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Characterization of norepinephrine-evoked inward currents in interstitial cells isolated from the rabbit urethra

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The urethra remains unknown, although a number of sharp microelectrode recordings made from the urethras of guinea pigs and rabbits demonstrated that exogenously applied norepinephrine (NE) could increase the frequency of slow waves (9, 12).

Recent studies in our laboratory (27, 28) demonstrated that a small population of freshly dispersed cells from the rabbit urethra shared similar characteristics with interstitial cells of Cajal (ICC) believed to be the pacemaker cells in the gastrointestinal (GI) tract (17, 19, 25, 26, 31, 32). These urethral interstitial cells were vimentin positive, noncontractile, and spontaneously active and were therefore postulated to be “pacemaker” cells in the rabbit urethra. A parallel could therefore be made between interstitial cells in the urethra and ICC in the GI tract. Because ICC act not only as pacemakers but also as mediators of neurotransmission (8, 26, 34, 35), it was of interest to investigate whether interstitial cells could play a similar role in the urethra. In support of this hypothesis is the observation that exogenously applied NE increased the frequency of spontaneous transient depolarizations (STDs) recorded from isolated urethral interstitial cells (27), demonstrating that NE could directly influence isolated urethral interstitial (pacemaker) cells. This novel finding may prove to have important implications with regard to mechanisms controlling urethral tone. In this study we have characterized the mechanisms underlying the effect of exogenous NE on isolated interstitial cells and investigated whether these cells contribute to neurogenic responses in the rabbit urethra.

**MATERIALS AND METHODS**

The bladder and urethra were removed from both male and female rabbits immediately after they had been killed by lethal injection of pentobarbitone. The most proximal 1 cm of the urethra was removed and placed in Krebs solution, and from this, strips were dissected for cell dispersal or for tension recording.

*Cell dispersal.* Strips of proximal urethra 0.5 cm in width were cut into 1-mm³ pieces and stored in Hanks’ Ca²⁺-free solution for 30 min before being incubated in dispersal me-

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10 μM Norepinephrine

10 μM Norepinephrine

10 μM Norepinephrine

Fig. 1. Effect of norepinephrine (NE) on isolated interstitial cells. In voltage clamp, NE (10 μM) could either increase the frequency of spontaneous transient inward currents (STICs; A), evoke a series of inward currents of diminishing amplitude (B), or evoke a large single inward current (C).

Solutions. The compositions of the solutions used were as follows (in mM): 1) Hanks’ solution, 129.8 Na+, 5.8 K+, 135 Cl−, 4.17 HCO3−, 0.34 HPO42−, 0.44 H2PO4−, 1.8 Ca2+, 0.9 Mg2+, 0.4 SO42−, 10 glucose, 2.9 sucrose, and 10 HEPES, pH adjusted to 7.4 with NaOH; 2) 49 mM Cl− Hanks’ solution, 129.8 Na+, 5.8 K+, 86 glutamate, 49 Cl−, 4.17 HCO3−, 0.34 HPO42−, 1.8 Ca2+, 0.9 Mg2+, 0.4 SO42−, 10 glucose, 2.9 sucrose, and 10 HEPES, pH adjusted to 7.4 with NaOH; and 3) Ca2+-free Hanks’ solution, 129.8 Na+, 5.8 K+, 4.17 HCO3−, 0.44 H2PO4−, 1.8 Ca2+, 0.9 Mg2+, 0.4 SO42−, 10 glucose, 2.9 sucrose, and 10 HEPES, pH adjusted to 7.4 with NaOH; and 4) Krebs solution 146.2 Na+, 5.9 K+, 133.3 Cl−, 25 HCO3−, 1.2 H2PO4−, 2.5 Ca2+, 1.2 Mg2+, and 11 glucose, pH maintained at 7.4 by bubbling with 95% O2-5% CO2.

Perforated-patch recordings from single cells. Currents were recorded by using the perforated-patch configuration of the whole cell patch-clamp technique (15, 24). This circumvented the problem of current rundown encountered using the conventional whole cell configuration. The cell membrane was perforated by using the antibiotic amphotericin B (600 μg/ml). Patch pipettes were initially front-filled by being dipped into pipette solution and were then backfilled with the amphotericin B-containing solution. Pipettes were pulled from borosilicate glass capillary tubing (1.5-mm outer diameter, 1.17-mm inner diameter; Clark Medical Instruments) to a tip of diameter ~1–1.5 μm and resistance of 2–4 MΩ.

Series resistance and capacitative currents were usually uncompensated for in this study. Voltage-clamp commands were delivered via an Axopatch one-dimensional patch-clamp amplifier (Axon Instruments), and membrane currents were recorded by a 12-bit analog-to-digital/digital-to-analog converter (Axodata 1200 or Labmaster-Scientific Solutions) interfaced to an AT-type computer running PCLAMP software. During experiments, the dish containing the cells was continuously perfused with Hanks’ solution (see Solutions) at 37 ± 1°C. Additionally, the cell under study was continuously superfused by means of a custom-built close delivery system with a pipette of tip diameter 200 μm placed ~300 μm from the cell. The Hanks’ solution in the close delivery system could be switched to a drug-containing solution with a dead-time of <5 s. The effect of NE was quantified by examining the peak amplitude of the inward currents evoked by NE. Where stated, data were corrected for junction potentials of ~3 mV in standard Hanks’ solution and +2 mV in low-chloride Hanks’. Summary data are presented as means ± SE, and statistical comparisons were made on raw data by using Students’ paired t-test, taking the P < 0.05 level as significant.

Isolated tissue recording. Circularly orientated strips (8 × 1 × 1 mm) of smooth muscle were removed from rabbit urethra, placed in a water-jacketed organ bath maintained at 37°C, and perfused with warmed Krebs solution that was bubbled with 95% O2-5% CO2. Strips were adjusted to a tension of 2–4 mN and allowed to equilibrate for 50 min before experimentation began. Contractions were measured by using Statham UC3 and Dynamometer UF1 transducers, the outputs of which were recorded on a Grass 7400 chart recorder. Transmural nerve stimulation was applied by a Grass S11 stimulator, which delivered 0.3-ms pulses of 50 V (nominal) at a frequency of 10 Hz. Pulses were applied for 0.1, 0.3, 1, 3, and 10 s. Stimuli were applied ~1 min after the preceding neurogenic contraction had returned to baseline levels. All experiments were carried out in the presence of atropine (1 μM) and Nω-nitro-L-arginine (10 μM) to block contributions from cholinergic and nitricergic nerves.

Drugs. The following drugs were used: amphotericin B (Sigma), 2-aminoethoxydiphenyl borate (2-APB; Sigma), 9-anthracenecarboxylic acid (9-AC; Sigma), 2-nitro-4-carboxyphenyl-N,N-diphenylcarbamate (NCDC; Sigma), atropine (BDH Laboratories), caffeine (Sigma), cyclopiazonic acid (CPA; Calbiochem), dimethyl sulfoxide (DMSO; Sigma), Nω-nitro-L-arginine (Sigma), niflumic acid (Sigma), NE HPO42−, 0.44 H2PO4−, 1.8 Ca2+, 0.9 Mg2+, 0.4 SO42−, 10 dextrose, 2.9 sucrose, and 10 HEPES, pH adjusted to 7.4 with NaOH; 3) Ca2+-free perfused-patch pipette solution, 133 Ca2+, 135 Cl−, 1.0 Mg2+, 0.5 EGTA, and 10 HEPES, pH adjusted to 7.2 with CsOH; and 4) Krebs solution 146.2 Na+, 5.9 K+, 133.3 Cl−, 25 HCO3−, 1.2 H2PO4−, 2.5 Ca2+, 1.2 Mg2+, and 11 glucose, pH maintained at 7.4 by bubbling with 95% O2-5% CO2.

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(Levophed; Sanofi Winthrop, UK), penitrem A (Sigma), phenotolamine (Ciba), and prazosin (Tocris).

Stock solutions of 9-AC (0.1 M), niflumic acid (0.01 M), penitrem A (0.1 mM), and CPA (0.01 M) were made in DMSO. 2-APB and NCDC were made in stock solutions of ethanol (0.1 M). Drug vehicles had no effect on the currents studied. Prazosin was dissolved in distilled water to give a 0.01 M stock solution. All drugs were then diluted to their final concentrations in Hanks’ or Krebs solution. NE and phenotolamine were dissolved directly into experimental solutions to give the required concentrations.

RESULTS

Digestion of rabbit urethral smooth muscle strips yielded both interstitial cells and smooth muscle cells. Although the majority of cells were smooth muscle, interstitial cells could be easily distinguished from smooth muscle cells by using a number of criteria (27). Thus interstitial cells were highly branched, failed to contract in response to depolarizing current injection or application of NE, possessed abundant Ca2+-activated Cl− current, and normally fired spontaneous transient inward currents. In a previous study we demonstrated that NE (10 μM) enhanced the frequency of slow waves in isolated urethral interstitial cells, but the mechanisms underlying this effect were not examined (27). In this study we carried out a series of experiments to establish the mechanisms through which NE mediates its effects on these cells.

Effect of NE. Figure 1 shows typical examples of the effect of NE (10 μM) on spontaneous activity in interstitial cells held under voltage clamp. When cells were held at −60 mV, spontaneous transient inward currents (STICs) were apparent, and application of 10 μM NE evoked a large inward current followed by a series of currents of diminishing amplitude. These results suggest that the elevation of STIC frequency underlies the increase in slow wave activity observed previously (27). The effects of NE were reproducible within the same cell if a period of 80 s was allowed between successive applications, presumably reflecting a combination of the time taken for the NE-sensitive store to refill and the cell to recover from desensitization. Although the effects of NE on individual interstitial cells were reproducible, variable responses to NE were elicited in different cells. In 16 of 35 cells examined, NE evoked currents similar to those shown in Fig. 1, A and B. They consisted of a large transient inward current that was followed by a series of smaller inward currents with a mean frequency of 7 ± 1 min−1. In the remaining 19 cells, application of NE evoked large single inward currents similar to that shown in Fig. 1C.

Effect of α-adrenoceptor antagonists. Having demonstrated that NE could evoke large transient inward currents, we assessed the involvement of α-adrenoceptors in this response by examining the effect of phenotolamine and prazosin on the response to NE. Figure 2A shows a typical response to NE before, during, and after application of the nonspecific α-adrenoceptor antagonist phenotolamine (1 μM). Figure 2B shows a summary bar chart for six experiments in which phenotolamine decreased the mean peak inward current from

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**Fig. 2.** Blockade of NE response by α-adrenoceptor antagonists. Application of NE evoked a series of inward currents, which were reversibly abolished by phenotolamine (A) or prazosin (C). B and D show summary data in which the effect of NE was examined before (control), during, and after (wash) application of phenotolamine (1 μM) or prazosin (1 μM), respectively.
We next examined the effect of the STICs by releasing Ca$^{2+}$ from intracellular stores, we examined the effects of the Ca$^{2+}$-ATPase inhibitor CPA (10 μM) on the response. Figure 3A shows a typical experiment in which the response to repeated applications of NE was examined in the presence of CPA. Before application of CPA, NE evoked large inward currents that became progressively smaller in the presence of CPA. Figure 3B shows a summary for six similar experiments in which the peak inward current was reduced from −604 ± 149 pA to −116 pA ± 63 pA after 240 s in CPA ($P < 0.05$). Within 160 s of the removal of CPA, NE evoked large inward currents that were not significantly different from control (−501 ± 180 pA). These data support the idea that NE caused the release of Ca$^{2+}$ from intracellular stores.

**Inhibition of phospholipase C with NCDC.** Previously, we demonstrated (21) that the inhibitor of inositol 1,4,5-trisphosphate (IP$_3$)-dependent release, 2-APB, abolished the response to NE in urethral interstitial cells (28). More recent studies (23) have questioned the specificity of 2-APB and suggested that it may act by inhibiting store refilling rather than inhibiting store release. Therefore, we examined the effects of the phospholipase C inhibitor NCDC (100 μM) on NE-evoked currents. Figure 4A shows a typical experiment in which NE evoked large inward currents before the addition of NCDC. In its presence, spontaneous activity was abolished as reported previously (28), and the response to NE was inhibited (Fig. 4B). Figure 4C shows a summary for six cells in which NE evoked mean inward currents that were −396 ± 97 pA in amplitude before, compared with −14 ± 9 pA after, application of NCDC ($P < 0.05$). To test directly for the possibility that NCDC blocked Cl$^{-}$ channels, we examined its effects on Cl$^{-}$ currents evoked by caffeine. Figure 4D shows a typical experiment in which application of caffeine (10 mM) to a cell held at −60 mV evoked a large inward current. In the presence of NCDC (100 μM; Fig. 4E), application of caffeine evoked a current of similar amplitude to control. Figure 4F shows a summary of four experiments in which caffeine application evoked large inward currents of −383 ± 77 pA. In the presence of 100 μM NCDC, the caffeine-evoked currents were slightly increased to −439 ± 67 pA, but this was not significant ($P = 0.08$).

The above data suggest that the effects of NCDC are not mediated through blockade of Cl$^{-}$ channels and are consistent with our previous findings that the effects of NE are mediated by an IP$_3$-dependent mechanism (28).

**Involvement of Ca$^{2+}$-activated Cl$^{-}$ channels.** The data presented so far support the idea that activation of α$_1$-adrenoceptors stimulates phospholipase C, leads to the production of IP$_3$, and consequently releases Ca$^{2+}$ from intracellular stores. This, in turn, activates Ca$^{2+}$-sensitive channels on the cell membrane and elicits large transient inward currents. To test whether the NE-induced currents were carried by Cl$^{-}$, we utilized solutions with Cl$^{-}$ as the only anion in the intracellular solution, as these studies reported that the inward currents were carried by Cl$^{-}$ (27). We found that NE-evoked inward currents were carried by Cl$^{-}$ ions through Ca$^{2+}$-activated Cl$^{-}$ channels previously characterized in these cells (27), we examined the reversal potential of the NE-induced current and the effects of different Cl$^{-}$ channel blockers on this current. Figure 5A shows the effects of NE on a cell recorded with symmetrical Cl$^{-}$ solutions [chloride equilibrium potential ($E_{Cl}$) = 0 mV] and held at a variety of voltages ranging from −60 to +40 mV. When the cell was held at −60 mV, NE application evoked inward currents that became progressively smaller in amplitude as the cell was depolarized toward 0 mV. At potentials positive to 0 mV, the currents were outward and increased.
in amplitude as the cell was held at more depolarized potentials. A summary of six similar experiments is shown in Fig. 5B where the peak amplitude of the inward current was plotted at different holding potentials. Under these conditions, the NE-evoked current reversed close to 0 mV, suggesting that the current was carried by Cl\(^-\) ions. In a separate set of experiments, we examined the effects altering the external Cl\(^-\)/H\(^+\) concentration on the reversal potential of the NE-evoked current. A series of 400-ms voltage ramps from -50 to +50 mV were applied to cells every 500 ms before and during application of NE. Figure 5C shows a typical experiment in which the NE-sensitive currents were obtained and plotted against voltage. Under control conditions (\(E_{Cl} = 0\) mV), the NE-sensitive current reversed at -1 mV, and when the external Cl\(^-\) concentration was decreased to 49 mM (\(E_{Cl} = 27\) mV), the reversal potential of the inward current shifted to 25 mV. In four similar experiments, the reversal potential of the NE-sensitive currents shifted from 0 to 23 ± 2 mV when external Cl\(^-\) was reduced from 135 to 49 mM (\(P < 0.01\), data corrected for junction potentials of -3 mV and +2 mV in normal and reduced external Cl\(^-\) solutions, respectively).

The above data are consistent with the idea that NE-evoked currents are due to the stimulation of Cl\(^-\) currents. Further evidence to support this is presented in Fig. 6. In Fig. 6, A and C, application of NE induced currents that were markedly reduced in the presence of either 9-AC (1 mM) or niflumic acid (10 \(\mu\)M), respectively. Figure 6B shows a summary for six experiments in which 9-AC (1 mM) significantly reduced the inward currents evoked by NE from -1,253 ± 510 pA to -299 ± 89 pA (\(P < 0.05\)). Similarly, niflumic acid decreased the NE-evoked inward currents from -813 ± 185 pA to -140 ± 26 pA (\(P < 0.05\), \(n = 6\)). After washout of both drugs, the response to NE returned and was not significantly different from control.

**Tension recordings.** Having characterized the effects of NE on isolated urethral interstitial cells, we next wanted to establish whether they played any role in

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**Fig. 4. Inhibition of phospholipase C abolishes the response to NE.** A: effect of NE in the absence of 2-nitro-4-carboxyphenyl-N,N-diphenylcarbamate (NCDC). B: inhibition of phospholipase C in the presence of NCDC (100 \(\mu\)M) abolished both spontaneous activity and the response to NE. C: a summary of 6 similar experiments in which NCDC abolished both spontaneous activity and the response to NE. D and E show the lack of effect of NCDC on caffeine (Caff)-evoked Cl\(^-\) currents. F: a summary of 4 similar experiments in which NCDC failed to block caffeine-evoked Cl\(^-\) currents.
mediating noradrenergic neurotransmission. To test this possibility, the effect of transmural nerve stimulation was examined on strips of urethra in the presence of atropine (1 μM) and Nω-nitro-L-arginine (10 μM) to block the contribution of cholinergic and non-adrenergic noncholinergic (NANC) transmitters, respectively. Pulses (0.3-ms duration, 10 Hz) were applied for 0.1, 0.3, 1, 3 and 10 s. Figure 7, A and C, shows typical responses of a urethral strip to each stimulation. Increasing the duration of stimulation increased both the amplitude and duration of contraction. In the presence of phentolamine (1 μM), the contractile responses to nerve stimulation were abolished, suggesting that only noradrenergic responses were evoked under these conditions (n = 3, data not shown). We next examined the effects of blocking Cl− channels on the response to nerve stimulation. Figure 7A shows a typical experiment in which nerve-evoked contractions were elicited in the absence and presence of niflumic acid (100 μM). Because niflumic acid has been demonstrated to open large conductance Ca2+-activated K+ (BK) channels (11), all experiments were carried out in the presence of penitrem A (100 μM) to block BK channels (14, 18). Application of niflumic acid reduced spontaneous contractions and decreased the amplitude of nerve-evoked contractions. The bar chart in Fig. 7B shows a summary of six similar experiments where the peak contraction amplitude was measured before and during application of niflumic acid. Data were normalized to the response obtained by a stimulation duration of 10 s in the absence of Cl− channel blockers. In the presence of niflumic acid, the amplitude of contraction was significantly reduced at 0.3, 1, and 10 s to 14 ± 4, 16 ± 5, and 18 ± 4%, respectively (P < 0.05). The effects of 9-AC (1 mM) were also examined on neurogenic contractions. Figure 7C shows a typical response to nerve stimulation before and during application of 9-AC. Note again that both spontaneous and neurogenic contractions were depressed in the presence of 9-AC. Figure 7D shows summary data for seven similar experiments in which 9-AC reduced the amplitude of contraction in response to 0.3-, 1-, and 10-s stimu-
tion periods to 16 ± 4, 16 ± 3, and 30 ± 5%, respectively (P < 0.05).

**DISCUSSION**

A variety of studies have demonstrated that ICC play a central role in the initiation, coordination, and modulation of spontaneous activity in the GI tract (25, 26, 31). Cells with characteristics similar to the ICC in the gut have recently been isolated in a variety of tissues from the lower urinary tract (22, 27, 28). Although their function in these tissues has not been thoroughly established, we have hypothesized that urethral interstitial cells contribute to the generation of tone by acting as pacemakers to "drive" the surrounding smooth muscle (27). Two main pieces of evidence support this contention. First, isolated interstitial cells fire regular spontaneous slow waves, which resemble the electrical activity recorded in whole tissues with microelectrodes, whereas smooth muscle cells are electrically quiescent (12, 13, 27). Second, the slow waves recorded in whole tissues and isolated interstitial cells are abolished when Ca\(^{2+}\) release from IP\(_3\)-sensitive stores is inhibited or Ca\(^{2+}\)-activated Cl\(^{-}\) channels are blocked (12, 27, 28).

Modulation of this mechanism by excitatory neurotransmitters such as NE could provide an efficient means to alter tone, because an increase in slow wave frequency could lead to an enhancement of contractile force. Previous studies have demonstrated that NE is the major excitatory transmitter in the urethra (2, 3), and for this reason we examined the mechanisms of its action on rabbit urethral interstitial cells. Hashitani et al. (12) demonstrated that NE increases the frequency of STDs in the rabbit urethra by enhancing the oscillatory release of Ca\(^{2+}\) from intracellular stores, which results in the periodic activation of Cl\(^{-}\) channels. Because urethral smooth muscle cells in the rabbit possess little Ca\(^{2+}\)-activated Cl\(^{-}\) current (27), it is unlikely that the observed changes in electrical activity recorded with intracellular microelectrodes reflect an action on these cells and leaves the intriguing possibility that interstitial cells act as the primary target for neuronally released NE.

In a previous study (27), we demonstrated that under current-clamp conditions, NE produced responses in isolated interstitial cells that appear to be remarkably similar to those recorded in whole tissue preparations with sharp microelectrodes (12). Thus NE produced an initial large slow wave that was followed by a series of more frequent but shorter slow waves. The reason for such a pattern of activity became apparent when cells were held under voltage clamp, where NE typically induced a large inward current that was followed by more frequent, gradually diminishing inward currents. A number of lines of evidence suggest that the NE-evoked currents reflected the stimulation of Ca\(^{2+}\)-activated Cl\(^{-}\) currents that have been demonstrated in a variety of smooth muscles (1, 20). First, when cells were held under voltage clamp, the currents reversed close to \(E_{Cl}\) in symmetrical Cl\(^{-}\) solutions and shifted in a predictable manner when \(E_{Cl}\) was altered. Second, they were significantly decreased by either
niflumic acid or 9-AC. Although the concentration of 9-AC (1 mM) used to block the response is relatively high compared with other tissues (20), we have previously demonstrated that this concentration selectively blocks the Cl\(^{-}\)/H\(^{+}\) current in urethral interstitial cells (27).

When we examined the pharmacology of the NE response, it appeared to be similar to the pathway described in a number of smooth muscles (1, 4, 5, 30). Thus the response to NE was abolished by either phentolamine or prazosin, suggesting that it was mediated via activation of \(\alpha\)-adrenoceptors. Similarly, its effects were attenuated when phospholipase C was blocked with NCDC, which is consistent with our previous observation that the IP\(_3\) inhibitor 2-APB (21) inhibits NE responses in urethral interstitial cells (28). These data support the idea that NE mediates its effects by upregulating the normal pacemaking mechanism in these cells.

In recent years, a number of studies have demonstrated that ICC in the gut act as important intermediaries in both excitatory and inhibitory neurotransmission (8, 26, 34, 35). The evidence to support this role in the gut has been obtained from mutant mice (W/W') that failed to develop intramuscular ICC. In these animals, both excitatory and inhibitory nerve evoked responses were greatly attenuated (8, 34, 35). Unfortunately, similar rabbit knockouts are presently unavailable. However, we have previously demonstrated that in marked contrast to the smooth muscle cells, interstitial cells in the urethra possess abundant Ca\(^{2+}\)/H\(^{+}\)-activated Cl\(^{-}\)/H\(^{+}\) current (27). We exploited this difference in the present study to assess the contribution of interstitial cells to noradrenergic neurotransmission in strips of urethra by attempting to pharmacologically knock out interstitial cells using Cl\(^{-}\)/H\(^{+}\) channel blockers. Although this method lacked the elegance of genetic knockout technologies, it produced effects that were consistent with abolishing the contribution of interstitial cells. Thus, in the presence of either niflumic or 9-AC, both spontaneous and neurogenic contractions were attenuated. These data tentatively suggest that interstitial cells in the urethra not only contribute to spontaneous mechanical activity but also may play an important role in neurotransmission.

Although the idea that interstitial cells mediate neurotransmission in the urethra may be appealing, the results obtained on whole tissue strips should be interpreted with caution. This preparation consists of a

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**Fig. 7.** Neurogenic contractions are attenuated in the presence of Cl\(^{-}\)/H\(^{+}\) channel blockers. A: effect of field stimulation on tension in a strip of rabbit urethra. Pulses [0.3-ms duration, 50 V (nominal), 10 Hz] were applied for 0.1, 0.3, 1, 3, and 10 s. In the presence of 100 \(\mu\)M niflumic acid (bottom). B: a summary bar chart for 6 experiments in which the amplitude of contraction at each stimulation frequency was plotted in the absence (control) and presence of niflumic acid. C: a typical example of the effect of nerve stimulation before (top) and during (bottom) application of 9-AC (1 mM). D: a summary of 6 similar experiments.
complex arrangement of nerves, smooth muscle, and interstitial cells, all of which may be affected by the Cl\(^-\) channel blockers. The observed effects on neurogenic contractions may be caused by a presynaptic action of the Cl\(^-\) channel blockers, which could inhibit neurotransmitter output. However, inhibition of neuronal Cl\(^-\) channels should prolong neuronal action potentials and, consequently, enhance neurotransmitter output. Alternatively, the Cl\(^-\) channel blockers could inhibit neurotransmitter output by blocking neuronal Ca\(^{2+}\) channels. Although we cannot rule out this possibility, it is interesting to note that Jackson et al. (16) have failed to demonstrate any effect of niflumic acid on action potential-evoked axonal Ca\(^{2+}\) transients in the rat vas deferens, suggesting that Ca\(^{2+}\) influx is unaffected by niflumic acid.

Recent immunohistochemical evidence presented by Smet et al. (29) and Waldeck et al. (33) would also support our contention that urethral interstitial cells play a role in neurotransmission. These studies demonstrated the presence of branched interstitial cells in the human, guinea pig, and rabbit urethras, respectively, that were immunopositive for cGMP and support the idea that they may be important in mediating neurally released nitric oxide responses. Unfortunately, the detailed immunohistochemistry and electronmicroscopy data that have been provided to support the role of ICC as mediators of neurotransmission in the gut provided are not yet available in the urethra. Future studies should focus on the relationships among nerves, interstitial cells, and the surrounding smooth muscle in the urethra to determine whether the interstitial cells are selectively innervated.

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