G).

**Discussion**

Neuropathic pain is a multidimensional chronic pain condition that is frequently caused by peripheral or central nerve injury or disease. These damaged nerves continuously send incorrect signals to the spinal cord and
Figure 6. Pharmacological predictivity of pain-like behaviours after MPNL. At 3 or 14 days after MPNL, mice were treated systemically with (A,B) dexamethasone (2 mg/kg, s.c), (C,D) indomethacin (5 mg/kg, i.p), (E,F) morphine (5 mg/kg, i.p) or (G,H) gabapentin (100 mg/kg). The withdrawal threshold was measured using von Frey filaments after treatment. The data are presented as the mean ± S.E.M. (n = 5 per group). Time 0 represents the baseline before pharmacological treatments. ***P < 0.001 and **P < 0.01 to *P < 0.05 indicates a statistically significant difference when compared with the vehicle treated group.
Figure 7. **MPNL-induced pain-like behaviours depend on microglial cell activation.** The lumbar spinal cord samples were collected at indicated times after MPNL. (A) The mRNA for Aif1 (Iba1) and protein expressions were measured by RT-PCR (n = 6 per group), (B) western blotting (representative blotting and analyses of n = 4 per group) or (C) immunofluorescence (representative image in different magnifications, n = 4 per group). Scale bars of 500 and 100 μm. (D–F). The morphological subtypes of microglial cells (D) ramified, (E) hypertrophied and (F) amoeboid were quantified. (D) Mice (n = 5 per group) were treated with minocycline (i.t., 3–30 nmol/site) or vehicle before and 3 days after MPNL. The withdrawal threshold was assessed daily after surgery (3 h after drug administration). Time 0 represents the baseline value before surgery. The data are presented as the mean ± S.E.M. ***P < 0.001 and **P < 0.01 to *P < 0.05 indicates a statistically significant difference when compared with the sham or vehicle treated group.

Figure 8. **MPNL-induced pain-like behaviours depend on astrocyte activation.** (A) The lumbar spinal cord samples were collected at indicated times after MPNL or sham. The GFAP protein expression was measured using western blotting (representative blotting and analyses; n = 4 per group). (B) The mice (n = 5 per group) were treated with fluorocitrate (i.t., 1–3 nmol/site) 7 days after MPNL, and the withdrawal threshold was assessed before and up to 48 h after fluorocitrate treatment. The data are presented as the mean ± S.E.M. ***P < 0.001 and **P < 0.01 indicates a statistically significant difference when compared with the sham or vehicle treated group.
supraspinal sites, triggering changes in the structure, function and organization within the nervous system. One important cause of peripheral neuropathic pain is physical trauma to the peripheral nerves. However, the cellular and molecular mechanisms involved in peripheral neuropathic pain are not entirely understood. To better understand the mechanisms involved in peripheral neuropathic pain, in the present study, we developed a novel mouse model of this condition by using MPNL. To our knowledge, this is the most distal model of peripheral nerve injury described in mice that leads to a phenotype of pain hypersensitivity that might resemble human peripheral neuropathic pain syndromes.

One important sensory characteristic/symptom of neuropathic pain conditions is the development of mechanical hypersensitivity, which is clinically described as tactile allodynia. In this context, it was clearly observed that after MPNL, animals developed hypersensitivity to mechanical stimulation. Notably, mechanical hypersensitivity was not accompanied by any significant motor impairment even at the earlier time points (one day after surgery). This might be an advantage compared with other models of peripheral neuropathic pain in which motor deficits can influence the determination of the nociceptive threshold. This hypothesis is supported by our present findings in which the MPNL model did not damage motor neurons in the ventral horn of the spinal cord, whereas CCI did. In fact, the medial plantar nerve innervates the medial plantar region of the animal paw, which does not compromise the muscles responsible for the paw withdrawal reflex. Furthermore, the surgery performed to produce MPNL requires a tiny incision and did not involve any injury to muscles that are important for mouse gait. In addition, autotomy was not observed after MPNL. Signals of neuropathic pain without significant motor neuron damage and motor impairment has been also described in other models of peripheral nerve injury in rats. Another important characteristic of the present model of neuropathic pain is the intensity of mechanical hypersensitivity developed after MPNL. In contrast with other models of peripheral neuropathic pain, such as CCI (Fig. 2), MPNL did not induce a maximal response in the von Frey test, allowing the study of mechanisms...
The current understanding of neuropathic pain implies that the pain develops because of neuronal injury or diseases. To access whether the MPNL model induces neuronal damage, the expression of ATF3 was analysed. ATF3 is a member of the ATF/CREB family of basic leucine zipper (bZIP) transcription factors. It is not constitutively expressed. However, ATF3 expression can be induced in various tissues undergoing stress or injury including sensory neurons. Indeed, its expression in the cell body of primary sensory neurons has been used as an indirect marker of neuronal injury. In fact, after a physical injury to the peripheral nerves in several models of neuropathic pain, ATF3 is upregulated. MPNL induces an increase in ATF3 expression in the DRGs, supporting the hypothesis that MPNL might be a real model of neuropathic pain. Interestingly, it seems that the most affected neurons in the MPNL model are located in L4 DRG. In addition, it is important to mention that compared with CCI, MPNL is less severe in terms of promoting damage to the sensory neurons.

An important feature of a novel model for human diseases is the pharmacological predictivity. The MPNL model presents the predictive value to clinical drugs used for the treatment of neuropathic pain. Interestingly, MPNL-induced mechanical hypersensitivity was sensitive to classical anti-inflammatory drugs (NSAIDs) and opioids (morphine) at an earlier time point (3 days after surgery), suggesting an important inflammatory component in the induction phase. However, at later time points, only gabapentin exhibited a significant anti-nociceptive effect, which is similar to that observed when the neuropathic pain develops in human patients. In fact, NSAIDs and steroidal anti-inflammatory drugs are poorly effective or even ineffective at alleviating neuropathic pain in humans or animal models of neuropathic pain. In addition, in a recent systematic review and meta-analysis of pharmacological treatment of neuropathic pain, gabapentin was shown to be one of the first-line drugs. These results indicate that MPNL is a predictive model for drugs that could act against neuropathic pain.

Neuropathic pain is a result of neuroplasticity across the nociceptive system. Both peripheral and central sensitization are observed. In addition to intrinsic neural mechanisms, there is a growing recognition of the importance of non-neuronal cells, especially glial cells, in the genesis and maintenance of chronic pain. In particular, glial cells have been shown to be involved in the development of neuropathic pain. Indeed, their expression in the cell body of primary sensory neurons has been used as an indirect marker of neuronal injury.

Materials and Methods

Animals. The experiments were performed in C57BL/6 male mice (wild type, WT, 20–25 g) and mice deficient for the following proteins: TNF receptor type 1 and 2 (TNFR1/2−/−) and IL-6−/−. All animals were housed in the animal care facility of Ribeirão Preto Medical School, University of Sao Paulo. Animals were taken to the testing room at least 1 hour before the experiments and were used only once. Food and water were available ad libitum. The animal care and handling procedures were in accordance with the International Association for the Study of Pain guidelines for those animals used in pain research, and they were approved by the Committee for Ethics in Animal Research of the Ribeirão Preto Medical School-USP (Process no 128/2011).

Neuropathic pain models. MPNL model. Under isoflurane anaesthesia (1.5% in oxygen), the skin on the medial surface of the ankle was incised (0.5 cm) to expose the medial plantar nerve. After dissection, one ligature of this nerve was performed with 4–0 catgut suture (Ethicon, São Paulo, Brazil). The ligature stayed tightly bound to the nerve without throttling due to the feature of the wire used in the process. Then, 4–0 silk suture was used to join the two edges of the incision (Fig 1). The sham (control) group procedure involved an incision of...
CCI model. Mice were anaesthetized with isoflurane (1.5% in oxygen) followed by trichotomy in the surgical area. CCI was created as previously described\textsuperscript{19}. Briefly, the right sciatic nerve was exposed at the mid-thigh level, and four ligatures with 4–0 catgut suture (Ethicon, São Paulo, Brazil) were loosely tied around the nerve just proximal to the trifurcation. For the sham-operation controls, the mice underwent the same procedure without constriction of the nerve.

Compounds. The following compounds were used in this study: minocycline (Sigma, #M9511, St. Louis, MO, USA), fluorocitrate (Sigma, #F9634 St. Louis, MO, USA), indomethacin (Sigma, #I7378, St. Louis, MO, USA), dexamethasone (Sigma, #D1159, St. Louis, MO, USA), gabapentin (Tocris Bioscience, #0806, Bristol, United Kingdom), morphine (Cristália, São Paulo, Brazil), and infliximab (Schering-Plough, São Paulo, Brazil).

Drug Administration. Intraperitoneal (i.p.): the drugs were injected into the ventral portion of the mouse close to the abdominal midline using a 1 mL syringe with a needle length of 2.5 cm and a gauge of 0.7 cm.

Intrathecal (i.t.): under isoflurane anaesthesia (1.5% in oxygen), trichotomy was performed in the mice dorsal region. Using a 30 unit insulin syringe (BD Ultra Fine TM II), the drugs were administered (5 µL) between the L4 and L5 vertebrae. The animal’s tail reflex findings were expected to confirm the correct performance of the procedure\textsuperscript{41}.

Nociceptive Test. Animals were placed on an elevated wire grid, and the plantar surfaces of their hind paws were stimulated with a series of ascending force von Frey monofilaments. Mice were first habituated to the experimental environment (room and apparatus) for a period of at least 30 minutes. Mechanical allodynia was accessed by measuring the paw withdrawal threshold taken as the lowest force that evoked a brisk withdrawal response to one of five repetitive stimuli. Briefly, a logarithmic series of 10 calibrated monofilaments (von Frey hairs) was applied to the right hind paws to determine the stimulus intensity threshold stiffness required to elicited a paw withdrawal response. The log stiffness of the hairs was determined by log10 (milligrams) and ranged from 0.903 (8 mg or 0.008 g) to 3.0 (1000 mg or 1 g). The mechanical threshold is represented as previously described\textsuperscript{55,56}. An investigator who was blinded to the group allocation performed all of the behaviour experiments.

Rota-rod Test. The rotarod test was performed to check each animal’s motor coordination. The animals were placed on a rotating roller (20 rpm, initially). After 1 min, the roll speed was increased to a maximum of 36 rpm within 90 seconds. The residence time of each animal in the apparatus was measured by a maximum of 2 minutes. The test was performed three times for each animal with intervals of 20 minutes. The animals were acclimated one day before the experiment.

Western Blotting. The dissection and collection of spinal cord tissue from a region that included lumbar segments (L3–L6) and peripheral dorsal root ganglia (segments L3, L4, L5 and L6) ipsilateral to the nerve injury were performed. The tissue was homogenized and lysed in RIPA buffer (Sigma, #R0278, St. Louis, MO, USA) and protease inhibitor (Protease Inhibitor Cocktail Tablets - Roche Diagnostics, #04693116001, São Paulo, Brazil). The homogenate was centrifuged at 15,000 g for 10 minutes at 4 °C. An aliquot of the supernatant was separated for the determination of the protein concentration using the Coomassie colorimetric method (Bradford Reagent – Thermo Scientific, Rockford, IL, USA).

The protein samples were separated by electrophoresis on an SDS-PAGE acrylamide gel (gradient 4% to 20%) and transferred to nitrocellulose membranes. The incubation consisted of one hour at room temperature with blocking solution containing 5% blocking agent (GE Healthcare Life Science, Pittsburgh, PA, USA) dissolved in Tris-Buffered Saline - 10% Tween 20 (TBS-T), followed by incubation with primary antibodies anti-GFAP, 1:30000 (Merck Millipore, #AB5541, Darmstadt, Germany), anti-Iba1, 1:400 (Wako Chemicals, #019–19741, Osaka, Japan) and anti-ATF-3, 1:400 (Santa Cruz Biotechnology, #sc-188, CA, USA) overnight at 4 °C in TBS-T/5% bovine serum albumin (BSA). Then, the membranes were washed in TBS-T 3 times over 5 minutes and were incubated for one hour with secondary antibodies conjugated with peroxidase at 1:5000 in 5% blocking agent (GE Healthcare Life Science, #RPN2125, São Paulo, Brazil) dissolved in TBS-T. The membranes were incubated with HRP Substrate LuminalTM Fort Western (Millipore, #WBLUF0500, São Paulo, Brazil) and were then revealed, as indicated by the manufacturer. Image capture was performed with a Quemi TM-Doc XRS apparatus as directed by the manufacturer. The densitometric data were measured following normalization to the control using Scientific Imaging Systems (Image labTM 3.0 software, Biorad Laboratories, Hercules CA).

Real-time RT-PCR. After collection of tissue from the region that included the lumbar segments (L4–L6) of the spinal cord ipsilateral to the lesion, tissues were rapidly homogenized in TRIzol reagent (Invitrogen Technologies Corporation, Carlsbad, California, USA), and 0.2 mL of chloroform was added to the samples, which were shaken. The suspension was centrifuged for 13000 g at 4 °C for 15 min. The aqueous phase was transferred to a fresh tube to which 0.75 mL of isopropanol was added. After mixing, the samples were incubated for 2 hours at –20 °C. The samples were centrifuged for 15 min at 13000 g (4°C). The RNA precipitate was washed with 0.5 mL of ethanol, and the preparation was suspended in 20 µL of PCR-grade H2O. RNA concentrations were determined by optical density at a wavelength of 260 nm with the NanoVue® apparatus (GE Healthcare). One microgram of total RNA was transcribed into cDNA by reverse transcriptase enzyme action Improm Pre-II® (Promega, Madison, Wisconsin, USA). Quantitative RT-PCR in real time was performed on an ABI Prism® 7500 Sequence Detection System (Applied Biosystems) using SYBR-green System Fluorescence (Applied Biosystems,
Warrington, UK) for quantification during amplification. RT-PCR was performed with a final reaction volume of 6.25 μL; RT-PCR was initially kept at 95 °C (10 min), followed by 40 cycles of 94 °C (1 min), 56 °C (1 min) and 72 °C (2 min). The melting curve was analysed (65–95 °C) to verify that only one product was amplified. Samples with more than one peak were excluded. The results were analysed using the method of quantitative relative expression 2−ΔΔCt. The primer sequences were as follows in Table 1:

### Table 1. Primers used for real-time PCR.

| Gene name | Forward sequence | Reverse sequence |
|-----------|------------------|------------------|
| GAPDH | 5′-GGGTGTGACCCAGAAGGAAAT | 5′-CCTTCCACATGGGAAATT |
| TNF-α | 5′-AGGGATGGAGAATGCTCTCAATG | 5′-GACCTTGCTAGGAATTTGGAGA |
| IL-6 | 5′-TTTCATCCCAATTTCATTAC | 5′-CCTTCTGATCCTCGACTTAT |
| GFAP | 5′-AGGGCAAGAAACCGCCATCACTC | 5′-CTCAAGGGAGACGTGGCGAGGCT |
| Aif1 | 5′-TGGAGGAGCATTAGCCAAAG | 5′-GCTTCAGTTGGACGAGCCAG |

*Immunofluorescence staining and analyses.* The animals were deeply anesthetized with ketamine and xylazine (2:1) and perfused through the ascending aorta with PBS, followed by 4% paraformaldehyde. Spinal cord sections (60 μm, free-floating technique) and DRG sections (10 μm, one the glass technique) were obtained using a cryostat. The spinal cord sections were washed in PBS (0.01 M, pH 7.4) 3 times over 5 minutes and incubated in 5% normal goat serum and 1% BSA dissolved in PBS with 0.1% Triton X100 (PBS-T) for 1 hour. Subsequently, the sections were washed in PBS-T (0.01 M, pH 7.4) 3 times over 5 minutes and then subjected to immunofluorescence staining with overnight incubation at 4 °C with polyclonal anti-Iba1 (1:1000; Wako Chemicals, Osaka, Japan) or rabbit polyclonal anti-ATF3 (1:500; Santa Cruz Biotechnology, # sc-188, CA, USA). After incubation with the primary antibodies, the sections were washed in PBS-T 3 times over 5 minutes and incubated at room temperature for 2 hours with Alexa Fluor 488® goat anti-rabbit IgG (Molecular Probes, Molecular Probes, Carlsbad, CA, USA). The sections were washed with PBS-T as described earlier, mounted on glass slides with Fluoromount™ Aqueous Mounting Medium (Sigma, St. Louis, MO, USA), and then covered with cover slips. The DRG sections were fixed on slides and subsequently immersed in TBS and 50%, 70% and 50% ethanol solutions for 10 minutes each to increase the permeability. Then, the slides were washed in TBS 3 times over 10 minutes, and the blocking step was performed with 5% normal goat serum (NGS) in TBS for 60 minutes at room temperature. Subsequently, the slides were subjected to immunofluorescence staining with an overnight primary antibody incubation at 4 °C with rabbit polyclonal anti-ATF3 (1:500, Santa Cruz Biotechnology, # sc-188, CA, USA) antibody diluted in 5% NGS. As a control, primary antibodies were omitted from the reaction. They were washed in TBS 4 times over 5 min, incubated at room temperature for 60 min with Alexa Fluor 488® goat anti-rabbit IgG (Molecular Probes, Carlsbad, CA, USA) diluted in 5% NGS in TBS. The slides were washed in TBS 4 times over 5 minutes and further covered with a coverslip as described above. A confocal microscope (SP5, Leica, Wetzlar, Germany) was used to visualize and capture Iba1 and ATF-3 staining sections in the DRG and spinal cord under the same light conditions. For the quantitative analysis, at least five slices from the spinal cords and at least five from the DRGs from each mouse were used for the Iba1 and ATF-3 immunoreactive quantification. The percentage of ATF3-labeled neurons in the DRGs was calculated by dividing the number of ATF3+ neurons by the total number of neurons counted × 100. In the spinal cord, the number of ATF3+ neurons per slices was quantified.

*Microglia were classified as previously described*. In brief, ramified microglia were classified as having a normal pattern, with fine, mild and radially projecting processes. Hypertrophied microglia were defined as having an enlarged and dimmed cell body with denser, shorter and lesser projecting processes. Finally, amoeboid microglia were defined as densely stained and having enlarged soma with very few or no processes, which was usually accompanied with the presence of filopodia. The percentage of a specific morphological type was accessed by calculating the ratio of the total cell type morphology to the total number of microglial cells observed in the field.

*Cytokine measurement.* Mice were terminally anesthetized with ketamine and xylazine (2:1) and perfused through the ascending aorta with PBS at specific time points after MPNL induction. The collection of tissue from the region including lumbar segments (L4–L6) of the spinal cord ipsilateral to the lesion, these tissues were rapidly homogenized in 150 μL of buffer containing protease inhibitors. TNFα and IL-6 concentrations were determined by ELISA using commercial kits (DuoSet; R&D Systems, Minneapolis, MN, USA) and paired antibodies as instructed by the manufacturer.

*Data analysis and statistics.* All the results are presented as the means ± S.E.M. For the behaviour tests, we used 5 mice/group, and the results are representative of 2 independent experiments. For western blotting, immunofluorescence and ELISA assays, we used 4 mice/group, whereas for the PCR assays, we used 6 mice/group. Two-way analysis of variance (ANOVA) was used to compare groups and doses at all time points (curves) when the nociceptive responses were measured at different time points after the stimulus injection. The factors analysed were the treatments, time and time by treatment interactions. When there was a significant time by treatment interaction; one-way ANOVA followed by Bonferroni’s t test was performed for each time point. Alternatively, when the hypernociceptive responses were measured once after the stimulus injection, the differences between responses were evaluated by one-way ANOVA followed by Tukey’s t test. P < 0.05 was considered as significant.
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Author Contributions
M.B.S., J.C.A.F., F.Q.C., G.R.S. and T.M.C. designed the study, analysed the data and wrote the main manuscript text. M.B.S., R.K., G.S.B., T.A.B. and G.R.S. executed the experiments. M.B.S., R.K., G.S.B. and G.R.S. prepared all of the figures. All authors reviewed the manuscript.

Additional Information
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