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Autophosphorylated CaMKII Facilitates Spike Propagation in Rat Optic Nerve

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Repeated spike firing can transmit information at synapses and modulate spike timing, shape, and conduction velocity. These latter effects have been found to result from voltage-induced changes in ion currents and could alter the signals carried by axons. Here, we test whether Ca^{2+} /calmodulin-dependent protein kinase II (CaMKII) regulates spike propagation in adult rat optic nerve. We find that small-, medium-, and large-diameter axons bind anti-Thr286-phosphorylated CaMKII (pT286) antibodies and that, in isolated optic nerves, electrical stimulation reduces pT286 levels, spike propagation is hastened by CaMKII autophosphorylation and slowed by CaMKII dephosphorylation, single and multiple spikes slow propagation of subsequently activated spikes, and more frequent stimulation produces greater slowing. Likewise, exposing freely moving animals to flickering illumination reduces pT286 levels in optic nerves and electrically eliciting spikes *in vivo* in either the optic nerve or optic chiasm slows subsequent spike propagation in the optic nerve. By increasing the time that elapses between successive spikes as they propagate, pT286 dephosphorylation and activity-induced spike slowing reduce the frequency of propagated spikes below the frequency at which they were elicited and would thus limit the frequency at which axons synaptically drive target neurons. Consistent with this, the ability of retinal ganglion cells to drive at least some lateral geniculate neurons has been found to increase when presented with light flashes at low and moderate temporal frequencies but less so at high frequencies. Activity-induced decreases in spike frequency may also reduce the energy required to maintain normal intracellular Na^+ and Ca^{2+} levels.

Key words: CaMKII; conduction velocity; interspike interval; optic nerve; retino-geniculate synapse

Significance Statement

By propagating along axons at constant velocities, spikes could drive synapses as frequently as they are initiated. However, the onset of spiking has been found to alter the conduction velocity of subsequent (“follower”) spikes in various preparations. Here, we find that spikes reduce spike frequency in rat optic nerve by slowing follower spike propagation and that electrically stimulated spiking *ex vivo* and spike-generating flickering illumination *in vivo* produce net decreases in axonal Ca^{2+} /calmodulin-dependent protein kinase II (CaMKII) autophosphorylation. Consistent with these effects, propagation speed increases and decreases, respectively, with CaMKII autophosphorylation and dephosphorylation. Lowering spike frequency by CaMKII dephosphorylation is a novel consequence of axonal spiking and light adaptation that could decrease synaptic gain as stimulus frequency increases and may also reduce energy use.

Introduction

Axons generate, conduct, and, under some conditions, also modulate spikes. At low and moderate stimulus frequencies, spikes

typically propagate without changes in amplitude, firing probability, and velocity. By contrast, higher frequencies can modulate

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spike amplitude, duration, propagation speed, and/or firing probability (Swadlow et al., 1980; Debanne et al., 1997; Williams and Stuart, 1999; Geiger and Jonas, 2000; Raastad and Shepherd, 2003; Khaliq and Raman, 2005; Meeks et al., 2005; Monsivais et al., 2005). At synapses that transmit information efficaciously if more than one spike arrives at the synaptic terminal within a narrow range of interspike intervals (Singer et al., 1972; Mastrojarre, 1987; Stevens and Wang, 1995; Usrey et al., 1998; Sincich et al., 2007; Weyand, 2007), axons that maintain steady firing probabilities and conduction velocities would be particularly useful because they preserve the time (i.e., interspike interval) that elapses between spikes while they propagate. On the other hand, if the firing of one spike altered the conduction velocity of a subsequently elicited spike, then interspike intervals would increase or decrease as the spikes propagate, depending on whether the first (cardinal) spike travels more quickly or slowly than the next (follower) spike. This, in turn, could modulate information transfer at synapses that require particular interspike intervals to transmit signals to postsynaptic cells.

What cellular mechanisms control interspike intervals in axons? For three reasons, we begin to address this question by recording from optic nerve. First, some studies have found that single cardinal spikes alter the conduction velocity of subsequently elicited spikes in retinal ganglion cell (RGC) axons (Bishop et al., 1953; Foster et al., 1982), whereas other studies have reported that this does not occur (George et al., 1984). Second, voltage-gated Ca^{2+} channels, Ca^{2+} /calmodulin-dependent protein kinase II (CaMKII), and changes in intracellular Ca^{2+} concentration have been detected in RGC axons (Brown et al., 2001; Calkins et al., 2005; Weitz et al., 2013) and these alter spike propagation in other preparations (Hodgkin and Keynes, 1956; Lüscher et al., 1996; Tsubokawa et al., 2000). Third, previous studies have shown that illumination activates CaMKII in RGC somata (Ogata et al., 2012) and that electrical stimulation elevates Ca^{2+} concentration in optic nerve axons (Lev-Ram and Grinvald, 1987; Verbny et al., 2002). However, these studies did not identify conditions that activate CaMKII in optic nerve and did not show that changes in CaMKII activation alter spike propagation.

Here, we test whether cardinal spikes alter the propagation speed of follower spikes, whether illumination and electrical stimulation activate CaMKII, and whether changes in CaMKII autophosphorylation alter spike propagation, all in adult rat optic nerve. We find that single cardinal spikes slow the propagation of follower spikes; that repetitive spiking reduces the level of autophosphorylated CaMKII (pT286) and slows follower spike propagation further; that an inhibitor of CaMKII activation slows cardinal and follower spike propagation; that a protein phosphatase inhibitor increases pT286 levels and accelerates cardinal and follower spike propagation; that optic nerve axons bind antibodies against pT286 and protein phosphatase 2A; and that pT286 levels are reduced by spiking elicited *in vivo* by flickering illumination. In addition, we electrically stimulate adult rat optic chiasm and optic nerve *in vivo* and find that spiking slows propagation of subsequent spikes in the optic nerve. By increasing interspike intervals between cardinal and follower spikes, activity-induced spike slowing would tend to limit signal transmission, as observed at retinogeniculate synapses during responses to light flashes at high temporal frequencies. By reducing Na^+ and Ca^{2+} influx, interspike interval increases could also reduce the energy used by RGCs to maintain normal intracellular Na^+ and Ca^{2+} levels.

Materials and Methods

Animals. Adult rats were used for this study based on the presence and autophosphorylation of CaMKII in their RGC somata (Bronstein et al., 1988; Terashima et al., 1994; Ogata et al., 2012) and the presence of CaMKII in their optic nerves (Lund and McQuarrie, 2001). Long-Evans rats [male and female; postnatal day 60 (P60)–P120; 150–250 g] were obtained from a commercial supplier (Envigo Bioproducts) and housed in standard cages at $\sim 23^\circ\text{C}$ on a 12 h/12 h light/dark cycle. For the optic nerve experiments reported here, each rat was anesthetized by intraperitoneal ketamine and xylazine (70–100 and 5–10 mg/kg, respectively; see below for the source of chemicals used in this study) and decapitated. Each optic nerve was isolated by dissecting it from the back of each eye and from the optic chiasm and passing it through the optic foramen. These nerves were Western blotted, immunohistochemically stained, or electrophysiologically tested, as described below. For the retina experiments reported here, each rat was anesthetized as described above, enucleated, and killed by a lethal dose of Fatal Plus (150 mg/kg, intraperitoneal). After hemisectioning the eyes, the retinae were dissected away from the posterior eyecups and electrophysiologically tested as described below. All animal care and experimental protocols were approved by the Animal Use and Care Administrative Advisory Committee of the University of California–Davis. The *in vivo* optic nerve spikes were recorded by modification of a protocol described in detail previously (Tang et al., 2016). Briefly, adult Brown–Norway rats (male, 300–400 g; Harlan Laboratories) were anesthetized by intraperitoneal injection of ketamine hydrochloride (75 mg/kg) and xylazine (7.5 mg/kg), supplemented as needed. Animals were rested on a heating pad (37°C) and maintained under anesthesia for the remainder of the experiment via an intravenous infusion of ketamine (30 mg/kg · h), xylazine (1.5 mg/kg · h), and 0.9% physiological saline. A tracheotomy was performed to provide mechanical ventilation if necessary. Body temperature and heart rate were monitored with a rectal thermometer and ECG electrodes, respectively, and data were collected while vital signs remained at normal levels. Animals were killed with Euthasol (50 mg/kg) at the end of experiments. All procedures were approved by the Institutional Animal Care and Use Committee of the University of South Florida in accordance with National Institutes of Health guidelines.

Protein isolation. For Western blotting, optic nerves were placed in a capped microcentrifuge tube, frozen by dropping into liquid nitrogen, and homogenized in solubilization buffer (320 mM sucrose, 50 mM MgCl_2 , 7 M urea, 10 mM dithiothreitol, 100 mM Tris-HCl, pH 7.4). The homogenates were centrifuged at $600 \times g$ for 10 min at 4°C . The resulting pellets were discarded and the supernatant was concentrated in centrifugal filters (30 kDa, Amicon). The concentrated homogenate was diluted in 200 μl of homogenization buffer (320 mM sucrose, 50 mM MgCl_2 , 100 mM Tris-HCl, pH 7.4), concentrated again, and then diluted in a final volume of 65 μl of homogenization buffer. The total protein concentration was determined using a bicinchoninic acid protein assay kit. All solutions were ice-cold and supplemented with protease and phosphatase inhibitor tablets.

Western blot. Aliquots of optic nerve homogenates were loaded into 4–12% gradient gel lanes (Bis-Tris; Invitrogen), electrophoretically separated in MOPS running buffer at 200 V for 1 h, and transferred to polyvinylidene difluoride (PVDF) membranes (Bio-Rad) at 30 V for 1 h. Protein standards (SeeBlue Plus2) were run in lanes adjacent to the samples. Each PVDF membrane was immersed for 1 h in a protein-supplemented buffer solution (Superblock) and then incubated with primary antibodies on a rocker overnight at 4°C . As a loading control, lanes were also immunostained for β -actin, myelin basic protein, or CaMKII. The membranes were then rinsed with TBS (supplemented with 0.5% Tween 20) and incubated in species-specific secondary antibody conjugated to fluorophores for 1 h at room temperature. The fluorescence of these fluorophores was visualized and imaged in a digital imager (FluorChem Q; Alpha Innotech). The fluorescence intensities of the immunostained protein bands were measured with software of the imager (AlphaView Q, ProteinSimple) and normalized to that of the loading control protein.

Fixation and immunohistochemistry. For immunostaining, optic nerves were immersed overnight at 4°C in a fixative containing formal-

dehyde (4% w/v) in PBS, pH 7.4, rinsed, embedded in 5% (w/v) low-melt agarose at 37°C, allowed to set for at least 60 min at 4°C, and cut into blocks with a razor blade. These blocks were sectioned transversely or longitudinally at a thickness of 75–100 μm on a vibrating microtome (VT1000S; Leica Microsystems). Free-floating sections were blocked and permeabilized for 24 h at 4°C in TBST (TBS plus 0.3% v/v Triton X-100, pH 7.4) supplemented with 5% normal donkey serum. The sections were incubated in primary antibody (either overnight at 37°C or for as long as 5 d at 4°C), rinsed with PBS, incubated in secondary antibody (overnight at either 37°C or 4°C), rinsed with PBS, and mounted in Prolong Diamond Antifade Mountant between glass coverslips (no. 1.5) and glass slides. Primary and secondary antibodies were diluted in TBST.

Antibody characterization. The mouse monoclonal anti-neurofilament antibody (BioLegend, clone SMI-312, RRID:AB_2566782) was generated against pan axonal neurofilaments, validated on Western blots by the manufacturer, and used at a dilution of 0.5 $\mu\text{g}/\text{ml}$.

The rat monoclonal anti-myelin basic protein (MBP) IgG2a antibody (Abcam, clone 12, RRID:AB_305869) was generated against full-length cow MBP, binds to a region defined by aa 82–87, and used at a dilution of 1:500. Antibody specificity was tested by the manufacturer using a synthetic peptide consisting of aa 82–99.

The mouse monoclonal anti-protein phosphatase 1 α (PP1 α) IgG1 antibody (Thermo Fisher Scientific, clone 10C6-3, RRID:AB_2533541) was generated against a mixture of two peptides from the C-terminal regions of human PP1 α protein and diluted to 2 $\mu\text{g}/\text{ml}$. The manufacturer demonstrated antibody specificity by siRNA-mediated knock-down of the target protein.

The rabbit polyclonal anti-protein phosphatase 2A α catalytic subunit (PP2A α C subunit) IgG antibody (Thermo Fisher Scientific, RRID:AB_10983779) was generated against a synthetic peptide corresponding to amino acids at the C-terminus of human PP2A catalytic subunits and used at a dilution of 1:200. The manufacturer verified specificity by Western blot and immunohistochemistry.

The rabbit monoclonal anti-phospho-CaMKII IgG antibody (Cell Signaling Technology, clone D21E4, RRID:AB_2713889) was generated against a synthetic phosphopeptide corresponding to residues surrounding Thr287 of human CaMKII- β protein and diluted to 1.85 $\mu\text{g}/\text{ml}$. The rabbit polyclonal anti-phospho-CaMKII IgG antibody (Abcam, RRID:AB_305050) was generated against a synthetic phosphopeptide derived from a region of human CaM Kinase II- α that contains threonine 286 and diluted to 1.5 $\mu\text{g}/\text{ml}$. The immunizing peptide (Cell Signaling Technology, catalog #1712S) was used to control against nonspecific binding by both primary antibodies.

The mouse monoclonal anti-CaMKII α IgG_{2a} antibody (Santa Cruz Biotechnology, clone A-1, RRID:AB_626789) was generated against a synthetic peptide corresponding to residues 303–478 of mouse CaMKII α protein and diluted to 1:200. The manufacturer ran positive controls on rat brain extracts.

The goat polyclonal anti- β -actin antibody (Abcam, RRID:AB_306374) was generated against a synthetic peptide within human β actin amino acids 1–100 conjugated to keyhole limpet hemocyanin and diluted to 1:400. The manufacturer ran positive controls on Western-blotted HeLa, NIH3T3, PC12, and CHOK1 whole-cell lysates.

Signals attributed to the primary antibodies were visualized with donkey, species-specific, fluorophore-conjugated secondary antibodies (Jackson ImmunoResearch Laboratories, RRID:AB_2340684, RRID:AB_2492289, RRID:AB_2307351, RRID:AB_2340863) diluted typically to 1:500. The fluorophores were Alexa Fluor 488, Cy3, or Alexa Fluor 647.

Confocal imaging and image handling. Confocal images were acquired on a Leica TCS SP8 STED 3 \times high-sensitivity confocal microscope coupled with Leica Application Suite X software. The results reported here were collected by use of excitation lines for AF488 (499 nm), Cy3 (554 nm), and AF647 (653 nm), an oil-immersion 100 \times objective (numerical aperture = 1.4), optical sections at step sizes of 0.180 μm , and 5-frame Kalman averaging for each optical section. For most imaging sessions, some sections were processed without primary antibody (i.e., stained with secondary antibody alone) and the laser intensity, pinhole diameter, and photomultiplier gain were selected so that the secondary

antibody fluorescence intensity was indistinguishable from background fluorescence.

Datasets were imported into the ImageJ package Fiji (<http://imagej.net/Fiji>, RRID:SCR_002285). Changes in color space, adjustments to brightness or contrast (if any), and image overlays were made in Photoshop CS6 (Adobe Systems, RRID:SCR_014199).

Ex vivo optic nerve stimulation and recording. To test whether and how spiking alters spike propagation in rat optic nerve, we measured spikes elicited by depolarizing current pulses and repeated these measurements before, during, and after conditions that increase or decrease CaMKII autophosphorylation. Because effects of the reagents that we used typically develop over the course of several minutes and no study we know of has recorded intra-axonally from optic nerve axons for more than a few minutes (Eng et al., 1990), we used glass suction electrodes and carbon fiber electrodes to record continuously for hours while superfusing various media. In these recordings, each stimulation elicits a compound action potential (CAP) that typically displays two or three positive peaks (Sefton and Swinburn, 1964; Stys et al., 1991). The different peaks sum spikes propagating along axons of different diameters (Bishop et al., 1953) and these arise from different populations of RGCs (Boycott and Wässle, 1974). Ranked by the “time-to-peak” between the stimulus artifact and the maximum amplitude of each peak, these peaks will be denoted hereafter as P1, P2, and (if present) P3. The ratios of the time-to-peak of P2 to P1 and of P3 to P1 in our recordings agree with those reported for rat optic nerve responses to optic tract stimulation (Sefton and Swinburn, 1964).

For electrophysiological stimulation and recording, optic nerves were transferred to a Plexiglas recording chamber and connected to electrodes, with one end of each nerve gently drawn into a stimulating glass suction electrode (Stys et al., 1991) and the other end either drawn into a recording glass suction electrode or impaled with a recording carbon fiber electrode (Carbostar-1; Kation Scientific). Before commencing data collection, all optic nerves were equilibrated to Ames medium by continuous superfusion for 1 h (Stys et al., 1991). Spikes were then elicited while superfusing the nerves with control Ames medium and “test” (reagent-containing) media. All solutions were bubbled with carbogen, warmed by a substage heater to 30–32°C (Stys et al., 1992b) and flowed through the recording chamber at ~ 2 ml/min.

The stimulus pulses were delivered from an optically isolated stimulator (STG 4002; Multi Channel Systems) that was controlled by a PC computer program (MC_Stimulus II, version 3.0.1; Multi Channel Systems, RRID:SCR_014955). The stimuli consisted of 100- μs biphasic pulses of ± 1 –16 V (50:50 duty cycle, negative first). To test the effect of single spikes on subsequent spikes, we used interstimulus intervals (ISIs) that exceeded the duration of CAPs that we recorded (i.e., ≥ 3 ms). To test the effect of multiple spikes on subsequent spikes, we varied the number of “conditioning” pulses that preceded the “test” pulse. CAPs were amplified by a high impedance differential amplifier (npb EXT-02B; npi Electronic), recorded at a bandwidth of 100 Hz to 3 kHz, and sampled at 100 kHz using another PC program (pCLAMP, version 9.2.1.9; Molecular Devices, RRID:SCR_011323).

In vivo optic nerve stimulation and recording. To record spikes *in vivo*, a craniotomy was performed over the optic chiasm with the head supported in a stereotaxic apparatus. Brain tissue was removed and a bipolar needle electrode was lowered into the optic nerve at the base of the chiasm. The optic nerve was then transected behind the globe, the eye was discarded, and the glass tip of a suction electrode was positioned near the optic nerve stump. The nerve stump was gently aspirated into the suction electrode, which was filled with artificial CSF containing the following (in mM): 126 NaCl, 3 KCl, 2 CaCl₂, 2 MgCl₂, 1.2 NaH₂PO₄, 26 NaHCO₃, and 10 glucose, pH 7.45. A silver chloride wire coiled around the suction electrode tip served as the reference electrode. The pair of bipolar electrode leads was connected to an electrical stimulator (S48; Grass Instruments) and the pair of suction electrode leads was connected to an amplifier (Xcell-3x4; FHC). Spikes were activated antidromically by applying current pulses (0.1 ms) to the optic chiasm until a CAP was recorded in the optic nerve. In one preparation, spikes were also activated orthodromically by switching the connections of the leads (to stimulate the optic nerve and record at the optic chiasm). Evoked signals were

amplified differentially (10,000 \times), filtered (10 Hz to 1 kHz), digitally sampled at 100 kHz, stored on a hard drive, and processed offline.

These *in vivo* CAPs consisted of a larger peak and, in most cases ($n = 2$ of 3 preparations recorded from), a smaller peak on the falling phase of the large peak. Based on the order of appearance after the stimulus artifact (i.e., as in our *ex vivo* recordings), we termed the large peak P1 and the shoulder-like peak P2. The CAP elicited by each stimulus was found to be fitted by sums of two four-parameter Gaussian functions. Therefore, the time-to-peak values were measured from the beginning of the stimulus artifact to the maximum amplitude of the entire response (i.e., the unfitted peak) and to the maximum amplitude of the Gaussian functions. Each function was of the following form:

$$y = a + be^{-[(x-c)/2d]^2}$$

where a is a y -offset, b is the height at the peak of the Gaussian, c is time-to-peak, and d is a sharpness factor. Using the “fmincon” function of MatLab, the following constraints were incorporated in these fits: a ranged between -5 and $+5 \mu\text{V}$, b of P1 ranged from $20 \mu\text{V}$ to the maximum amplitude of the recording, b of P2 ranged from $5 \mu\text{V}$ to the maximum amplitude of the recording, c of P1 preceded the timing of the large peak, c of P2 followed the timing of the large peak, d ranged between 3 and 50 for P1, and d ranged between 3 and 75 for P2.

Multielectrode array stimulation and recording. In some experiments, axonal spikes were activated and recorded on multielectrode arrays (60MEA200/30iR-ITO; Multi Channel Systems) from the vitreal side of whole, isolated retinæ. These spikes were recorded in nine retinæ from a total of 13 pairs of stimulating and recording electrodes that were separated by a few hundred micrometers (cf. Sekirnjak et al., 2006). The stimuli were biphasic current pulses and the analog data were acquired at a sampling frequency of 25 kHz and stored on a hard disc. The time-to-peak values were measured from digital averages of the spikes elicited by ~ 100 repetitions of the stimuli per pair of stimulating and recording electrodes.

Effects of changing CaMKII autophosphorylation. Effects of reagents and experimental conditions that increase or decrease CaMKII autophosphorylation were tested by Western blotting optic nerves that were incubated in either control Ames medium or “test” (reagent-containing) medium; by recording spikes that were elicited while superfusing the optic nerves with control medium, test medium, and control medium again; or by recording spikes before, during, and after eliciting spikes. In the early stages of this study, to screen reagent effectiveness, to compare optic nerves over identical periods of time *ex vivo*, and to minimize any possible effect of day-to-day differences in dissection, single optic nerves were transversely sliced into two segments (each 4–6 mm in length) and the segments were incubated in test solution or in control solution. In a few pilot experiments that used okadaic acid, the optic nerve segments were incubated in fixed volumes of medium and these incubations were performed at room temperature (23°C) to minimize evaporation. Effects found were confirmed in subsequent experiments at 30–32°C.

Dark and light adaptation. Rats were dark and light adapted as described in detail previously (Ogata et al., 2012). Briefly stated, rats were placed in light-tight boxes at 7:00 P.M. At 1:00 P.M. of the following day, one of the boxes was kept dark, while the interior of the other box was illuminated by two arrays of 505 nm LEDs (Optek Technology) attached to the box ceiling. The light exposures were 250-ms flashes presented at 2 Hz over a period ranging from 30 s to 60 min. Each rat was then killed as described above and, under dim red light ($<1 \mu\text{W}/\text{cm}^2$ provided by a 650 nm LED), enucleated. The optic nerves were dissected under room light and processed by Western blotting.

Reagents. Reagents were obtained from the following sources: Abcam Biochemicals: tetrodotoxin; Bio-Rad: Tween 20; Calbiochem: dimethylsulfoxide, okadaic acid; Fisher Scientific: dimethylsulfoxide, PBS (Ca^{2+} - and Mg^{2+} -free, pH 7.4), TBS (pH 7.4), urea; Jackson ImmunoResearch: normal donkey serum; Life Technologies: ProLong Diamond Antifade Mountant; Roche: cOmplete Mini, PhosSTOP; Sigma-Aldrich: ASB-14, dithiothreitol, ethanol, formaldehyde, MgCl_2 , sucrose; Thermo Fisher Scientific: bicinchoninic acid protein assay kit, low-melting-point agarose, MOPS running buffer, Restore PLUS Western blot Stripping Buf-

fer, SeeBlue Plus2 Pre-Stained Protein Standard, SuperBlock, Triton X-100; United States Biological: Ames medium, Ca^{2+} - and Mg^{2+} -free Ames medium; Virbac: Euthasol; Vortech: Fatal Plus; and Western Medical Supply: ketamine, xylazine.

Experimental design and statistical analysis. Standard electrophysiological methods were used to measure the time-to-peak of elicited CAPs and spikes *ex vivo*. Each reported value is the time elapsing between the beginning of the stimulus capacitive current and the maximum amplitude of the axonal response analyzed using pCLAMP. We used optic nerves that were typically ~ 8 mm long and stimuli that elicited 2 or 3 peaks in the CAPs. Also, the data for a given condition were measured with only one type of recording electrode (glass or carbon fiber). However, some nerves fit better than others in the suction electrodes (cf. Stys et al., 1991). Moreover, we positioned the stimulating and recording electrodes where we obtained stable recordings even if this resulted in different interelectrode distances. As might therefore be expected, time-to-peak values differed from nerve to nerve. To minimize contributions of length differences to effects we tested for, we calculated changes for each optic nerve separately as a percentage increase or decrease relative to the control value and report the mean \pm SEM of the differences. For sufficiently large sample sizes, we used paired t tests to assess whether differences were statistically significant and report the mean, SEM, and value range, sample size, and p -value. For datasets consisting of relatively small sample sizes, we used signed-rank tests to assess whether differences were statistically significant and formed box plots to illustrate the median, mean, and range of the raw data.

During analysis of the *in vivo* recordings, individual CAPs were considered “unfittable” and deleted if the p -value of the correlation coefficient of the fit with the data was not significant at 0.01 or if the p -value of the coefficient of determination was $>1e^{-4}$. Fewer than 0.2% of the CAPs that we recorded *in vivo* were discarded for these reasons. Also, the CAPs in one preparation lacked a discernible P2 peak. Because the sum of two or more Gaussian functions did not yield adequate fits, analysis for this preparation was limited to locating the raw peak.

Western blotting was used to detect proteins of interest in optic nerves collected under various experimental conditions. Because this entailed tissue homogenization, optic nerve samples were processed in pairs, with one maintained under the test condition and the other under a control condition. The pairs were either cut from a single optic nerve, were the left and right optic nerves of a single animal, or were the optic nerves of animals kept under different lighting conditions and collected on the same day. The value reported for single optic nerve samples is the fluorescence intensity measured by a digital imager normalized to that of a loading control protein (see above). The difference between each pair of nerves is expressed as the ratio of the test value to the control value ($100 \times \text{experimental/control}$). The difference for each condition is the mean of the differences for the pairs used in each experiment. To test whether spiking alters CaMKII autophosphorylation levels, Western blots were formed from optic nerves that were snap frozen immediately after stimulation by pulse protocols that slow spike propagation (e.g., a single epoch of 16 conditioning pulses). In other cases, the stimuli were pairs of pulses delivered for a total of 10–120 s. The first and second pulse of each pair was separated by 1, 2, 3, or 5 ms (Fig. 1A). These intervals were presented in a randomized sequence and each pair of pulses was separated from the next pair by 1 s. Because the net effects of these protocols on pT286 were indistinguishable, the results were pooled.

Immunostaining experiments were performed on at least three optic nerves per reported observation. Axon diameter and myelin thickness were measured in an average of 300 axons in a total of 11 optic nerves. These measurements were made from transverse and longitudinal sections that included axons of various diameters. The axon diameters were assessed in longitudinal sections to sample the diameter of axon segments that were more than a few micrometers long. The inner and outer diameters of myelin were assessed in transverse sections to sample round, perpendicularly sliced myelin profiles.

All observations reported here are based on measurements made in one or both optic nerves or retinæ of at least three animals. Statistical analyses of the data were performed using Excel 2013 (Microsoft), MatLab R2014A (The MathWorks, RRID:SCR_001622), and R version

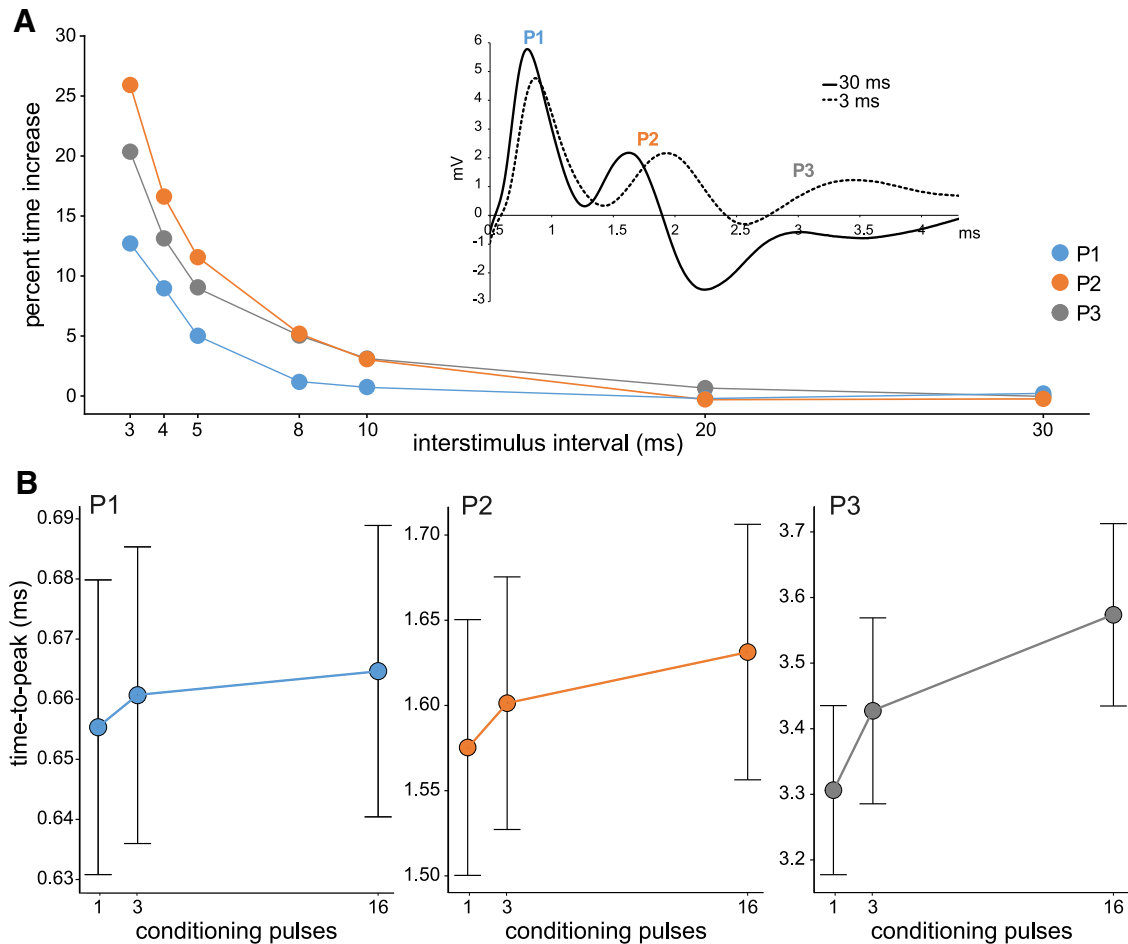


Figure 1. Slowing of optic nerve spike propagation by prior spiking. **A**, Inset, CAPs recorded with a glass suction electrode from an isolated rat optic nerve. CAPs were elicited by pairs of stimulus pulses; these stimuli were repeated 3 times for each ISI. The solid and dotted lines show the average of the “follower” CAPs (i.e., those elicited by the second pulse of each pulse pair) when the ISI was 30 ms (solid line) or 3 ms (dotted line). The peaks of the CAP are labeled P1, P2, and P3. **A**, Time-to-peak of P1, P2, and P3 at different ISI values. For each optic nerve, the time-to-peak of P1, P2, and P3 in the cardinal and follower CAPs were measured, normalized by the cardinal values, and multiplied by 100. Each dot plots the difference between the mean of the percentage cardinal and follower values for the optic nerves recorded from ($n = 15$, $n = 15$, and $n = 11$ for P1, P2, and P3, respectively). Note the lack of spike slowing at 30 ms and the monotonic increase in slowing as the ISI decreases. **B**, Increase in the time-to-peak of P1, P2, and P3 as the number of conditioning pulses increases. The data for P1, P2, and P3 are plotted in **A** and **B** by blue, orange, and gray dots, respectively. Dots and error bars plot the mean and SEM of the measured values.

3.4.1 (R Foundation for Statistical Computing, RRID:SCR_001905). Measurements presented without statistical analyses are given as mean \pm SEM and sample size.

Results

This study used extracellular stimulation and recording, Western blotting, and immunohistochemistry to test, in adult rat optic nerve, whether: (1) spiking alters the speed at which subsequent spikes propagate, (2) autophosphorylated CaMKII and associated protein phosphatases are present, (3) flickering illumination and repeated depolarizations alter CaMKII autophosphorylation levels, and (4) CaMKII autophosphorylation and dephosphorylation alter spike propagation.

Spike-induced slowing of spike propagation

Illumination elicits as many as a few hundred spikes per second (Hz) in rat RGCs *in situ* (Brown and Rojas, 1965; Heine and Passaglia, 2011). To test whether repetitive spiking alters conduction velocity in RGC axons, we electrically stimulated excised optic nerves with pairs of current pulses separated by intervals ranging between 3 and 30 ms. In response to a pair of pulses separated by 20 ms or more, the time-to-peak values of the dif-

ferent peaks (P1, P2, and P3) of the CAP elicited by the second pulse did not significantly differ from those elicited by the first pulse (Fig. 1A). Therefore, in response to a pair of pulses at an instantaneous stimulus frequency of 50 Hz or less, single cardinal CAPs did not alter the time-to-peak of single follower CAPs. By contrast, the time-to-peak of P1, P2, and P3 elicited by the second pulse increased monotonically as the stimulus intervals decreased from 20 to 3 ms (Fig. 1A, Table 1). The time-to-peak increases of P1, P2, and P3 all reached $\sim 50\%$ of the increases at 3 ms at stimulus intervals of ~ 5 ms and the difference between the time-to-peak values at 3 and 20 ms were statistically significant (paired t test) for P1 (0.08 ± 0.01 ms, $n = 15$, range 0.04–0.15 ms, p -value = 5×10^{-7}), P2 (0.37 ± 0.03 ms, $n = 15$, range 0.20–0.65 ms, p -value = 1×10^{-8}), and P3 (0.60 ± 0.07 ms, $n = 11$, range 0.23–1.01 ms, p -value = 1×10^{-5}).

Because Figure 1A and Table 1 show that the time-to-peak of a follower spike is not increased if a cardinal spike precedes it by 20 ms, one might not expect conditioning spikes at 20-ms intervals to alter the time-to-peak of test spikes that follow them. We tested this by eliciting a pair of CAPs at a 5-ms interval preceded by a variable number of CAPs at 20-ms intervals and found that

Table 1. Change in time-to-peak at different ISIs

Peak no.	ISI (ms)	C–F time-to-peak difference (ms)	Range (ms)
P1 (<i>n</i> = 15)	3	0.08 ± 0.01	0.04–0.15
	4	0.06 ± 0.01	0.02–0.10
	5	0.03 ± 0.01	0.00–0.08
	8	0.01 ± 0.01	–0.01 to 0.02
	10	0.00 ± 0.01	–0.01 to 0.02
	20	0.00 ± 0.01	–0.01 to 0.00
P2 (<i>n</i> = 15)	3	0.37 ± 0.03	0.21–0.66
	4	0.23 ± 0.03	0.12–0.44
	5	0.16 ± 0.02	0.07–0.31
	8	0.07 ± 0.01	0.01–0.13
	10	0.04 ± 0.01	–0.02 to 0.08
	20	0.00 ± 0.01	–0.07 to 0.02
P3 (<i>n</i> = 11)	3	0.62 ± 0.07	0.27–1.02
	4	0.40 ± 0.04	0.17–0.65
	5	0.27 ± 0.03	0.15–0.47
	8	0.15 ± 0.02	0.07–0.32
	10	0.09 ± 0.01	0.03–0.15
	20	0.02 ± 0.01	–0.05 to 0.11

Stimulus pulses were presented as pairs, the first and second eliciting a cardinal (C) and follower (F) CAP, respectively. The C–F difference for each ISI is the mean ± SEM of the differences between the time-to-peak of the P1, P2, and P3 peaks in the C and F CAPs recorded from each of the indicated number of optic nerves.

the P1, P2, and P3 time-to-peak values of the test CAP increased (Fig. 1*B*). Increasing the number of conditioning CAPs produced further increases (Fig. 1*B*). Therefore, our recordings show a slowing effect that increases gradually and accumulates. Moreover, the slowing effect appears to asymptote after 50–300 ms of conditioning stimulation; that is, increasing the number of conditioning CAPs from 3 to 16 produced smaller increases in the slowing effect than increasing the conditioning CAP number from 1 to 3 (Fig. 1*B*). Repeated CAPs at lower frequencies (e.g., 10–20 Hz) did not induce slowing of test CAPs even if relatively large numbers of conditioning pulses were used. Also, we did not find that conditioning pulses decrease the time-to-peak of follower CAPs. This is of interest because studies of other systems have reported that spikes increase the conduction velocity of some axons during transient “supernormal” phases (Swadlow et al., 1980; Taylor et al., 1992) and that spikes reduce conduction velocity without supernormal increases in other preparations (Lüscher et al., 1994; George et al., 2007; Meeks and Mennerick, 2007).

Change in interspike interval and instantaneous spike frequency

Because the distance traveled by cardinal and follower spikes are the same, spike-induced increases in time-to-peak are consistent with the possibility that follower spikes propagate more slowly than cardinal spikes (Fig. 1). Due to this slowing, the interspike interval between cardinal and follower spikes at the recording electrode in single axons would be longer than the ISI at the stimulating electrode.

The changes in time-to-peak shown in Figure 1 were measured in segments of optic nerve that averaged 8 mm in length. In adult rat, the optic nerve length from the rear of the eye to the optic chiasm is ~10 mm (Matheson, 1970) and the optic tract length from the chiasm to the lateral geniculate nucleus (LGN) is 9–10 mm (Sumitomo et al., 1969). To estimate how much the interspike interval between cardinal and follower spikes might increase between the eye and a subcortical brain target, we assumed a distance of 20 mm from eye to LGN, that follower spike delays increase linearly with distance, and that adding the delays

and stimulus intervals yields the cardinal–follower spike intervals at the LGN. Inverting these sums to express the changes in units of instantaneous spike frequency (Hz) yields values of 313, 255, and 220 Hz for P1, P2, and P3, respectively, at a stimulus interval of 3 ms (333 Hz); 197, 185, and 176 Hz at a stimulus interval of 5 ms (200 Hz), and 100, 99, and 98 Hz at an interval of 10 ms (100 Hz). This suggests that optic tract spike frequencies will increase as stimulus frequencies increase >100 Hz, although not as much as RGC spike frequencies, and that stimulus frequencies and optic tract spike frequencies will not noticeably differ at <100 Hz. These results would not change qualitatively if the distance between the eye and target were a few millimeters longer or shorter than 20 mm. However, if the distance is a few-fold different (e.g., 100 mm, as in human; Lang and Reiter, 1985; Bernstein et al., 2016), then spike slowing could depress the instantaneous spike frequency at the shortest stimulus intervals more than at intermediate stimulus frequencies.

Beside slowing of propagation, spiking might appear to increase interspike intervals for two other reasons. One is that a cardinal spike might increase the “rise time” between the beginning of the depolarizing phase of a follower spike and the moment the follower spike reaches its peak (Meeks and Mennerick, 2007). Although we could not directly test this possibility on single axons in optic nerve, we could elicit and record axonal spikes on multielectrode arrays. When pairs of pulses at one electrode elicited pairs of spikes at a recording electrode a few hundred micrometers away, the latencies from the stimulus to the beginning of the two spikes elicited were indistinguishable. We also detected no difference in the rise times of cardinal and follower spikes in the recordings from some of these electrode pairs. However, on other electrode pairs, the follower spike rise times increased at high pulse frequencies. For example, at an ISI of 5 ms, the rise time of the follower spikes was as much as 0.28 ms longer than that of the cardinal spikes. Because we could not measure the length of axons that these measurements were made from, we could not identify which recordings corresponded to P1, P2, and P3 of the CAPs. Nonetheless, the changes that we measured (0–0.28 ms) are smaller than the increases in time-to-peak values of P2 and P3 in the CAPs (0.07–0.31 ms and 0.15–0.47 ms, respectively; Table 1) and as small as the increases found in P1 (0.00–0.08 ms, Table 1).

A second possible reason is that, among the axons that contribute to a single CAP peak, the firing of more rapidly conducting axons might be suppressed by the first stimulus and the firing of more slowly conducting axons might either be spared or further slowed. This seems unlikely because recordings of isolated CAP peaks in optic tract (by recording from positions populated by axons of relatively uniform diameters) have shown that spike-induced changes in time-to-peak do not split peaks into more rapidly and slowly propagating subgroups (Bishop et al., 1953). However, we cannot yet exclude this possibility for optic nerve. We used various fine-tipped metal and glass electrodes to record from optic nerve, but did not capture isolated peaks. This is presumably because axons of different diameter are not segregated into different layers or sectors within the rat optic nerve (Figs. 2, 3; see also Sugimoto et al., 1984; Reese, 1987).

Autophosphorylated CaMKII in rat optic nerve

Previous reports of voltage-gated Ca²⁺ channels and CaMKII in rat optic nerve (Brown et al., 2001; Lund and McQuarrie, 2001) raise the possibility that pT286 might be present under certain conditions and thus positioned to modulate the excitability of RGC axons. We began to test this possibility by probing Western

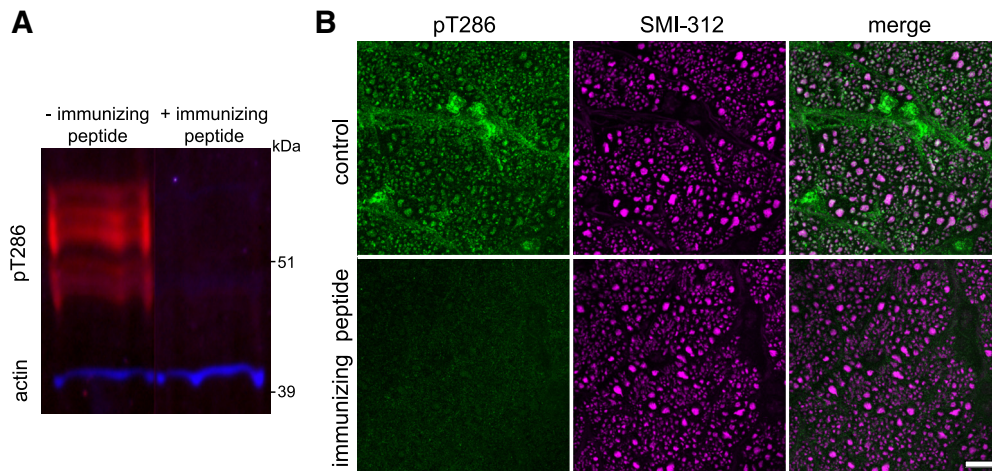


Figure 2. pT286-immunopositive optic nerve protein bands and axons, and reduction of pT286 staining intensity by immunizing peptide. **A**, Western blots of a rat optic nerve, processed side-by-side in adjacent lanes. Left lane, “– immunizing peptide” probed with anti-pT286 (red) and anti- β -actin (blue) antibodies. Right lane, “+ immunizing peptide” probed with anti- β -actin antibody and anti-pT286 antibody that was preincubated with immunizing peptide. The latter shows no detectable pT286 signal above background intensity. Molecular weight and migration distance of protein standards that were run in an adjacent lane are shown at right edge of **A**. **B**, Transverse sections of an optic nerve processed side-by-side and imaged at identical laser intensity, photomultiplier gain, and pinhole diameter. Left, middle, and right panels of top row show pT286 (green), SMI-312 (magenta), and merge thereof, respectively. pT286 detected in axons (round, SMI-312-immunopositive profiles) and in baffle-like structure filling dark, SMI-312-immunonegative region between clusters of SMI-312-immunopositive profiles. Left, middle, and right panels of bottom row show green only (anti-pT286 antibody that was preincubated with immunizing peptide), magenta only (SMI-312), and merge thereof, respectively. This row shows no detectable pT286 signal above background intensity. Scale bar at lower right in **B** indicates 10 μ m and applies to all panels.

blots of optic nerve homogenates with an anti-pT286 antibody and by immunostaining transverse and longitudinal sections of optic nerve for pT286. Both the rabbit polyclonal and the rabbit monoclonal anti-pT286 antibodies used bound to three protein bands in these blots, with apparent molecular weights of \sim 49, 53, and 55 kDa (Fig. 2A). These values agree well with those found in other preparations (Matsuyama et al., 2014; Sheng et al., 2015). No staining above background was detectable in Western blots with primary antibody that was preincubated with an excess of immunizing peptide (Fig. 2A). Opposite staining patterns were obtained when adjacent lanes were stained with blocked and unblocked primary antibody, stripped of primary and secondary antibodies with a stripping buffer, and then restained with the antibodies switched (e.g., blocked antibody applied to the lane that had been bound by unblocked antibody before stripping). Moreover, antibodies directed against the α isoform (CaMKII- α) bound to the 49 and 53 kDa bands in our blots, as found in blots prepared from retina, hippocampus, and cerebellum (Ochiishi et al., 1994; Laabich and Cooper, 1999; Liu et al., 2000; Menegon et al., 2002). In blots stained with some anti-CaMKII antibodies, the immunofluorescence intensity of the 49 kDa band exceeded that of the other bands (Fig. 4A).

Within the optic nerve, an anti-pT286 antibody (rabbit polyclonal directed against an amino acid sequence that surrounds and includes the pT286 of CaMKII- α) bound to round, elongated profiles (Figs. 2, 3). These profiles were identified as axons because they also bound antibody directed against pan axonal neurofilaments (SMI-312; Ulfing et al., 1998). All, or very nearly all, of the SMI-312-immunopositive axons were pT286-immunopositive and we did not notice a size, number, or distribution of axons that consistently lacked pT286 immunopositivity. The pT286- and SMI-312-immunopositive profiles ranged in diameter from \sim 0.3 to 4.3 μ m in both transverse (Fig. 3H) and longitudinal (Fig. 3A) sections. This diameter range agrees with previously reported dimensions of rat optic nerve axons (Forrester and Peters, 1967; De Juan et al., 1978; Foster et al., 1982; Sugimoto et al., 1984; Reese, 1987). The mean and median values

of the diameters were 1.0 and 0.8 μ m, respectively ($n = 390$ axon profiles from a total of eight optic nerves). No immunostaining above background was detectable with primary antibody that was preincubated with an excess of immunizing peptide (Fig. 2B). Moreover, a different antibody (rabbit monoclonal, directed against an amino acid sequence that surrounds and includes the phosphorylated T287 of CaMKII- β) produced similar staining of SMI-312-immunopositive profiles in longitudinally and transversely sectioned preparations.

In preparations triple-labeled for pT286, SMI-312, and MBP (Dixon and Eng, 1984), pT286 was found in SMI-312-immunopositive profiles and not in MBP-immunopositive profiles (Fig. 3). Moreover, cylindrical, pT286-immunopositive profiles were circumscribed by annular, MBP-immunopositive profiles (Fig. 3H). The MBP-immunopositive annuli that we could measure in our confocal images ranged in outer diameters between 1.2 and 4.9 μ m (mean \pm SEM = $2.6 \pm 0.1 \mu$ m; $n = 164$). These values are intermediate between light and electron microscopically measured values reported previously for rat (Forrester and Peters, 1967; Matheson, 1970). The ratio of the diameter of each pT286-immunopositive profile to the outer diameter of the MBP-immunopositive profiles that circumscribed it (g-ratio; Fig. 3P) ranged between 0.4 and 0.8, and averaged 0.7 ± 0.1 (mean \pm SEM, $n = 164$). Our images also suggested that smaller-diameter MBP-immunopositive annuli were present (arrowheads in Fig. 3H). However, the outer and inner diameters of these were too small for us to measure. We also could not resolve whether any fibers lacked MBP-immunopositive annular wrappings, especially small fibers that might correspond to the unmyelinated fibers reported by other studies (Foster et al., 1982; Sugimoto et al., 1984).

Net reduction of pT286 levels by flickering illumination and by spiking

Based on changes in the intensity and location of arrestin, transducin, and cAMP immunostaining in photoreceptors and of cAMP in retinal ganglion cells, we previously found that flicker-

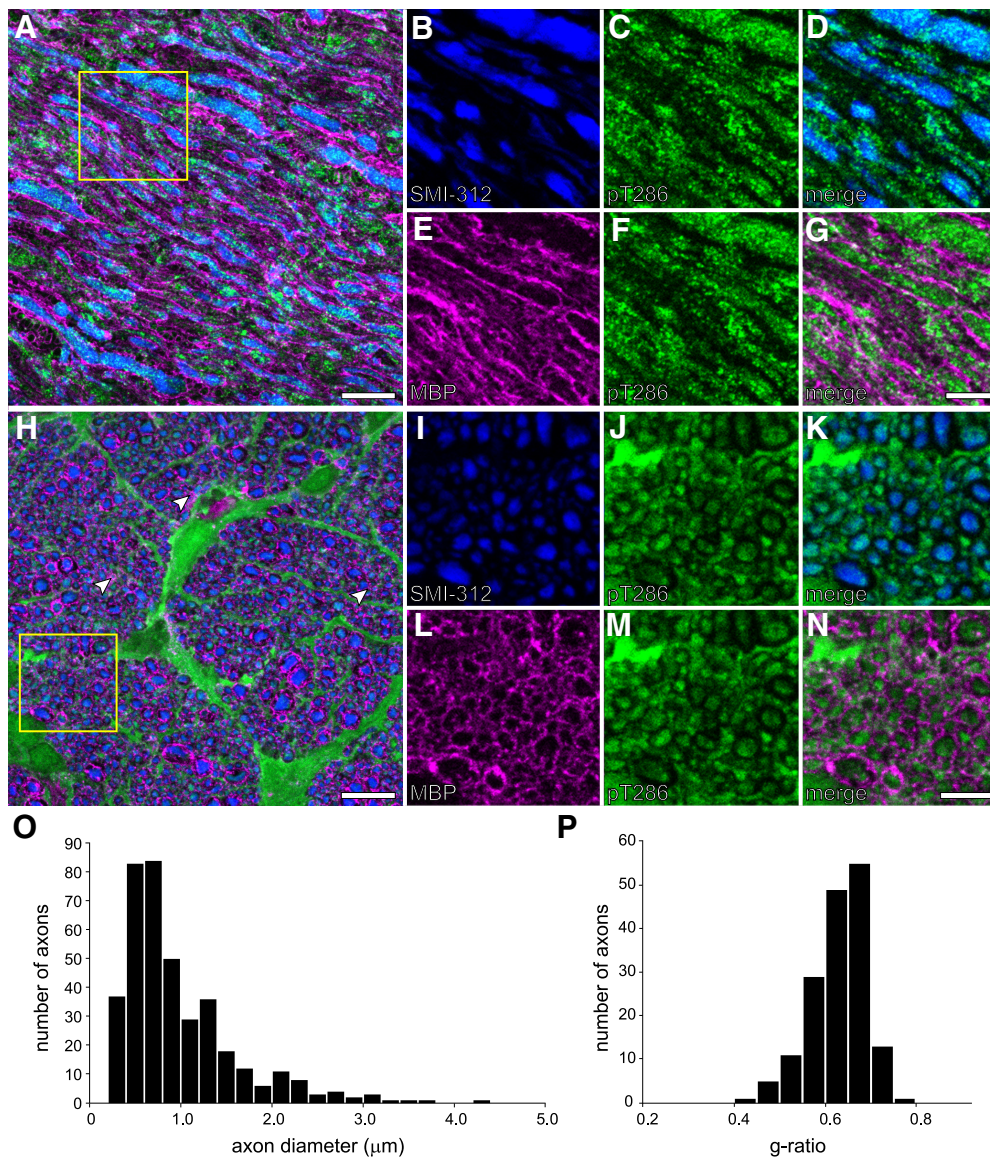


Figure 3. Diameters and myelin of pT286-immunopositive optic nerve axons. Longitudinal (**A**) and transverse (**H**) sections triple-labeled for SMI-312 (blue), pT286 (green), and MBP (magenta). **B–D**, SMI-312 (**B**) and pT286 (**C**) of the area within the yellow square in **A** and merge thereof (**D**). SMI-312 and pT286 colocalization generates cyan. **E–G**, MBP (**E**), pT286 (**F**), and merge thereof (**G**) of the same field as **D**. Separation of magenta and green in **G** shows that pT286 localizes to axons and not to myelin. The axon diameters (e.g., **A** and **G**) range between small and large. **H–N**, Comparable information in cross-section, including colocalization (cyan in **K**) of SMI-312 (**I**) and pT286 (**J**), separation (**N**) of MBP (**L**) and pT286 (**M**), and wide range of axon diameters (**N**) and inner diameter of myelin sheaths (**L**). **I–N**, Field outlined by yellow square in **H**. Scale bars: **A**, **H**, 10 μm ; **G**, 5 μm (applies to **B–G**); **N**, 5 μm (applies to **I–N**). Histograms of axon diameters (**O**) and g-ratio (**P**; ratio of inner and outer diameters of MBP-immunopositive profiles) of optic nerves of which **A** and **H** are part.

ing illumination light adapts adult rat retinae and increases pT286 immunostaining intensity in rat RGC somata (Ogata et al., 2012). Because CaMKII can be detected in optic nerve (Lund and McQuarrie, 2001; Calkins et al., 2005), electrically stimulated spiking elevates free Ca^{2+} levels in optic nerve axons (Lev-Ram and Grinvald, 1987; Verbny et al., 2002) and alternating increments and decrements of illumination increases spike firing in RGCs (Hartline, 1938; Baden et al., 2016), we anticipated that flickering illumination and electrical stimulation would increase pT286 immunostaining intensity in the optic nerve.

Contrary to these expectations, flickering illumination (for details, see Materials and Methods) lowered the staining intensity of the pT286-immunopositive protein bands in Western blots (Fig. 4). Because illumination reduced the intensity of all three pT286-immunopositive bands (~ 49 , 53, and 55 kDa; Fig. 4A, B),

the scatter plot and bar graph in Figure 4, C and D, compare the sum of the intensities of the bands in the dark versus light. The fluorescence intensities of these bands normalized to that of loading control protein in each lane were $20 \pm 7\%$ higher (mean \pm SEM, $n = 9$) in the blots prepared from dark-adapted rat optic nerves than in those of concomitantly collected and processed light-adapted rat optic nerves. The mean intensities differed significantly (Fig. 4D, Wilcoxon signed-rank test, $p = 0.04$) and there was no statistically significant light/dark staining intensity difference between the CaMKII-immunopositive bands in the same blots (Wilcoxon signed-rank test, $p = 0.78$; Fig. 4D).

Similarly, the average of the fluorescence intensities of the pT286-immunopositive protein bands in a given blot lane normalized to that of the loading control protein was $30 \pm 5\%$ lower (mean \pm SEM, $n = 10$) in the blots prepared from electrically

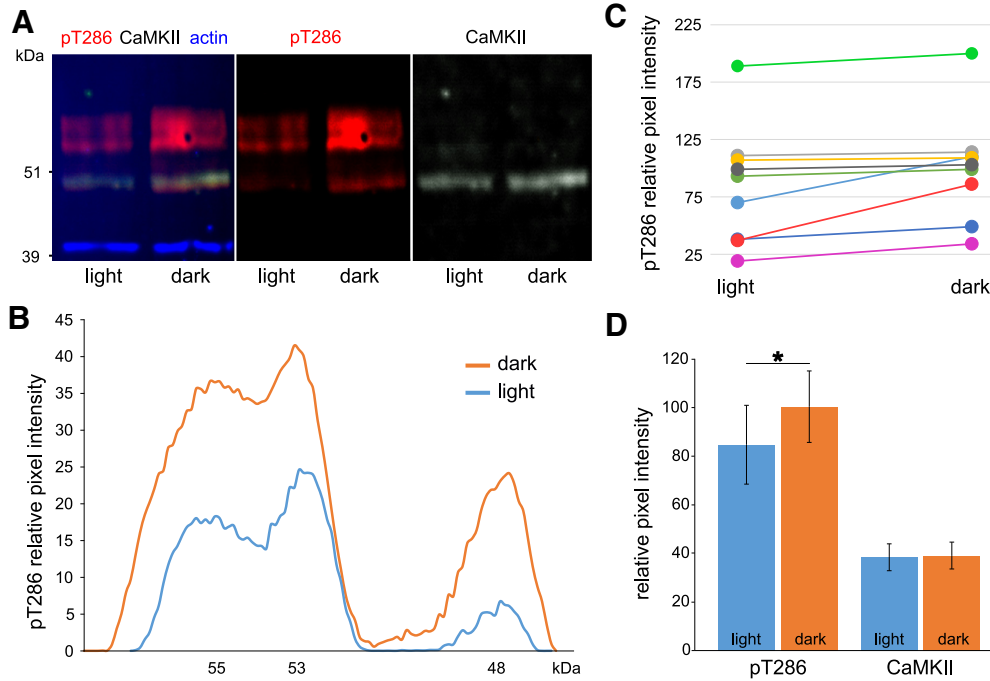


Figure 4. Illumination reduces optic nerve pT286 signal intensity. **A**, Western blot prepared from optic nerves of dark- and light-adapted rats collected, processed, and probed in pairs with antibodies directed against pT286 (red), CaMKII (gray), and β -actin (blue). The left, middle, and right pairs of lanes show all signals superimposed (red, gray, and blue color channels), pT286 signals only (red), and CaMKII signals only (gray), respectively. Molecular weight and migration distance of protein standards run in adjacent lane shown at left edge of **A**. **B**, pT286 pixel intensity in **A** normalized by β -actin band pixel intensity and plotted against estimated apparent molecular weight. The two tallest peaks correspond to the two bright bands >50 kDa in **A**. **C**, pT286 pixel intensity in Western blots of optic nerves collected from pairs ($n = 9$) of light- and dark-adapted rats. The average of the fluorescence intensities of the pT286 bands in each blot was normalized by the fluorescence intensity of the β -actin band in each lane. Values from each rat pair are plotted by dots of same color and connected by lines. **D**, Mean signal intensities (\pm SEM) of pT286 and CaMKII bands for the nerves as in **C**. pT286 signal variance is relatively large because light decreased the signal intensity in some preparations more than in others (**B**). However, the difference between means is statistically significant (*, Wilcoxon signed-rank test, $p = 0.04$). In **A–D**, “light” and “dark” denote light- and dark-adapted, respectively.

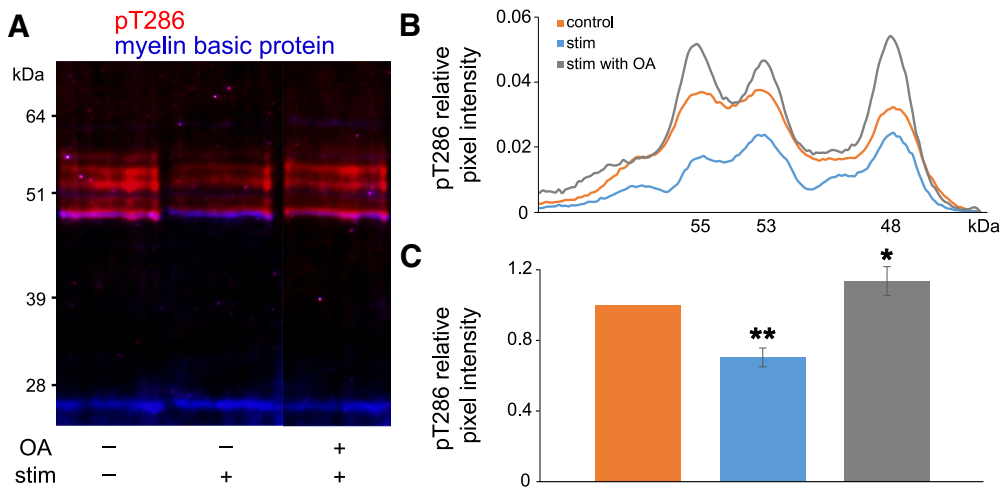


Figure 5. Okadaic acid increases optic nerve pT286 signal intensity. **A**, Western blots prepared from optic nerves that were electrically stimulated or not in the presence and absence of okadaic acid (as indicated by the lane labels below the blot lanes). Blots were probed with antibodies directed against pT286 (red) and MBP (blue). Molecular weight and migration distance of protein standards run in adjacent lane shown at left edge. **B**, Relative pixel intensity of pT286 signals in **A** normalized by pixel intensity of MBP bands and plotted against estimated apparent molecular weight. **C**, Relative pixel intensity of pT286 signals from $n = 3$ optic nerves processed and analyzed as in **A** and **B**. Differences between fluorescence intensity of protein bands from control and electrically stimulated optic nerves (***) and between those of control nerves and nerves stimulated in the presence of okadaic acid (*) are statistically significant ($p = 0.002$ and 0.025 , respectively).

stimulated optic nerves (for details, see Materials and Methods) than in those of concomitantly processed optic nerves that were not stimulated (Fig. 5). The unstimulated nerves were positioned between stimulating and recording electrodes and maintained under identical conditions except that electrical stimuli were not delivered. The difference between these fluorescence intensities

was statistically significant (Wilcoxon signed-rank test, $p = 0.002$; Fig. 5C).

The T286 of CaMKII is autophosphorylated during activation by Ca^{2+} /calmodulin complexes (Lisman et al., 2002) and can be dephosphorylated by protein serine/threonine phosphatases (Ishida et al., 2003). Therefore, one possible explanation for the

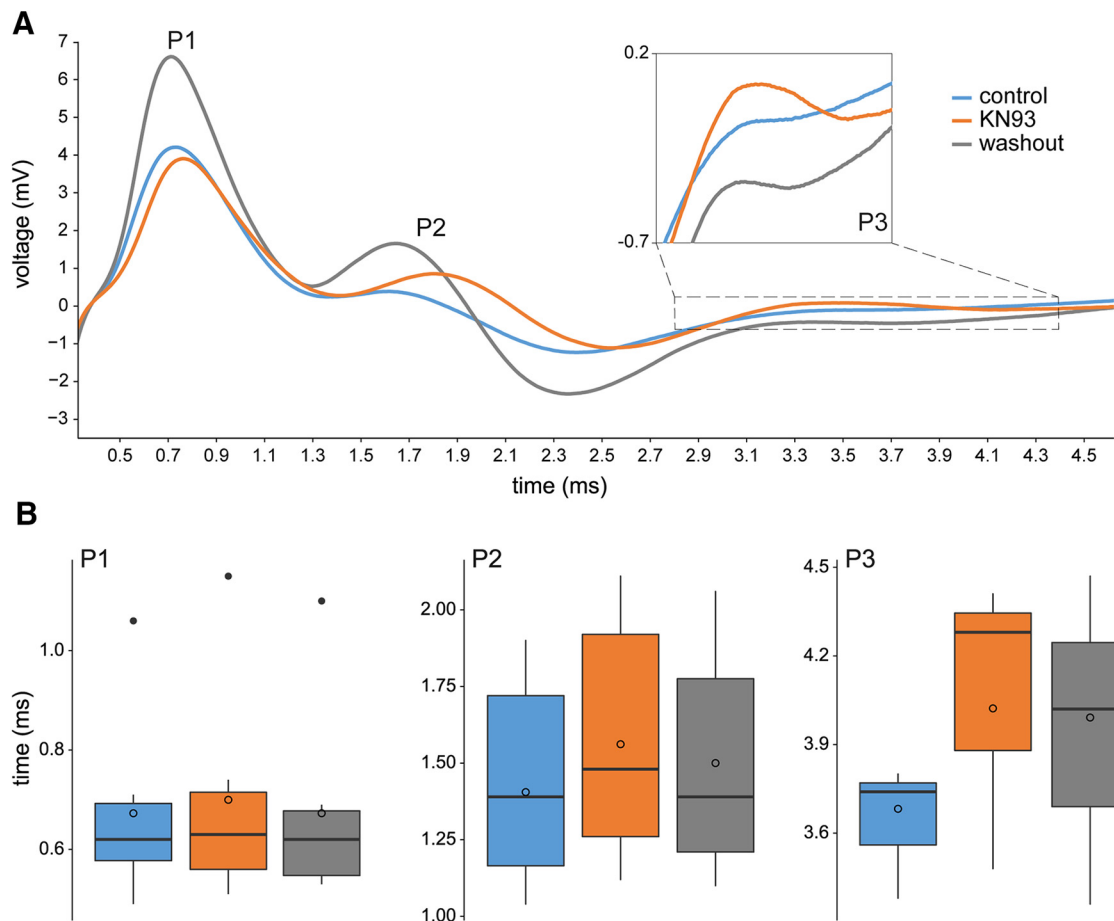


Figure 6. KN93 reversibly increases time-to-peak of optic nerve spikes. **A**, CAP recorded before (blue) and during (orange) inclusion of 2 μ M KN93 in medium superfusing an optic nerve and after washing with control medium (gray). Inset shows magnified view of P3. **B**, Box plots of the effect of KN93 on time-to-peak and its reversibility (same color coding as in **A**). These show the median (thick horizontal line), mean (circle), interquartile range (IQR, box), largest and smallest values no further than 1.5 \times IQR from the box (vertical lines) and outliers (single dots). The difference between the P2 time-to-peak in control and KN93 is statistically significant ($p = 0.015$).

illumination- and spiking-induced decrease in pT286 is that spiking activates a protein phosphatase that dephosphorylates the basal level of pT286. A second possibility is that spiking decreases CaMKII autophosphorylation and that a constitutively active protein phosphatase dephosphorylates the basal level of pT286. A third possibility is that spiking increases both CaMKII autophosphorylation and a protein phosphatase activity that dephosphorylates pT286 and that the net effect that can be detected by Western blots is a decrease in pT286 levels. Of these possibilities, the pT286 immunostaining intensities of Western blots normalized by the loading control protein staining intensities (Fig. 5A) showed effects that are consistent with the third. The simplest interpretations of the electrically stimulated decrease in pT286 (Fig. 5A–C) are that constitutive phosphorylation helps set the pT286 level in unstimulated optic nerve and that stimulation increases phosphatase activity. Moreover, the pT286 signal during electrical stimulation in the presence of okadaic acid (0.4 μ M) exceeds that in resting optic nerves by $17 \pm 7\%$ (mean \pm SEM, $n = 10$; Fig. 5A–C). The difference between these intensities was statistically significant (Wilcoxon signed-rank test, $p = 0.025$). This suggests that electrical stimulation increases CaMKII autophosphorylation in addition to increasing phosphatase activity and that dephosphorylation outweighs autophosphorylation in control medium.

Decreasing CaMKII autophosphorylation decreases the speed of spike propagation

Because repetitive stimulation slows spike propagation (Fig. 1) and lowers the pT286 fluorescence intensity in Western blots (Fig. 5), we next tested whether spike time-to-peak is affected by a water-soluble inhibitor of Ca^{2+} /calmodulin binding to CaMKII (KN93; Sumi et al., 1991). We compared CAPs recorded from optic nerves superfused with control medium, control medium supplemented with KN93 (2 μ M), and control medium again, each for 20–60 min. KN93 increased the P1, P2, and P3 time-to-peak values of CAPs activated by single pulses to $103 \pm 2\%$, $109 \pm 1\%$, and $111 \pm 4\%$ of the control values, respectively (mean \pm SEM of the changes measured nerve-by-nerve; $n = 6$, $n = 7$, and $n = 3$, respectively; Fig. 6). Washing with control medium decreased the P1, P2, and P3 values of the same nerves to $100 \pm 2\%$, $104 \pm 1\%$, and $108 \pm 6\%$ of the control values, respectively. Box plots (Fig. 6B) show that the median, mean, and majority of the time-to-peak values increased in KN93 and recovered at least partially during subsequent washing. The difference between the time-to-peak values (pooled across P1, P2, and P3) was statistically significant between control and KN93 ($p = 0.00036$) and between KN93 and wash ($p = 0.0065$, both Wilcoxon signed-rank test, $n = 16$). The difference between the measurements in control and KN93 was statistically significant for P2

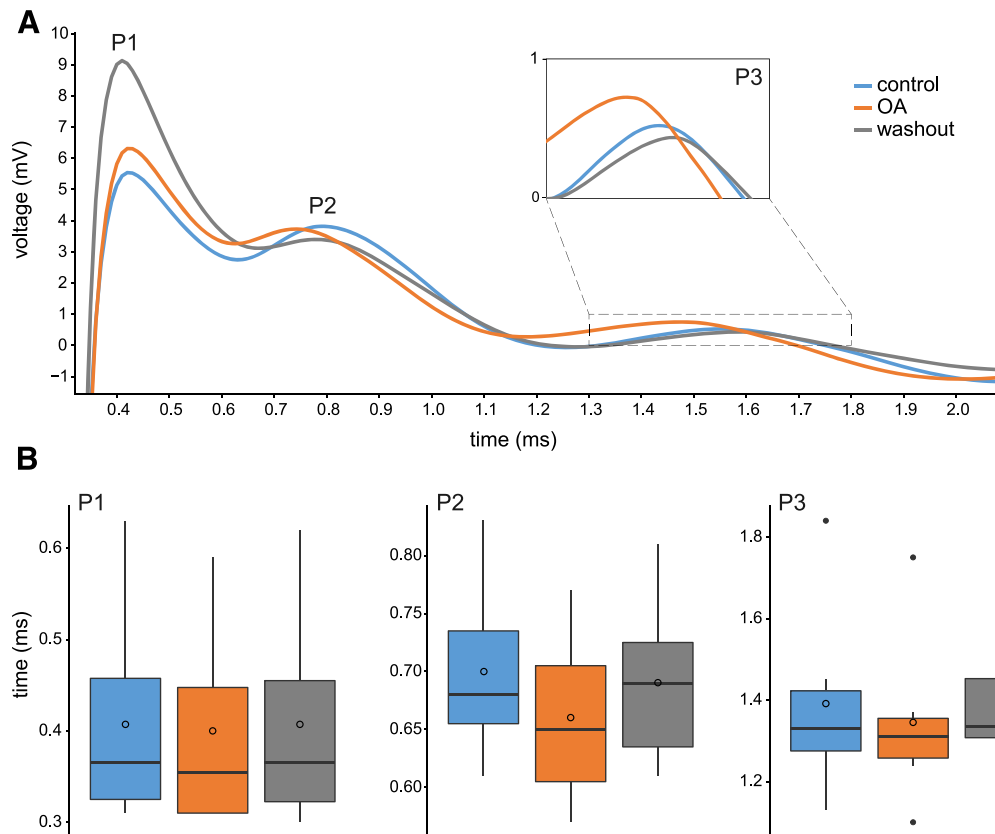


Figure 7. Okadaic acid reversibly decreases time-to-peak of optic nerve spikes. **A**, CAP recorded before (blue) and during (orange) inclusion of $0.4 \mu\text{M}$ okadaic acid in medium superfusing an optic nerve and after washing with control medium (gray). Inset shows magnified view of P3. **B**, Box plots of the effect of okadaic acid on time-to-peak and its reversibility in same color-coded conditions as in **A**. These show the median (thick horizontal line), mean (circle), interquartile range (IQR, box), largest and smallest values no further than $1.5 \times \text{IQR}$ from the box (vertical lines), and outliers (single dots).

(Wilcoxon signed-rank test, $p = 0.015$) and KN93 slowed P1 slightly and slowed P3 more than P2 or P1. However, the differences for P1 and P3 did not reach statistical significance due to small sample sizes.

Some previous studies used KN93 solubilized in dimethylsulfoxide (DMSO) to inhibit CaMKII. We found that DMSO-solubilized KN93 produced inconsistent (and, in some nerves, irreversible) changes in spike interval (result not illustrated). This is consistent with reports that DMSO-solubilized KN93 can produce nonspecific effects (Ledoux et al., 1999; Gao et al., 2006) and that DMSO is not an inert vehicle (Tamagnini et al., 2014). Based on these results, we did not test for the effects of other DMSO-solubilized reagents (e.g., KN92 and KN62).

Increasing CaMKII autophosphorylation increases the speed of spike propagation

We next tested whether increasing CaMKII autophosphorylation speeds up spike propagation. Given that okadaic acid and fostriecin inhibit protein phosphatase isoforms that dephosphorylate pT286 in various preparations (Ishida et al., 2003), CAPs were recorded from optic nerves superfused with control medium, control medium supplemented with either okadaic acid or fostriecin, and then control medium again, each for 20–60 min.

Okadaic acid ($0.4 \mu\text{M}$) reduced the P1, P2, and P3 time-to-peak values in response to single pulses to $97 \pm 1\%$, $93 \pm 2\%$, and $97 \pm 1\%$ of the control values (mean of the changes measured nerve-by-nerve; $n = 4$, $n = 5$, and $n = 4$, respectively; Fig. 7). Washing with control medium increased the mean P1, P2, and P3

values of the same nerves to $100 \pm 1\%$, $99 \pm 2\%$, and $103 \pm 6\%$ of the control values, respectively. Similarly, fostriecin ($1 \mu\text{M}$) decreased the time-to-peak of P1, P2, and P3 to $96 \pm 1\%$, $94 \pm 2\%$, and $92 \pm 3\%$ of the control values, respectively, and washing increased these values to $102 \pm 2\%$, $103 \pm 1\%$, and $99 \pm 5\%$ of the control values, respectively (mean, $n = 6$, $n = 6$, and $n = 5$, respectively; Fig. 8). Box plots show that the median, mean, and range of the time-to-peak values shifted to lower values in okadaic acid (Fig. 7B) and in fostriecin (Fig. 8B) and recovered during subsequent washes. Wilcoxon signed-rank tests indicated that the decreases in time-to-peak of P2 and P3 by okadaic acid and the decrease in time-to-peak of P2 by fostriecin were statistically significant (Wilcoxon signed-rank test, p -values of 0.0223, 0.0313, and 0.0355, respectively). Although fostriecin decreased the time-to-peak of P1 and of P3, the sample sizes did not allow calculations of Wilcoxon signed-rank statistics for those peaks. Also, the latency of P1 was so short (typically 0.5 ms) that it would have been difficult to detect changes similar in magnitude (5–10%) to those we found in P2 and P3.

To test whether a phosphatase inhibitor counteracts activity-induced slowing of spikes, we measured the time-to-peak of cardinal CAPs and of F3 CAPs (i.e., those elicited after three conditioning pulses; cf. Fig. 1B) before and during the application of fostriecin ($1 \mu\text{M}$) and after washing with control medium (Fig. 9A,B). These CAPs were elicited by trains of 4–5 pulses presented at an ISI of 5 ms with a 1-s pause between trains. Each cardinal CAP and F3 CAP was thus preceded by a 1-s and 5-ms stimulus interval, respectively. We calculated slowing as the dif-

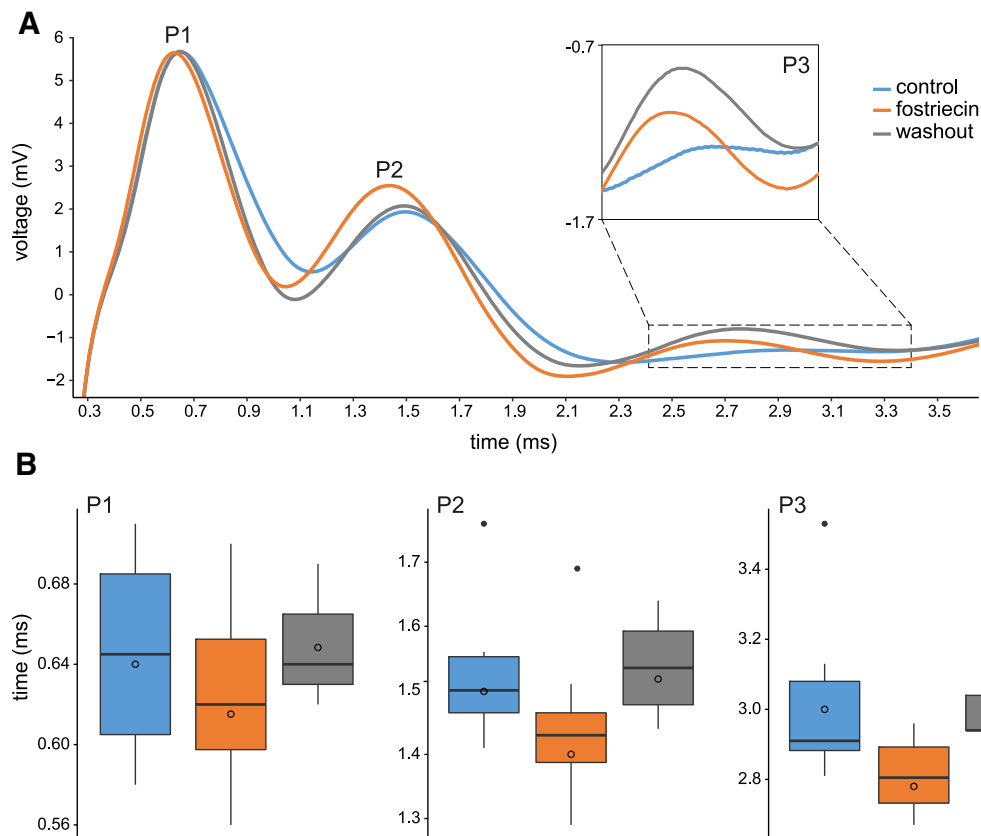


Figure 8. Fostriecin reversibly decreases time-to-peak of optic nerve spikes. **A**, CAP recorded before (blue) and during (orange) inclusion of 1 μ M fostriecin in medium superfusing an optic nerve and after washing with control medium (gray). Inset shows magnified view of P3. **B**, Box plots of the effect of fostriecin on time-to-peak and its reversibility in same color-coded conditions as in **A**. These show the median (thick horizontal line), mean (circle), interquartile range (IQR, box), largest and smallest values no further than 1.5 * IQR from the box (vertical lines), and outliers (single dots).

ference between the time-to-peak of the cardinal and F3 CAPs (Fig. 9C). Fostriecin reversibly reduced this difference for P1, P2, and P3. Fostriecin reduced the slowing to $91 \pm 15\%$, $73 \pm 19\%$, and $71 \pm 15\%$ of the control value, respectively. These reductions were statistically significant for P2 and P3 (Wilcoxon signed-rank tests, $p = 0.016$ and $p = 0.031$ and $n = 8$ and $n = 6$, respectively), but did not reach significance for P1 ($p = 0.477$, $n = 8$). After washing, the slowing of P1, P2, and P3 were $107 \pm 5\%$, $106 \pm 3\%$, and $112 \pm 2\%$ of the control values, respectively. The differences between the values measured in fostriecin and wash were statistically significant (Wilcoxon signed-rank tests, $p = 0.031$, $p = 0.008$, and $p = 0.031$, and $n = 8$, $n = 8$, and $n = 6$, respectively). As observed with okadaic acid (Fig. 5), the pT286 immunostaining intensity of Western blots prepared from optic nerves electrically stimulated in the presence of fostriecin (1 μ M) were greater than those of optic nerves that were electrically stimulated without fostriecin ($n = 2$; results not illustrated).

Protein phosphatases in optic nerve

Okadaic acid and fostriecin are commonly used to inhibit the protein phosphatase isoforms 1 and 2A (Bialojan and Takai, 1988; Walsh et al., 1997). We therefore attempted to localize these isoforms in transverse and longitudinal optic nerve sections by immunostaining with anti-PP1 and anti-PP2A antibodies. Preparations that were double-labeled for PP1 and MBP showed that optic nerve myelin bound anti-PP1 antibodies and that detectable levels of PP1 immunopositivity were not found in optic nerve axons (Fig. 10). This result is consistent with a previous report of PP1 in optic nerve homogenates (da Cruz e Silva et al.,

1995). By contrast, preparations that were triple-labeled for PP2A, MBP, and SMI-312 showed that axons bound anti-PP2A antibodies and that myelin did not (Fig. 11). Like the pT286-immunopositive axonal profiles shown in Figure 3, the PP2A-immunopositive axonal profiles ranged in diameter from small to large (Fig. 11). No previous study that we know of has detected PP2A in optic nerve axons.

Activity-induced spike slowing *in vivo*

Last, to test whether spike-induced spike slowing occurs in rat optic nerve *in vivo*, we used electrodes inserted into the optic chiasm to electrically stimulate spikes in ganglion cell axons and used a suction electrode to record these spikes from the optic nerve behind the eye. Stimuli were presented as a pair of pulses or as a train of 15 pulses with a fixed ISI separating the pulses. The pairs were repeated 13–14 times to form a series with a 2-s rest period between each pair. The trains were repeated the same way. For each ISI, 3 series of pairs were interleaved with 3 series of trains, resulting in 6 series (2, 15, 2, 15, 2, and 15) per preparation per ISI. This set of 6 series was first collected for the 5-ms ISI, then for 8-ms ISI, and then for 30-ms ISI. (The CAP durations we found *in vivo* precluded tests of shorter stimulus intervals.) These recordings showed that both single and multiple conditioning pulses increased the time-to-peak values of the CAP waveforms recorded in the optic nerve (Fig. 12). As found in our recordings from excised optic nerves (Fig. 1), the degree of propagation slowing increased as the ISI decreased from 30 to 5 ms (gray vs orange vs blue lines in Fig. 12C,D) and as the number of conditioning pulses increased from 1 to 15 (gray, orange, and blue lines

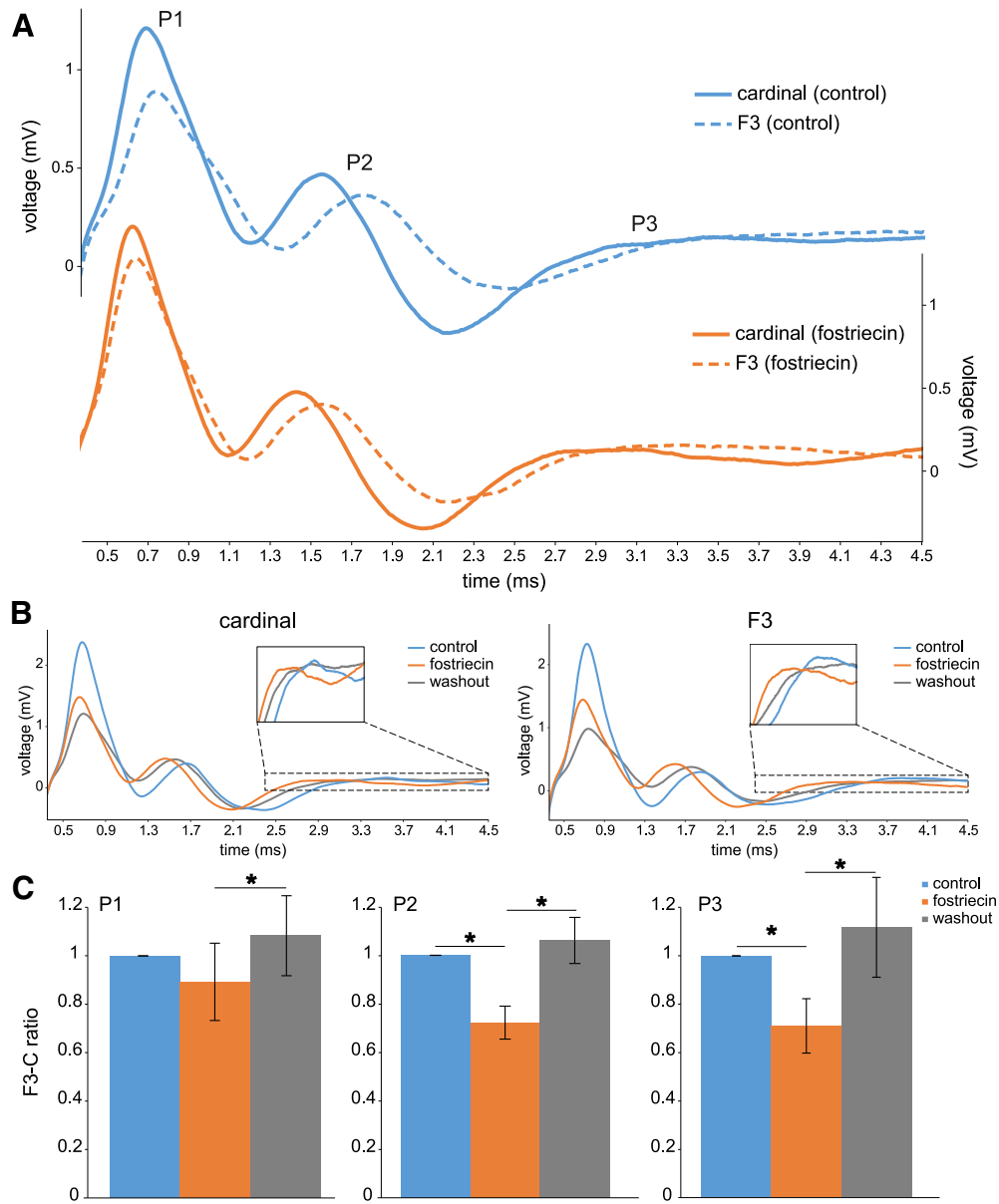


Figure 9. Fostriecin counteracts slowing by multiple conditioning pulses. **A**, CAPs elicited by the first (cardinal; C; continuous line) and fourth (F3; dotted line) of four pulses delivered at 200 Hz (5-ms ISI) in control (blue) and fostriecin (orange). **B**, Cardinal and F3 CAPs recorded before and during inclusion of 1 μ M fostriecin in medium superfusing the optic nerve and after washing with control medium. Digital average of $n = 3$ –5 CAPs recorded from one optic nerve in these media are plotted by blue, orange, and gray lines, respectively. **C**, Effect of fostriecin on the time-to-peak of the F3 CAPs normalized to those of the cardinal CAPs and its reversibility (same color coding as in **B**). The difference between control and fostriecin is statistically significant for P2 and P3 (*, Wilcoxon signed-rank, $p = 0.0156$ and $p = 0.0313$, respectively), but not for P1. The difference between fostriecin and the washes were significant for P1, P2, and P3 (*, Wilcoxon signed-rank; $p = 0.0313$, $p = 0.0078$, $p = 0.0313$, respectively).

in Fig. 12C,D). Also consistent with our *ex vivo* recordings, the increase in time-to-peak was larger for P2 than for P1 (Fig. 12D), the time-to-peak of both P1 and P2 was increased more by the first conditioning pulses than later in long trains of pulses (Fig. 12C,D), and none of the conditioning pulses decreased the time-to-peak values. Our data showed these trends regardless of whether the time-to-peak values were measured at the maximum amplitude of the raw waveform (dots in Fig. 12A,C) or at the maximum amplitude of Gaussian functions whose sums fitted the raw waveform (solid and dashed lines in Fig. 12B,D). These fits were used to estimate the time-to-peak values because distinct peaks were not well separated (cf. Fig. 1, 8) and, instead, the *in vivo* CAPs resembled the waveforms elicited by high stimulus intensities in previous studies (Evans

et al., 2010). By switching the recording and stimulating electrodes in one preparation, we also elicited CAPs in the optic nerve and recorded them in the optic chiasm. The time-to-peak values of these orthodromically stimulated CAPs also increased at short ISI values. However, these CAPs were smaller in amplitude than the antidromically stimulated CAPs described above, rendering the orthodromic propagation changes more difficult to measure than the antidromic propagation changes.

Discussion

Our results show that rat optic nerves can modulate the propagation of the spikes that they generate. Specifically, we found that conditioning spikes and changes in CaMKII autophosphoryla-

tion alter the propagation speed of subsequently elicited spikes. Below, we compare these results with CaMKII responses in other preparations and relate propagation speed changes to afferent signal transmission by RGCs.

Modulation of pT286 in optic nerve axons

Because flickering illumination increases pT286 levels in RGC somata (Ogata et al., 2012), we were initially surprised to find that illumination and electrical stimulation decrease optic nerve pT286 levels (Figs. 4, 5). These decreases could result from dephosphorylation occurring alone or from dephosphorylation that outweighs concomitant phosphorylation. Our observations are consistent with the latter in that protein phosphatase inhibition counteracts the decrease of pT286 by electrical stimulation and increases pT286 staining intensity to levels exceeding that in unstimulated optic nerve (Fig. 5). The stimulation-induced decrease in pT286 that we found differs from earlier findings that depolarization and excitation increase CaMKII autophosphorylation (Molloy and Kennedy, 1991; Ocorr and Shulman, 1991). However, excitation has been found to reduce CaMKII autophosphorylation in hippocampus and parietal cortex (Yamagata and Obata, 1998), low and high levels of neuronal activity have been found to favor CaMKII phosphorylation and dephosphorylation, respectively (Gu and Yan, 2004), and NMDA receptor activation by low and high agonist concentrations has been found to increase and decrease CaMKII phosphorylation, respectively (Zhou et al., 2012). Moreover, the differential pT286 changes that we have found in RGC somata (Ogata et al., 2012) and optic nerve axons (Fig. 4) resemble the finding that elevated extracellular KCl depolarizes somata and axons of embryonic rat hippocampal neurons, elevates the intracellular free Ca^{2+} concentration in both compartments, and increases the pT286 immunostaining intensity in only the somata (Menegon et al., 2002). The parallel with our results is underscored by the increases in axonal pT286 produced by depolarizing the same neurons in the presence of okadaic acid (Menegon et al., 2002).

Modulation of spike propagation by pT286

The pT286 immunopositivity in unstimulated optic nerve axons (Fig. 3) resembles the basal levels of autophosphorylated CaMKII found in other unstimulated preparations (Molloy and Kennedy, 1991; Ocorr and Schulman, 1991) and raises the possibility that increases and decreases in pT286 can influence functional properties of resting and stimulated optic nerves. Consistent with this, we found that okadaic acid and fostriecin increased pT286 levels, accelerated spike propagation, and decreased interspike interval, whereas electrical stimulation decreased pT286 levels, slowed spike propagation, and increased interspike interval. Our finding that illumination lowers pT286 levels in optic nerve axons (Fig. 4) at an intensity and flicker frequency that translocates photoreceptor arrestin (Ogata et al., 2012) identifies a novel mechanism by which light adaptation can lower RGC spike frequency (Hayashida et al., 2009).

How do single spikes and changes in pT286 slow follower spike propagation? The simplest possibility is that they modulate

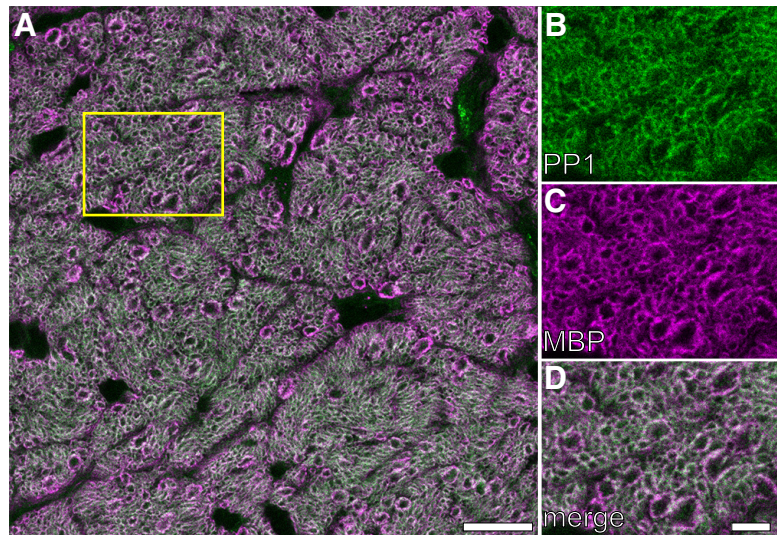


Figure 10. PP1 in optic nerve myelin. **A**, Transverse section double-labeled for PP1 (green) and MBP (magenta). **B–D**, PP1 (**B**), MBP (**C**), and merge thereof (**D**) of the area within the yellow square in **A**. PP1 and MBP colocalization generates white. Scale bars: **A**, 15 μ m; **D**, 5 μ m (applies to **B–D**).

one ion channel population that contributes to conduction velocity. Some studies have attributed propagation slowing to decreases of voltage-gated Na^+ conductance (Bliss and Rosenberg, 1979; De Col et al., 2008), low concentrations of tetrodotoxin reduce conduction velocity (Colquhoun and Ritchie, 1972), and depolarizations as brief as 2–3 ms inactivate RGC voltage-gated Na^+ current (Kim and Rieke, 2003; Hayashida and Ishida, 2004). However, the Na^+ current at adult rat optic nerve nodes of Ranvier appears to be carried by Nav 1.6 (Boiko et al., 2001) and this isoform is not significantly regulated by CaMKII (Chen et al., 2008; Aiba et al., 2010). A mixed cation current (I_h) is partially activated near resting potential in rat optic nerve (Eng et al., 1990), but RGC I_h is neither inactivated by depolarization (Lee and Ishida, 2007) nor known to be regulated by CaMKII. Last, a tetraethylammonium (TEA)-sensitive and 4-aminopyridine (4AP)-insensitive K^+ conductance reduces motoneuron conduction velocity after spikes (Baker et al., 1987), yet TEA has little (if any) effect on rat optic nerve spikes (Kocsis et al., 1986). By contrast, 4AP increases the time-to-peak of adult rat optic nerve CAPs (Foster et al., 1982; Devaux et al., 2002; Ogata, Partida, and Ishida, unpublished observations), 4AP-sensitive current can be inactivated by brief depolarizations (Debanne et al., 1997; Roeper et al., 1997; see also Geiger and Jonas, 2000), and K^+ currents through channels formed by Kv1.4 or Kv4.2 are inhibited by 4AP and increased by CaMKII (Roeper et al., 1997; Varga et al., 2004). Although Kv1.4 is absent in adult rat optic nerve (Rasband et al., 1999) and whether Kv4.2 is present is not yet known, voltage-induced inactivation of Na^+ current and 4AP-sensitive K^+ current plus decreases in 4AP-sensitive K^+ current by activity-induced decreases in pT286 could explain the propagation slowing that we observed. CaMKII inhibition and changes in pT286 levels might affect the response to single, paired, or multiple pulses by altering the 4AP-sensitive K^+ conductance and thereby altering the time and space constants of optic nerve axons (Jack et al., 1975). On the other hand, inhibition of either CaMKII or PP2A would not be expected to alter Na^+ channel inactivation because the Na^+ channels are Nav1.6. Therefore, Na^+ channel inactivation could alter the response to paired pulses even if CaMKII is inhibited.

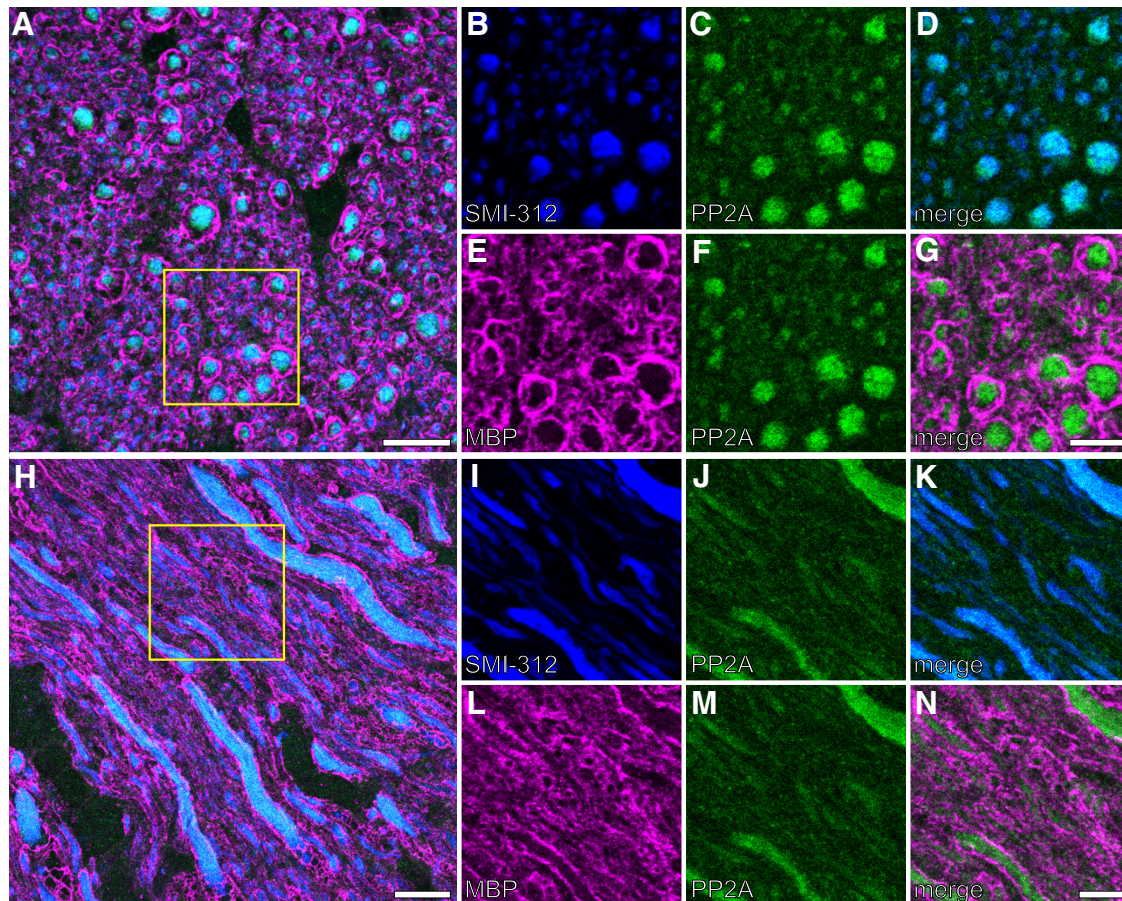


Figure 11. PP2A in optic nerve axons. Transverse (**A**) and longitudinal (**H**) sections triple-labeled for SMI-312 (blue), PP2A (green), and MBP (magenta). **B–D**, SMI-312 (**B**), PP2A (**C**), and merge thereof (**D**) of the area within the yellow square in **A**. SMI-312 and PP2A colocalization generates cyan. **E–G**, MBP (**E**), PP2A (**F**), and merge thereof (**G**) of the same field as **D**. Separation of magenta and green in **G** shows that PP2A localizes to axons and not to myelin. The axon diameters (e.g., in **A** and **G**) range between small and large. **H–N**, Comparable information in cross-section, including colocalization (cyan in **K**) of SMI-312 (**I**) and PP2A (**J**), separation (**N**) of MBP (**L**) and PP2A (**M**), and wide range of axon diameters (**N**). **I–N**, Field outlined by yellow square in **H**. Scale bars: **A**, **H**: 10 μm ; **G**, 3 μm (applies to **B–G**); **N**, 3 μm (applies to **I–N**).

Spike frequencies *in situ*

Our finding that single cardinal CAPs can slow the propagation of follower CAPs is consistent with recordings from optic tract (Bishop et al., 1953) and other preparations (Swadlow et al., 1980; Lüscher et al., 1994; George et al., 2007; Meeks and Mennerick, 2007).

These results suggest that, if spikes are elicited in RGCs intraretinally and recorded in the optic tract, then the spike intervals in the tract will be longer than the stimulus intervals in the retina. For example, we estimate that a stimulus frequency of 333 Hz at the retina (i.e., a pair of pulses separated by 3 ms) will elicit spikes at an instantaneous frequency as low as 220 Hz at a distance of 20 mm from the eye (see Results). Moreover, we found that cardinal CAPs slowed follower CAP propagation by greater amounts as the stimulus intervals became shorter than 20 ms (Figs. 1A, 12C,D). This predicts that, at synapses where signal transmission probability increases with stimulus frequency, the probability will not increase at high stimulus frequencies as much as it does at low or moderate frequencies. This trend can be seen in plots of efficacy versus interspike intervals of RGCs (Usrey et al., 1998; Carandini et al., 2007) and of optic tract fibers (Singer et al., 1972). The spike slowing that we describe here might thus limit the spike frequencies that propagate along the optic nerve, chiasm, and tract and, in turn, that retinogeniculate synapses can transmit.

Although we have no means of assessing relative contributions, we suppose that activity-induced spike slowing will combine with LGN receptor desensitization to transmitter released by RGCs (Chen et al., 2002), presynaptic inhibition (Chen and Regehr, 2003), and postsynaptic summation (Carandini et al., 2007) to shape the frequency dependence of retinogeniculate signal transmission. At the same time, spike-induced increases in spike interval might not reduce signal transmission at synapses where single spikes transmit with very high efficacies, e.g., from RGCs to rat LGN P cells (Burke and Sefton, 1966), some mouse and cat LGN cells (Cleland and Lee, 1985; Blitz and Regehr, 2003), and optic tectum cells in frog (George et al., 1984) and salamander (Gollisch and Meister, 2008). Also, some rodent RGC populations may be organized to ensure signal transmission by other means, such as sampling of individual points in space by cells with overlapping dendritic fields (Peichl, 1989; Zhang et al., 2012), synapses from more than one RGC of a given type onto single LGN cells (Hammer et al., 2015; Weyand, 2016; Rompani et al., 2017), and homologous electrical coupling (Völgyi et al., 2013). By correlating spike firing in multiple RGCs, these properties might increase transmission efficacy, as found in LGN inputs to visual cortex (Alonso et al., 1996) and modeled for primate RGC spikes (Pillow et al., 2005).

On average, the time-to-peak of P2 and P3 increased more than that of P1 as stimulus interval decreased (Figs. 1A, 12D).

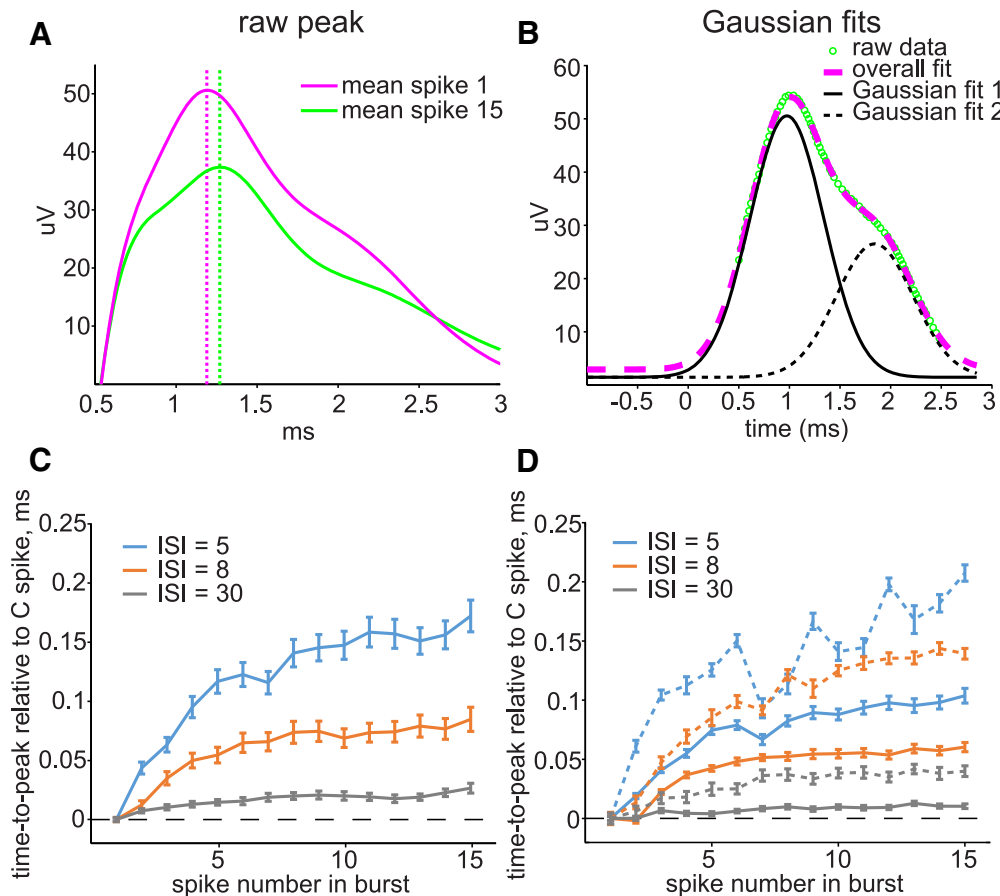


Figure 12. Spiking slows follower spike propagation *in vivo*. **A**, CAPs activated in optic chiasm and recorded in optic nerve (near optic foramen). Responses to the first (magenta) and fifteenth (green) pulse in a train of 15 stimulus pulses, all separated by 5 ms. Traces are means from 76 repeats of the stimuli aligned to the beginning of the stimulus artifacts. Dashed vertical lines drawn through the maximum amplitude of the responses showing that the time-to-peak of the fifteenth CAP is longer than that of the first CAP. **B**, CAP (traced by green dots) elicited by a single pulse, displaying a larger peak at $t \approx 1$ ms and a shoulder-like bulge at $t \approx 2$ ms. Plots of one Gaussian function peaking before (solid black line) and another peaking after (dashed black line) the large peak of the raw trace. The sum of these functions (dashed magenta line) follows the raw data trace on both the rising and falling phases. **C**, Effect of conditioning pulses on the time-to-peak of the large (maximum amplitude) peak recorded in three *in vivo* preparations. **D**, Effect of conditioning pulses on the time-to-peak of P1 (solid lines) and P2 (dashed lines) of CAPs in two preparations. **C, D**, Time-to-peak of the CAPs elicited by each of 15 pulses relative to that elicited by the first pulse. The values plotted are the means of the responses of each preparation to 40–50 presentations of 15 pulses at each of 3 ISIs (5, 8, and 30 ms). Lines connect the means of the delays at each ISI. Error bars indicate \pm SEM.

This predicts that the maximum spike frequencies that can be reached by slowly conducting RGC axons are lower than those of rapidly conducting units (see Results) and this has been observed *in vivo* (Fukada et al., 1966; see also Swadlow, 1985; Perge et al., 2012). Would reducing spike frequency be advantageous? Possibly yes because it would tend to reduce vesicle depletion in single RGC axon terminals, permit more recovery from postsynaptic receptor desensitization at individual synapses (Chen et al., 2002), and reduce the energy required to maintain normal intracellular Na^+ and Ca^{2+} (Stys et al., 1992a; Attwell and Laughlin, 2001) concentrations. Our recordings showed that the fastest conduction velocities changed the least and the slowest conduction velocities changed the most. These might reflect an inverse relation between fiber size and increase in Na^+ concentration and/or lower amounts of CaMKII, less CaMKII autophosphorylation, or more avid pT286 dephosphorylation in smaller-diameter axons.

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