Novel Potassium Channel Blocker, 4-AP-3-MeOH, Inhibits Fast Potassium Channels and Restores Axonal Conduction in Injured Guinea Pig Spinal Cord White Matter

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Sun W, Smith D, Fu Y, Cheng JX, Bryn S, Borgens R, Shi R. Novel potassium channel blocker, 4-AP-3-MeOH, inhibits fast potassium channels and restores axonal conduction in injured guinea pig spinal cord white matter. J Neurophysiol 103: 469–478, 2010. First published November 18, 2009; doi:10.1152/jn.00154.2009. We have demonstrated that 4-aminopyridine-3-methanol (4-AP-3-MeOH), a 4-aminopyridine derivative, significantly restores axonal conduction in stretched spinal cord white-matter strips and shows no preference in restoring large and small axons. This compound is 10 times more potent when compared with 4-AP and other derivatives in restoring axonal conduction. Unlike 4-AP, 4-AP-3-MeOH can restore axonal conduction without changing axonal electrophysiological properties. In addition, we also have confirmed that 4-AP-3-MeOH is indeed an effective blocker of \( I_h \) based on patch-clamp studies using guinea pig dorsal root ganglia cells. Furthermore, we have also provided the critical evidence to confirm the unmasking of potassium channels following mechanical injury. Taken together, our data further supports and implicates the role of potassium channels in conduction loss and its therapeutic value as an effective target for intervention to restore function in spinal cord trauma. Furthermore, due to its high potency and possible low side effect of impacting electrophysiological properties, 4-AP-3-MeOH is perhaps the optimal choice in reversing conduction block in spinal cord injury compared with other derivatives previously reported from this group.

INTRODUCTION

Traumatic spinal cord injury (SCI) results in severe functional deficits that are mainly due to damage in white matter. In most cases, axons are not completely severed but rather confused and/or compressed after SCI (Blight 1983a,b; Blight and DeCrescito 1986; Hayes and Kakulas 1997; Kakulas 1999). These anatomically continuous but functionally silent axons offer a realistic opportunity for functional restoration through therapeutic interventions. It is well accepted that myelin damage, a well-known pathology in SCI, could expose the fast potassium channels that directly contribute to conduction failure (Blight 1989; Blight and Gruner 1987; Shi et al. 1997; Waxman 1992, 1994). Specifically, activated potassium channels allow leakage of positive charges carried by potassium during depolarization and prevent the initiation of an action potential. Therefore blocking such potassium channels has been identified as an effective means to restore axonal conduction (Blight et al. 1991; Hansebout et al. 1993; Shi et al. 1997; Waxman 1989).

For the last two decades, researchers have identified 4-aminopyridine (4-AP), a known fast potassium channel blocker, as an effective agent in restoring axonal conduction in traumatically injured spinal cord both in laboratory settings and in clinical applications (Blight 1989; Fujihara and Miyoshi 1998; Haghighi et al. 1995; Hayes et al. 1993, 1994; Jensen and Shi 2003; Jones et al. 1983; Stefoski et al. 1987). However, despite its effectiveness, the overall benefit resulting from 4-AP treatments remained modest. The main reason is that the achievable concentration of 4-AP in vivo is two orders of magnitude below its maximal effective dosage seen in vitro due to severe side effects seen in higher concentrations in live animals and human (Blight et al. 1991; Donovan et al. 2000; Halter et al. 2000). The common side effects include respiratory distress, anxiety, and epileptiform seizures (Pena and Tapia 1999, 2000; Stork and Hoffman 1994). Therefore alternative blockers that target the same channel but perhaps with greater efficacy and fewer side effects are highly desirable and warranted.

To pursue such a goal, we have in the last few years successfully synthesized three 4-AP derivatives that can significantly enhance axonal conduction following spinal cord trauma (McBride et al. 2006, 2007; Smith et al. 2005; Sun et al. 2009). More importantly, it was demonstrated that axons rescued by 4-AP derivatives can conduct action potentials in a manner that resembles normal axons more than those rescued by 4-AP (McBride et al. 2007). The minimal effective dosage of these 4-AP derivatives, however, is similar to that of 4-AP (at 1 \( \mu \)M) which signifies the similar potency between 4-AP and its derivatives at the achievable level in vivo (McBride et al. 2006).

To continue this line of experimentation, we now report a new and fourth blocker in this series, 4-aminopyridine-3-methanol (4-AP-3-MeOH; Fig. 1). This compound can significantly enhance axonal conduction following mechanical trauma. Similar to the previous three carbamates, 4-AP-3-MeOH also restores conduction of injured axons in a manner similar to normal uninjured axons (McBride et al. 2007). However, a unique feature of 4-AP-3-MeOH is its high potency: the lowest effective dosage is 0.1 \( \mu \)M, which is one order of magnitude lower than 4-AP and its derivatives (Jensen and Shi 2003; McBride et al. 2006; Shi and Blight 1997; Shi et al. 1997). In addition, we have found that 4-AP-3-MeOH is indeed capable of blocking fast potassium channels, which is likely responsible for restoring axonal conduction. We have also confirmed that potassium channels are indeed exposed following mechanical stretch in the current study. This further
strengthens the notion that potassium channels are a key pathology in myelin damage that contribute to conduction block and an important therapeutic target to restore axonal conduction.

METHODS

Isolation of spinal cord

The adult female guinea pig (300–450 g) was anesthetized with a combination of ketamine (80 mg/kg), xylazine (12 mg/kg), and acepromazine (0.8 mg/kg). After anesthesia the guinea pig was perfused with oxygenated, cold Krebs’ solution [containing (in mM) 124 NaCl, 2 KCl, 1.2 KH2PO4, 1.3 MgSO4, 2 CaCl2, 10 dextrose, 26 NaHCO3, and 10 sodium ascorbate]. The vertebral column was extracted, and spinal cords were isolated as previously described (Shi and Blight 1996; Shi and Pryor 2000; Shi et al. 2000). The protocol was approved by Purdue University Animal Care and Usage Committee (PACUC).

Double sucrose gap recording

A double sucrose gap chamber was used for compound action potential (CAP) recording as previously described (Shi and Blight 1996; Shi and Borgens 1999; Shi and Whitebone 2006). A 4.5-cm-long spinal cord ventral white-matter strip was placed across the chamber. The central compartment was continuously perfused with oxygenated Krebs’ solution. The side compartments of the chamber were filled with isotonic potassium chloride (120 mM). Sucrose solution (320 mM) was perfused in the gaps for isolation. The temperature was maintained at 37°C for the duration of recording by a line heater and temperature probe (Warner Instruments, Hamden, CT). A bridge amplifier (Neurodata Instruments, Delaware Water Gap, PA) was used for recording, and data analysis was performed using customized Labview software (National Instruments, Austin, TX). 4-AP-MeOH was dissolved in Krebs’ solution and directly delivered to the spinal cord strip by circulating it into the central compartment. 4-AP-MeOH powder was purchased from Alfa Aesar (Ward Hill, MA). According to the MSDS provided by Alfa Aesar, the purity of 4-AP-MeOH is 100%. The MSDS sheet of 4-AP-MeOH could be obtained from http://www.alfa.com/content/msds/USA/H50033.pdf.

Stretch injury model

An injury device that has a constant stretch magnitude and falling speed was used to induce stretch injury in this study (Jensen and Shi 2003; McBride et al. 2007; Shi and Pryor 2002). A stretch rod was released and falling at a speed ~1.5 m/s to induce the stretch injury to the spinal cord strip. A flat raised stage with a central hole was built in the central compartment of the double sucrose chamber, and the stretch rod would fall through the hole. A nylon mesh was laid on top of the chamber to stabilize the strip of spinal cord. The rod and nylon mesh were removed immediately from the spinal cord after injury (Jensen and Shi 2003; McBride et al. 2007; Shi and Pryor 2002).

Coherent anti-Stokes Raman scattering imaging and immunofluorescence

Coherent anti-Stokes Raman scattering (CARS) imaging was used to visualize the myelin sheath directly without any staining (Fu et al. 2007; Wang et al. 2005). For immunofluorescence imaging, a 1-cm post-injury segment was cut from the central injury site and fixed in 4% paraformaldehyde for 24 h and followed with cryoprotection (incubation in 20% glycerol and 2% paraformaldehyde in PBS at 4°C) for 48 h. The strip was then sectioned into smaller pieces (~50 μm) using an oscillating tissue slicer (OTX-4000, Electron Microscopy Sciences, Hatfield, PA). After that, sections were incubated in 5% bovine serum albumin (BSA) with 0.5% Triton X-100 in PBS (pH 7.4) for blocking for 1 h at the room temperature. After rinsing with PBS, the sections were then incubated in 1:100 diluted Rabbit raised anti-Kv-1.2 (Alomone Labs, Jerusalem, Israel; 1% BSA with 0.1% Triton X-100 in PBS) at 4°C for 24 h. After being rinsed three times in PBS for total 30 min, the sections were then incubated in goat anti-rabbit IgG conjugated with 1:100 diluted FITC-goat anti-rabbit IgG (H+L) (Invitrogen, Carlsbad, CA; 1% BSA with 0.1% Triton X-100 in PBS) for 1 h at the room temperature. Control sections were incubated with the secondary antibody only. After rinsing three times in PBS for 30 min, the sections were imaged by a laser scanning confocal microscope (FV3000/IX70, Olympus) with a 488-nm laser. The nodal ratio is calculated as nodal length divided by axon diameter. The nodal length was defined as the distance between paranodal myelin with strong CARS contrasts at the two ends along the node, and the axon diameter was defined as the distance between paranodal myelin at two sides across the node (Fig. 10B).

Acute dissociation of guinea pig DRG cells

The guinea pig DRG cell dissociation process has been described in detail previously (Sun et al. 2009). Briefly, dorsal root ganglia are collected from all accessible segments of guinea pig vertebra and digested with trypsin (type I, 0.25 mg/ml, Sigma-Aldrich) and collagenase (type I, 0.5 mg/ml, Sigma-Aldrich) at 37°C for 30 min. The dissociated dorsal root ganglion cells were plated on poly-L-lysine (0.2 mg/ml, coating overnight, Sigma-Aldrich)-coated dishes. They were cultured in medium [containing 48.5% DMEM (Sigma-Aldrich), 48.5% F-12 Nutrient Mixture (Gibco), 2% horse serum (Sigma-Aldrich), and 1% penicillin-streptomycin solution (Sigma-Aldrich)] at 37°C (5% CO2) balance air, for 2.5 h before recording.

Whole cell patch-clamp recording

Currents were recorded using the Axopatch 200B amplifier (Molecular Devices, Union City, CA). Borosilicate glass tubing (Sutter Instrument, Novato, CA) was pulled using P-97 horizontal puller (Sutter Instruments) to make a recording pipette with resistances at ~4 MΩ. Pipette solution included (in mM) 140 KCl, 1 MgCl2, 5 EGTA, and 10 HEPS titrated to pH 7.4 with KOH. Bath solution included (in mM) 145 NaCl, 2 KCl, 1 MgCl2, 0.03 CaCl2, and 10 HEPS and 1 μM TTX titrated to pH 7.4 with NaOH. Holding potential was ~70 mV. The currents were filtered at 1–5 kHz, and the data were acquired and analyzed with pCLAMP 9.2 software (Molecular Devices).

Isolation of voltage-gated potassium currents

Na+ currents were blocked by 1 μM tetrodotoxin (TTX) in bath solution, and Ca2+ currents and Ca2+-dependent potassium currents were suppressed with low Ca2+ concentration (0.03 mM) in bath solution.
solution. There are mainly two voltage-gated potassium (Kv) currents on guinea pig DRG cells: a transient fast activating A-type current ($I_{A}$) and a slow sustained delayed rectifier type current ($I_{K}$) (Xu et al. 2006). $I_{A}$ can be separated from the total potassium currents recorded from DRG cells using a traditional electrophysiological method that is widely used in previous studies reported in the literature (Everill et al. 1998; Tan et al. 2006). The total outward potassium currents ($I_{total}$) were acquired from a series of 400-ms voltage stimuli ranging from $-50$ to $40$ mV with 10-mV steps. Before the step stimuli, the cell was held at $-100$ mV (hyperpolarization) instead of $-70$ mV for 2 s. When the membrane potential was held at $-30$ mV (depolarization) for 2 s before the step stimuli, the fast-activating component $I_{A}$ stays in the inactivation phase during the 400-ms step stimuli, and all the collected potassium currents were slow delay rectified component $I_{K}$. All currents were off-line leak subtracted. 4-AP (5 mM) and 5 mM 4-AP-3-MeOH were applied into the bath solution respectively. 4-AP and 4-AP-3-MeOH were both freshly made with bath solution before recording.

**Statistical analysis**

Paired t-test was used to compare dose responses of 4-AP-3-MeOH after stretch injury, inhibition of potassium currents by 4-AP and 4-AP-3-MeOH as well as some electrophysiological tests analysis. Student’s t-test was applied to analyze nodal ratio elongation.

**RESULTS**

**4-AP-3-MeOH enhances compound action potential propagation in stretch-injured spinal cord white matter**

After the stabilization of CAP recorded using double sucrose gap (Shi and Blight 1996; Shi and Borgens 1999; Shi and Whitebone 2006), a stretch injury was applied that initially completely abolished CAP conduction (Jensen and Shi 2003). The CAP then steadily recovered to reach a plateau after an average of 45 min (Fig. 2). The average amplitude of recovered CAP is $\sim 21\%$ of original CAP before the injury. 4-AP-3-MeOH was then applied through Krebs’ circulation in the central compartment of double sucrose gap chamber. On average, the 4-AP-3-MeOH was applied for 40–50 min followed by a period of wash at 30–40 min.

As demonstrated in Fig. 3, 4-AP-3-MeOH significantly increased CAP conduction in the following four concentration groups (Fig. 3): 0.1 $\mu$M (12.73 $\pm$ 3.2$,\%$, mean $\pm$ SE; $P < 0.01, n = 7$), 1 $\mu$M (11.27 $\pm$ 4.2$,\%$, $P < 0.05, n = 6$), 10 $\mu$M (13.35 $\pm$ 5.9$,\%$, $P < 0.05, n = 11$) and 100 $\mu$M (21.51 $\pm$ 4.5$,\%$, $P < 0.01, n = 6$). CAP was not significantly increased at 0.01 $\mu$M (6.44 $\pm$ 3.8$,\%$, $P > 0.05, n = 5$) and 1 mM (6.72 $\pm$ 8.3$,\%$, $P > 0.05, n = 5$). Similar to 4-AP and other 4-AP derivatives, 4-AP-3-MeOH at 10 mM suppressed CAP conduction (18.48 $\pm$ 8.1$,\%$, $P > 0.05, n = 5$).

**4-AP-3-MeOH has no significant effect on activation threshold**

A range of stimulus voltages was employed to assess CAP activation threshold. Stimulus intensities from 1.85 to 6.5 V were applied to injured spinal cord tissue pre and post 100 $\mu$M 4-AP-3-MeOH treatment. It is clear that 4-AP-3-MeOH increased CAP conduction at all stimuli intensity levels (Fig. 4). In addition, the CAP amplitude-stimulus intensity curves were similar in pre and after 4-AP-3-MeOH treatment (Fig. 4B). Figure 4C demonstrated the linear correlation between pre-drug and drug amplitude (%max) and 4-AP-3-MeOH treated amplitude (%max). The slope of curve 1 indicates that 4-AP-3-MeOH induced conduction enhancement was not biased toward axons with different thresholds for activation.

**4-AP-3-MeOH has no significant effect on axonal response to dual and multiple stimuli**

To detect the effect of 4-AP-3-MeOH on spinal cord’s response to multiple stimuli, we measured the refractory period and the response to train stimulus, respectively. For refractory period, spinal cord tissue was stimulated by dual stimuli with various interval times, ranging from 0.5 to 15 ms. Absolute refractory period (interval time that second peak starts to appear) and relative refractory period (interval time that 2nd peak amplitude is no <95% of the 1st peak) were determined. It is demonstrated that 4-AP-3-MeOH at 100 $\mu$M has little effect on both absolute and relative refractory periods of injured spinal cord tissue ($P > 0.05$, Fig. 5).

We also measured the response of injured spinal cord in response to multiple (or train) stimuli after 4-AP-3-MeOH administration. A train stimulus at the duration of 100 ms with high and low frequency (1,000 and 500 Hz, respectively) was applied to the injured spinal cord strip. The representative CAP response is shown in Fig. 6A. The CAP response is quantified by averaging the last four CAP peaks as the percentage of first peak. Figure 6B demonstrates the comparison of CAP responses in pre-drug and drug conditions. The CAP response following train stimuli (both high and low frequency) is not significantly affected by application of 4-AP-3-MeOH ($P > 0.05$ in comparisons of pre-drug and drug, 1,000 and 500 Hz).
4-AP-3-MeOH blocks fast potassium channels in guinea pig DRG neurons

Following the assertion that 4-AP-3-MeOH enhances CAP conduction in injured spinal cord, we investigated whether 4-AP-3-MeOH indeed blocks fast outward potassium currents as speculated. This was carried out using a patch-clamp technique (whole cell patch configuration) on guinea pig DRG cells where fast potassium currents are known to exist (Xu et al. 2006). Figure 7A shows the total voltage-gated potassium currents ($I_{\text{total}}$) recorded from guinea pig DRG cells elicited with a series of 400-ms step pulses. With a 2-s prepulse hyperpolarizing holding potential ($/H_{11002}$100 mV), the potassium currents are activated to a peak (transient) and then partially decayed to a sustained level. There are two significant and distinguishable components contributing to this current: one is fast activated and fast inactivated current called $I_A$, represented by the transient peak in the $I_{\text{total}}$; the other component, delay rectifier, $I_{\text{dr}}$, is relatively slowly activated but sustained during the step pulse duration (Stewart et al. 2003; Tan et al. 2006; Xu et al. 2006).

When we applied 5 mM 4-AP through the bath solution, we found the initial peak was largely blocked while the sustained delayed rectifier component remained relatively unchanged (Fig. 7B). 4-AP-sensitive potassium current is shown in Fig. 7C by subtracting the current after treatment from pretreatment (A and B). Quantification analysis of drug effect was carried out at a particular command potential of $/H_{11002}$30 mV (Fig. 7D). When compared at the initial peak current (which is largely $I_A$), the $I_{\text{total}}$ amplitude decreased from 4,306.38 ± 810.2 pA (peak value) to 2,574.91 ± 446.8 pA as a result of 4-AP application (Fig. 7, D and E, $P < 0.05$, $n = 6$, paired $t$-test). This has confirmed the existence of 4-AP-sensitive $I_A$.

FIG. 3. The change of CAP amplitude in injured spinal cord tissue in response to various concentrations of 4-AP-3-MeOH. The 4-AP-3-MeOH was applied after 30–40 min after injury and recovery. The increase in CAP amplitude as a result of 4-AP-3-MeOH was normalized to the pre-4-AP-3-MeOH level. CAP amplitude significantly increased in 4 concentration groups (from 0.1 to 100 μM). 4-AP-3-MeOH at 1 mM did not significantly enhance CAP amplitude ($6.72 ± 8.3%, n = 5, P > 0.05$). At 10 mM, 4-AP-3-MeOH began to show toxic effects ($−18.48 ± 8.1%, n = 5, P > 0.05$). All concentration groups were compared with control with a paired $t$-test. Error bars represent SE. *$P < 0.05$, **$P < 0.01$.

FIG. 4. A voltage test was performed to chart CAP responses (stretched cord) at different stimulus intensities and at 2 conditions: pre-drug and 100 μM 4-AP-3-MeOH-treated. CAP responses were shown in both superimposed form (A) and graphic plot form (B). Stimulus intensities ranged from 1.85 to 6.5 V, $n = 5$. Normalized CAP responses (as % of max CAP amplitude in per-drug condition) of injured spinal cord were plotted as pre-drug against 100 μM 4-AP-3-MeOH-treated (C). Original data in C are the same as for B. Overall trends demonstrated a linear relationship between pre-drug and the 4-AP-3 MeOH-treated group. Negligible difference in axon activation thresholds was found after 100 μM 4-AP-3 MeOH application. $n = 5$. Error bars represent SE.
Following the confirmation of the presence of 4-AP-sensitive fast potassium current, we then tested the ability of 4-AP-3-MeOH to inhibit the same fast potassium currents. Using the same protocol as 4-AP, we have shown that 4-AP-3-MeOH inhibits the fast potassium currents as well (Fig. 8, A–C). Specifically, at +30 mV, 5 mM 4-AP-3-MeOH decreased the fast activated component of potassium currents from 3,501.08 ± 592.5 to 2,434.20 ± 543.4 pA (Fig. 8, D and E, P < 0.05, n = 6, paired t-test). In addition to +30 mV, 4-AP-3-MeOH also suppresses $I_A$ at a wide range of step voltage commands from −50 to 40 mV (Fig. 9).

In addition to $I_{total}$, we also isolated delayed rectifier current $I_{dr}$, the slow-activated and long sustained component of the potassium current in DRG using a depolarization holding potential (Sun et al. 2009). It was clear that both 4-AP and 4-AP-3-MeOH did not significantly inhibit $I_{dr}$ (data not shown). Thus potassium current that is sensitive to 4-AP and 4-AP-3-MeOH are mostly the fast potassium current $I_A$.

**Stretch injury induces myelin damage at paranodal region and the exposure of voltage-gated potassium channels**

The abnormality of myelin structures at the paranodal region were examined using both CARS microscopy and immunohistochemistry. As a novel technique employed in this study, CARS provides excellent resolution in detecting myelin structures, particularly in the node of Ranvier without the need of any exogenous dye labeling (Wang et al. 2005). Figure 10A shows the CARS images of nodes of Ranvier in isolated spinal cord ventral white matter under normal conditions and 1.5 h after stretch injury. There is an obvious lengthening or widening of the node of Ranvier that is likely due to the retraction of myelin toward internodal regions. We also noted a weak...
The correlation between node ratios and axonal diameter in both control and stretch groups (Fig. 10B). However, there is an obvious increase in the node ratios in the stretched group compared with control group (Fig. 10B). Specifically, the nodal ratio, an indication of nodal lengthening, is increased from a normal level of $0.80 \pm 0.1$ ($n=121$) to $2.88 \pm 0.2$ ($n=94$; Fig. 10C, $P<0.01$, Student’s $t$-test).

In addition, there is also an obvious paranodal myelin split (or decompaction) in stretched spinal cord strip. Although varying in severity, this splitting or loosening of paranodal myelin can be seen in most of the stretched ventral white matter (Fig. 10A).

It has been reported that there are highly concentrated voltage-gated potassium channels (Kv1.1, Kv1.2, Kvβ2.1) in juxtaparade regions on myelinated axons (Poliak and Peles 2003; Rasband and Trimmer 2001; Vabnick and Shrager 1998). We hypothesize that the paranodal demyelination caused by stretch injury will expose those voltage-gated potassium channels. To test such hypothesis, we first examined the distribution of voltage-gated potassium channels after stretch injury using immunohistology. Specifically, we labeled Kv1.2 channels at 2 h after stretch injury. When observed using both CARS and two photon excitation fluorescence microscopy (TPEF), it is clear that such channels are clustered normally at juxtaparanodal region and covered by myelin. However, stretch-induced paranodal myelin damage clearly exposed the voltage-dependent potassium channels (Fig. 11, D–F).

**DISCUSSION**

4-AP-3-MeOH enhances CAP conduction after stretch

Using a well-established double sucrose gap recording technique, we have shown in the current study that 4-AP-3-MeOH, a 4-AP derivative, can significantly enhance action potential conductance in guinea pig spinal cord after mechanical stretch. This is the fourth 4-AP derivative in this line of investigation that demonstrates the capability to restore axonal conduction following SCI (McBride et al. 2006, 2007; Smith et al. 2005). Compared with previous three 4-AP derivatives, this com-
4-AP-3-MeOH preserves the advantage of retaining normal electrophysiological properties in restored axonal conduction (McBride et al. 2007). Furthermore, it possesses a unique benefit of being more potent than other three derivatives. First, 4-AP-3-MeOH showed no preference in restoring in axonal conduction in large or small axons that is similar to previous three derivatives, N-(4-pyridyl)-methyl carbamate, N-(4-pyridyl)-ethyl carbamate, and N-(4-pyridyl)-tertbutyl (Fig. 4) (McBride et al. 2007). Second, in 4-AP-3-MeOH-restored axonal conduction, relative or absolute refractory periods are not significantly altered compared with that of the normal axons (Fig. 5). Similarly, those axons rescued by 4-AP-3-MeOH retained the normal ability to follow repetitive or train stimuli (Fig. 6). It is worth mentioning that while restoring axonal conduction, 4-AP causes significant reduction in axonal responsiveness by increasing the absolute and relative refractory period as well as decreasing the ability of the cord to respond to repetitive stimuli (Jensen and Shi 2003; Targ and Kocsis 1986). In summary, similar to the other three 4-AP derivatives, the axons that are rescued by 4-AP-3-MeOH can conduct action potentials in a manner that is similar to normal axons and superior to those rescued by 4-AP.

Another unique feature of 4-AP-3-MeOH is its high potency compared with other three derivatives and 4-AP: the lowest effective concentration offering conduction enhancement is between 0.01 and 0.1 μM (McBride et al. 2006; Shi et al. 1997). This is about a 10-fold increase of potency compared with other three derivatives and 4-AP (McBride et al. 2006; Shi et al. 1997). Because the side effect of 4-AP is dose related (Shi et al. 1997), it is reasonable to speculate that a lower effective dosage may be accompanied by reduced possibility of inducing serious side effects.

**4-AP-3-MeOH inhibits fast potassium current**

In the current study, we showed that 4-AP-3-MeOH significantly inhibits an early and transient potassium current in guinea pig DRG cells. Based on its nature of fast activation as well as inactivation, it is likely a fast potassium channel or $I_A$.

**Fig. 9.** $I-V$ curve of total potassium currents ($I_{\text{total}}$) recorded from guinea pig dorsal root ganglion (DRG) cells. Potassium currents significantly decreased with 5 mM 4-AP-3-MeOH treatment (○) comparing to pretreatment (□) record at most command potential levels. Command potentials ranged from −50 to 40 mV with 10-mV steps. Error bars represent SE. n = 5.

**Fig. 10.** Stretch-injury-induced paranodal splitting and retraction. A: coherent anti-Stokes Raman scattering (CARS) imaging showed paranodal splitting and retraction after acute stretch injury to spinal cord white-matter strip. ←, node of Ranvier regions. Scale bar: 10 μm (for left and right). B: node ratios of both control (●) and stretch groups (○) were plotted against axonal diameter. Note the weak correlation of nodal ratios and axonal diameters. However, there is a conspicuous increase of node ratios in the stretched group compared with the control group. C: comparison of nodal ratio (node length divided by axon diameter, see inset) in control group and stretched group. The nodal ratio was significantly larger in the stretched group (2.88 ± 0.2, n = 94) vs. the control group (0.80 ± 0.1, n = 121; P < 0.01, Student’s t-test). Error bars represent SE.
In the current study, using a combination of traditional and novel multimodal imaging techniques (CARS and 2-photon excitation fluorescence microscopy), we have provided unequivocal anatomical evidence demonstrating the exposure of potassium channels at the juxtaparanodal region. This is particularly true in acute spinal cord trauma when the myelin damage begins to emerge (Karimi-Abdolrezaee et al. 2004). In the current study, using a combination of traditional and novel multimodal imaging techniques (CARS and 2-photon excitation fluorescence microscopy), we have provided unequivocal anatomical evidence demonstrating the exposure of potassium channels at the juxtaparanodal region. This is particularly true in acute spinal cord trauma when the myelin damage begins to emerge (Karimi-Abdolrezaee et al. 2004).
4-AP-3-MeOH restores axonal conduction in spinal cord

Hypothesis and observation under similar situations (Karimi-Abdolrezaee et al. 2004; Nashmi et al. 2000).

In summary, we have demonstrated the ability of 4-AP-3-MeOH, a 4-aminopyridine derivative, to significantly restore axonal conduction and act as an effective IA blocker. This compound has the advantage of being 10 times more potent when compared with 4-AP and other derivatives. Furthermore, unlike 4-AP, 4-AP-3-MeOH can restore axonal conduction without changing their electrophysiological properties. We have also provided the critical evidence to confirm the exposure of potassium channels after mechanical injury. Taken together, our data further support the role of potassium channels after mechanical injury. Without changing their electrophysiological properties, we are able to restore axonal conduction and act as an effective blocker. This compound has the advantage of being 10 times more potent when compared with 4-AP and other derivatives. Furthermore, unlike 4-AP, 4-AP-3-MeOH can restore axonal conduction without changing their electrophysiological properties. We have also provided the critical evidence to confirm the exposure of potassium channels after mechanical injury. Taken together, our data further support the role of potassium channels after mechanical injury.

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