Interaction of Atriopeptin III and Vasopressin on Calcium Kinetics and Contraction of Aortic Smooth Muscle Cells

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Abstract

The cellular mechanism of the vasodilatory action of atriopeptin III (APIII) on vasopressin (AVP)-induced Ca2+ mobilization and cell shape change in cultured vascular smooth muscle cells (VSMC) was studied. APIII (10−8 M) attenuated the increase of intracellular free Ca2+, [Ca2+]i, induced by 10−8 M AVP (234.0±14.8 vs. 310.0±28.4 nM, P < 0.01). Similar results were obtained in 45Ca2+ efflux experiments. APIII (10−7 M), however, did not alter AVP-induced inositol trisphosphate (IP3) production, although the levels of inositol-1-phosphate were significantly reduced. The effect of APIII to block or attenuate AVP-induced Ca2+ mobilization was associated with an inhibition of AVP-stimulated cell shape change. The effect of atrial natriuretic factor (ANF) on cell shape, however, occurred at lower ANF concentrations than the effect on the Ca2+ mobilization. APIII stimulated production of cyclic guanosine monophosphate (cGMP) in VSMC. The effect of APIII on AVP-stimulated Ca2+ mobilization was partially mimicked by the stable nucleotide 8-bromo cGMP and was not affected by the soluble guanylate cyclase inhibitor, methylene blue (10−4 M). These results suggest that APIII exerts its vasodilatory effect, in part, by interference with vasopressin-stimulated Ca2+ mobilization in vascular smooth muscle cells, perhaps by stimulating particulate guanylate cyclase and cGMP. However, an effect of ANF on the contractile mechanism at a site independent of Ca2+ release is also suggested by the present results.

Introduction

The isolation, characterization, and amino acid sequencing of the atrial natriuretic peptides from atrial cardiocytes has stimulated considerable investigation into the mechanism of action of these substances (1–3). In addition to the diuretic and natriuretic effects of these peptides, specific binding sites have been demonstrated for these atrial peptides in a variety of vascular tissues (4–8). Although a vasoconstrictor effect of atrial natriuretic factor (ANF)1 has been shown in certain experimental settings (9), in general ANF has been found to exert a vasodilatory action (3–14). This vasodilation by ANF, however, seems to be most readily demonstrable in the presence of vasoconstrictors (3, 9, 12).

This latter observation, therefore, suggests that ANF may exert its vasodilatory effect, at least in part, by interfering with the cellular action of vasoconstrictors. This possibility was tested in the present study by examining the interaction of arginine vasopressin (AVP) and atriopeptin III (APIII), a 24-amino acid peptide with well-established vasodilatory properties (5), on 45Ca2+ efflux, 45Ca2+ influx, cytosolic free Ca2+, and cellular contraction of rat vascular smooth muscle cells in culture. In addition, since it has been suggested that cyclic guanosine monophosphate (cGMP) may be an intracellular mediator of ANF (15, 16), the effect of the stable nucleotide analogue, 8-bromo cGMP, on AVP-mediated cellular Ca2+ kinetics and cell contraction was also examined. Finally, in view of results showing a decrease of Ca2+ mobilization by AVP in the presence of ANF, we studied the AVP-produced production of inositol trisphosphate (IP3), the putative mediator of Ca2+ release, both in the presence and absence of ANF.

Methods

APIII was obtained from Peninsula Laboratories, Inc. (Belmont, CA). 45CaCl2 was purchased from New England Nuclear (Boston, MA). 8-bromo cGMP, AVP, methylene blue, quin 2, and quin 2-aceoxy methylester (quin 2-AM) were purchased from Sigma Chemical Co. (St. Louis, MO) and fura 2 was from Molecular Probes, Inc. (Eugene, OR). [3H]inositol and radiolabeled inositol phosphates were purchased from Amersham Corp. (Arlington Heights, IL). Digitonin was obtained from Calbiochem-Behring Corp. (LaJolla, CA).

Preparation of aortic smooth muscle cells and cell culture. Rat aortic smooth muscle cells were isolated using a modified method originally described by Chamley et al. (17). The thoracic aortas from six to eight male Sprague-Dawley rats were incubated at 37°C for 30 min in 7.5 ml of Eagle’s minimum essential medium (MEM) containing 2 mg/ml collagenase (Cooper Biomedical, Inc., Malvern, PA; 218 U/mg). After dissecting the adventitia, the aortas were minced with sterile razor blades and incubated again at 37°C in Eagle’s MEM (2 mg/ml collagenase) for 2.5–3 h under continuous stirring. After this incubation the resulting single cell suspension was centrifuged for 5 min at 1,000 rpm. The cell pellet was resuspended in fresh incubation medium without collagenase. This procedure was repeated twice. The cells were plated onto 35-mm cultures dishes at a density of 2.5–3.0 × 104 cells/dish and grown at 37°C in a humidified atmosphere of 95% air and 5% CO2. Eagle’s MEM, supplemented with 10% fetal calf serum, 100 U/ml penicillin, and 100 μg/ml streptomycin was used as culture medium. Cell viability was checked by the exclusion of trypan blue (0.3%) and always exceeded 95%. The cultures reached confluence after 7–10 d and were then used for experiments. Only primary cultures were used because vascular smooth muscle cells may lose their contractility when subcultured (18). Quin 2 and fura 2 loading and fluorescent measurements. For measurements of intracellular free Ca2+, [Ca2+]i, levels, aortic smooth muscle cells were grown on round glass coverslips (13-mm diam.) Upon confluence, the coverslips were washed twice with MEM without fetal calf serum. The coverslips were then incubated with serum-

1 Abbreviations used in this paper: ANF, atrial natriuretic factor; APIII, atriopeptin III; AVP, arginine vasopressin; 8-bromo cGMP, 8-bromo cGMP; IP, inositol phosphate(s); MIX, 3-isobutyl-1-methylxanthine; PSS, physiological saline solution.
free MEM containing 15 μM quin 2-AM for 60 min at 37°C in 95% air and 5% CO2. At the end of the loading period, the coverslips were washed three times with physiological saline solution (PSS; 140 mM NaCl, 4.6 mM KCl, 1.0 mM MgCl2, 2.0 mM CaCl2, 10 mM glucose, 10 mM Hepes, pH 7.4) and inserted into disposable fluorescence coverslites containing 3 mM PSS. The coverslites were placed in the thermostated holder of a fluorescence spectrophotometer (650-105, Perkin-Elmer Corp., Norwalk, CT). Fluorescence of quin 2-loaded cells was measured using an excitation wavelength of 393 nm (slit 5 nm) and an emission wavelength of 492 nm (slit 15–20 nm) (19).

The complete intracellular hydrolysis of quin 2-AM was judged by scanning the emission spectrum of quin 2 from 400 to 600 nm in the loaded cells. The cells exhibited a complete spectral shift peaking at 490–500 nm at the end of the loading procedure. None of the agents tested affected the fluorescence emission of nonloaded cells, nor did they have any significant fluorescence of their own.

A slow spontaneous decrease in fluorescence emission of quin 2-loaded cells [3±0.5% (mean±SE, n = 11)] of total Ca2+-dependent fluorescent emission after 2 min was observed due to photo bleaching of the probe and fluorescence tracings were corrected (during each experimental period) for the calculation of [Ca²⁺].

Cellular quin 2 concentration was estimated using a quin 2 free activated standard in the presence of nonloaded cells. Assuming an intracellular quin volume of ~0.25 μl/10⁶ cells (20), the calculated cellular quin 2 concentrations were 1.35±0.24 mM (mean±SE, n = 11).

A value of 115 nM was used as the dissociation constant (Kd) of the quin 2-Ca²⁺ complex and [Ca²⁺] was calculated as described (19, 21).

Leakage of quin 2 from the cells was excluded by adding 5 × 10⁻⁴ M MnCl₂ to quench the fluorescence of any extracellular quin 2. This maneuver did not affect the basal fluorescence.

For the fura 2 measurements, the monolayers on coverslips were incubated for 45 min at 37°C in MEM containing no FBS and with 4 μM fura 2 AM. The monolayers were then washed three times with MEM containing no fura 2 and incubated another 15 min at 37°C to allow time for any nonhydrolyzed fura 2-AM to diffuse from the cells. Before measurement, the coverslips were rinsed again with PSS containing 2 mM Ca²⁺. Fluorescence was measured at 37°C using the same equipment and conditions as for quin 2 determinations, at an emission wavelength of 500 nm and excitation wavelengths of 342 and 380 nm. Autofluorescence was measured in similar cells which had not been loaded with fura 2 and was <10% of the total fluorescence of fura 2-loaded cells in all the experiments. [Ca²⁺] was calculated as described by Grynkiewicz et al. (22). In this and the following sets of experiments, unless otherwise stated, cells were preincubated in the presence or absence of ANF by 5 min, before the addition of the effectors and ANF was simultaneously added with the effectors. Preincubation with ANF avoids any possible interference by AVP on other mediators of ANF-mediated cGMP generation (23).

4Ca²⁺ efflux. The culture medium was removed by aspiration and cell monolayers were rinsed twice with 1 ml PSS and loaded with 8 μCi of ⁴⁰Ca²⁺ in 1 ml of PSS at 37°C for 3 h. Then the cultures were rapidly rinsed 10 times with 1 ml of PSS and another 1 ml of PSS was added. The medium was removed and replaced with 1 ml of PSS at 1, 2, 3, and 4 min and thereafter at 30-s intervals. Hormones and other compounds were added as indicated during the time course of ⁴⁰Ca²⁺ efflux. The amount of ⁴⁰Ca²⁺ released from the cells in each time interval was measured by liquid scintillation counting.

4Ca²⁺ uptake. The culture medium was removed by aspiration and the cells were washed twice with 1 ml of PSS. Then the cells were incubated for 5 min with 2 μCi of ⁴⁰Ca²⁺ in the presence or absence of effectors in 1 ml of PSS. To terminate the uptake, external ⁴⁰Ca²⁺ was removed by rinsing the dish five times at 4°C with Ca²⁺-free PSS containing 2 mM EGTA. Intracellular radioactivity was extracted with 1 ml of sodium dodecyl sulfate-containing alkaline solution and measured by liquid scintillation counting.

Measurement of cGMP and cAMP. The cells were incubated for 10 min at 37°C in PSS containing 0.5 mM 3-isobutyl-1-methylxanthine (MIX), with or without APIII. The reaction was terminated by adding 100 μl of 0.1 N HCl to disrupt the cells. The cells were washed with a rubber policeman and centrifuged. The supernatants were saved for determination of cGMP and cAMP which was done by radioimmunoassay with commercial kits by New England Nuclear.

Protein was determined in all studies by the method of Lowry et al. (24).

Measurement of inositol phosphates. After cells were grown to confluence, the culture media of each dish was replaced with 1 ml of inositol-free medium containing 10 μCi of myo-2-³H]inositol (specific activity 16.3 Ci/mmol), and cells were incubated during 24 h. It has been previously shown that steady-state labeling occurs by 24 h. At the moment of the experiment, culture media was aspirated and replaced with 500 μl of 37°C PSS containing 10⁻⁸ or 10⁻⁷ M AVP. The reaction was stopped after 30 s using 500 μl of ice-cold trichloroacetic acid (20% solution). Cells were washed using an Eppendorf pipette top and aliquots were taken for protein determination. In the experiments using ANF, cells were preincubated with 10⁻⁷ M ANF in PSS for 5 min; after this period, this solution was replaced by PSS containing 10⁻⁷ M ANF and 10⁻⁸ or 10⁻⁷ M AVP.

The samples were centrifuged at 1,000 g for 10 min. The supernant containing inositol phosphates, was washed three times with an equal volume of ether and stored at −20°C until analysis, which was performed during the same week of the extraction. The water-soluble fraction was thawed and brought to pH 6.0 using Tris-base (50 μM). The extracts were applied to columns containing 1 ml of Dowex-I (X8, Formate form) and serially eluted with 1-ml aliquots of CH₃OH, Borax (5 mM disodium tetraborate), 60 mM sodium formate), 0.2, 0.5, and 1.0 M ammonium formate (in 0.1 M formic acid) separating, respectively, inositol, inositol, glycerophosphatidylinositol, inositol-1-phosphate, inositol bisphosphate, and IP_3 (25); this last fraction includes 1,3,4-, and 1,4,5-IP_3 and 1,3,4,5-inositol phosphate. Samples were collected and counted in a scintillation counter. The extraction procedure has been previously validated with radioactive phosphoinositides. Results were expressed as number of IP₃ counts per milligram of protein.

Smooth muscle cell shape change. The change in cell shape of aortic smooth muscle cells was evaluated by using phase-contrast microscopy (IM, Carl Zeiss, Oberkochen, Federal Republic of Germany) and a digital imaging analysis system (Zidas, Carl Zeiss). The digitizer was calibrated with a micrometer scale for each experiment. Cell surface area was measured before and 20 min after the addition of any agent. These measurements were done in triplicate on each cell. Repeated measurements of the same cell yielded an average coefficient of variation of ~1.5%. Taking into account the spontaneously occurring change of cell shape and the variability in the image analysis, only a decrease in cell surface area of > 15% was considered a positive contractile response. A ×20 objective was used for all contraction studies. The microscopic field was chosen randomly. At the time of the experiment, culture medium was aspirated and cells were washed twice with PSS. Cells were then incubated with 37°C PSS in the presence or absence of the effectors and cell shape change was evaluated at room temperature. In experiments with ANF, a 5 min preincubation was performed, as described previously.

Statistical analysis. Data are presented as mean±SEM and statistical analysis was performed with Student’s t test for paired and unpaired data. The Bonferroni correction was used for multiple comparisons. Results of the contraction studies were evaluated with the Wilcoxon rank sum test.

Results

Intracellular free Ca²⁺. The basal [Ca²⁺] in resting aortic smooth muscle cells using quin 2 was 152.3±9.4 nM (n = 27). Upon stimulation with AVP, there was a dose-dependent rapid increase in [Ca²⁺] (Fig. 1), which returned to basal levels within 5–7 min. APIII (10⁻⁸ M) alone did not affect the basal free Ca²⁺ levels (145.4±8.7 vs. 152.3±9.4 nM, NS) either when the cells were preincubated with APIII for 10 min or when
AVP-induced increases in \([\text{Ca}^{2+}]_i\). Points represent the mean of the maximal change in \([\text{Ca}^{2+}]_i\), expressed as percent increase over baseline. Each point represents the mean±SEM of 11–27 determinations in at least five separate cell cultures.

\([\text{Ca}^{2+}]_i\) was measured directly after addition of APIII. In the presence of \(10^{-7}\) M AVP, ANF \((10^{-8}\) M) had no effect on stimulated \([\text{Ca}^{2+}]_i\) \((436.3±21.7\) vs. \(418.4±19.1\) nM, NS). However, APIII \((10^{-8}\) M) significantly attenuated the increase in cytosolic free \([\text{Ca}^{2+}]_i\) induced by \(10^{-8}\) M AVP (Fig. 2) in a dose-dependent manner. The mean peak value of \([\text{Ca}^{2+}]_i\) with AVP \((10^{-8}\) M) was \(310.0±28.4\) nM; it decreased to \(234.0±14.8\) nM \((P < 0.01)\) in the presence of APIII \((10^{-8}\) M). No measurable lag time was observed before the onset of the effect of AVP in the presence of APIII. The following similar results were found in experiments measuring \([\text{Ca}^{2+}]_i\) using fura 2. Whereas APIII \((10^{-8}\) M, 5-min preincubation) did not affect the \([\text{Ca}^{2+}]_i\) increase by \(10^{-7}\) M AVP \((\Delta503.08±18.4\) vs. \(\Delta362.48±33.0\) nM, NS, \(n=3)\), \(10^{-7}\) M APIII significantly reduced the \([\text{Ca}^{2+}]_i\) peak by \(10^{-8}\) M AVP \((\Delta168.9±20.25 \text{ to } \Delta104.32±22.0\) nM, \(P < 0.025, n=8)\). Basal levels of \([\text{Ca}^{2+}]_i\) were not significantly affected by ANF \((86.6±5.7\) vs. \(80.5±8.3\) µM). In control experiments using angiotensin II (All) as the cell activator, no effect of \(5 \times 10^{-8}\) M APIII was found in the \([\text{Ca}^{2+}]_i\), peak produced by \(10^{-7}\) All \((\Delta857.17±82.7\) without ANF vs. \(\Delta954.08±54.3\) nM with ANF) and \(10^{-8}\) M All \((\Delta857.16±90.0\) without ANF vs. \(\Delta504.25±11.4\) nM with ANF).

\(45\text{Ca}^{2+}\) efflux. Preincubation (5 min) of the cells with \(10^{-8}\) M APIII did not affect the spontaneous \(45\text{Ca}^{2+}\) efflux (data not shown). AVP \((10^{-8}\) M) markedly increased \(45\text{Ca}^{2+}\) efflux within 30 s \((1.64±0.15\) vs. \(2.59±0.37 \times 10^3\) cpm/mg protein per 30 s, \(n=5, P < .005)\) (Fig. 3). The maximal increase was observed after 60 s. Basal efflux rates were reached again after about 3 min. In cells preincubated for 5 min with APIII \((10^{-8}\) M), the stimulatory effect of AVP \((10^{-8}\) M) was attenuated and only a slight insignificant increase was observed. APIII \((5 \times 10^{-9}\) M) significantly attenuated the response to AVP (Fig. 3). However, no effect of APIII \((10^{-8}\) M) on \(10^{-7}\) M AVP-mediated \(45\text{Ca}^{2+}\) efflux was detected \((4.69±0.41\) vs. \(4.51±0.33 \times 10^3\) cpm/mg protein per 30 s, \(n=5)\).

\(4\text{Ca}^{2+}\) uptake. In Fig. 4, the \(4\text{Ca}^{2+}\) uptake in response to AVP alone, APIII alone, and AVP plus APIII are depicted. The uptake of \(4\text{Ca}^{2+}\) by aortic smooth muscle cells was measured over a period of 5 min. AVP \((10^{-8}\) M) significantly stimulated \(4\text{Ca}^{2+}\) uptake above spontaneous influx from \(9.58±0.75 \text{ to } 13.04±0.92 \times 10^3\) cpm/mg protein per 5 min \((n=5, P < 0.01)\). APIII \((10^{-8}\) M) alone did not affect the spontaneous uptake, but preincubation with APIII returned the stimulatory effect of AVP to baseline (AVP \(13.04±0.92\) to AVP + ANF \(9.80±0.64 \times 10^3\) cpm/mg protein per 5 min, \(n=5, P < 0.01)\). APIII \((10^{-8}\) M) did not affect \(4\text{Ca}^{2+}\) uptake by \(10^{-7}\) M AVP (data not shown). APIII \((10^{-9}\) M) did not affect AVP-stimulated uptake significantly \((12.70±0.48\) vs. \(13.04±0.92 \times 10^3\) cpm/mg protein per 5 min, NS).

Vascular smooth muscle cell shape change. Addition of AVP to the medium-induced cell shape change in a dose-dependent manner (Fig. 5) (26). Cell shape change began at 3–5 min after the addition of AVP and was maximal after 20 min. These changes in shape were characterized by a rounding of the cell body and the retraction of cell processes. With \(10^{-8}\) M AVP, \(24.3±6.5\%\) of the cells changed in response to the hormone. This percentage increased to \(53.9±5.6\%\) with \(10^{-6}\) M AVP. The average decrease in cell surface area for different hormone concentrations varied between 30.7% and 34.1%; these differences were not significant and did not correlate with the concentration of AVP. APIII \((10^{-8}\) M) virtually abolished contraction induced by \(10^{-8}\) and \(10^{-7}\) M AVP. APIII \((5 \times 10^{-9}\) M) decreased the percentage of cells responding to \(10^{-7}\) M AVP from \(34.8±4.8\%\) to \(19.4±3.5\%\) \((P < 0.05)\) (Fig. 6).

Cyclic nucleotide production. The production of cGMP and cAMP in aortic smooth muscle cells was measured in the...
presence of the phosphodiesterase inhibitor MIX (0.5 mM) under basal conditions and in the presence of APIII. Basal cGMP production was 5.65±1.60 pmol/mg protein per 10 min and increased in a dose-dependent manner with APIII doses of 10^-9 to 10^-6 M (Fig. 7). Basal cAMP production was 16.45±2.69 pmol/mg protein per 10 min (n = 3) and was not stimulated by 10^-7 M APIII (16.32±2.36 pmol/mg protein 10 per min, n = 4).

**Effects of 8-bromo cGMP on AVP-induced Ca^2+ mobilization and cell shape change.** Since the effect of APIII to attenuate AVP-induced Ca^2+ mobilization and shape change in aortic smooth muscle cells was associated with increased cGMP production, the stable nucleotide 8-bromo cGMP (8bcGMP) was used to mimic the effects of APIII. Preincubation for 5 min with 8bcGMP (10^-3 and 5 x 10^-3 M) significantly attenuated AVP-stimulated increase (10^-8 M) of [Ca^2+]i (Fig. 8). 8-bcGMP (10^-7 M) also reduced the 10^-7 M AVP-induced increase of [Ca^2+]i by 45.2% (P < 0.05), n = 4. 10^-3 M 8bcGMP had no effect on basal [Ca^2+]i (data not shown). 8bcGMP also attenuated the effect of AVP on 45Ca^2+ efflux (Fig. 9). The AVP-induced 45Ca^2+ uptake was also diminished by 8bcGMP in a dose of 10^-3 M (13.04±1.42 vs. 11.21±0.94 x 10^3 cpm/mg protein, n = 6, P < 0.05) (Fig. 10). 8bcGMP alone did not affect spontaneous 45Ca^2+ efflux and uptake (data not shown). Furthermore, 8bcGMP (10^-3 M) attenuated AVP (10^-7 M)-induced cell contraction (20.4±0.9% vs. 32.9±2.3%, n = 5, 10-17 cells per experiment, P < 0.05).

**Methylene blue.** Methylene blue, an inhibitor of soluble guanylate cyclase, was used in an attempt to inhibit the effects of APIII on AVP-stimulated Ca^2+ mobilization and cell contraction. Methylene blue (10^-4 M) did not block APIII (10^-8 M)-stimulated cGMP production (54.0±7.8 vs. 55.1±12.3 pmol/mg protein per 10 min, n = 4, NS). Methylene blue also did not prevent the inhibitory effect of APIII on AVP-stimu-
Figure 5. Dose-response curve of vascular smooth muscle cells in the presence of different concentrations of AVP. Each bar represents the mean±SEM of at least five experiments.

Radiolabeled 45Ca2+ efflux (Fig. 8) or 45Ca2+ uptake by APIII (Fig. 9). Furthermore, methylene blue had no effect on the inhibition of cell contraction by APIII (data not shown).

Inositol phosphates. The effect of APIII (10^-7 M) on AVP-induced inositol phosphate's production is shown in Table I. APIII did not interfere with the AVP-mediated increase in IP2 and IP3, whereas a decrease in IP1 levels was observed (P < 0.05).

Discussion

The present in vitro studies were undertaken to examine whether APIII alters the AVP-induced cellular Ca2+ kinetics and cell contraction of rat vascular smooth muscle cells in culture. The vascular (V1) receptors for AVP are known to increase the formation of IP3, which stimulates the intracellular release of Ca2+ from the sarcoplasmic reticulum (27-29).

Figure 6. Effect of APIII on AVP (10^-7 M)-stimulated contraction of aortic smooth muscle cells (n = 5, 10-17 cells per experiment). Cells were preincubated with APIII for 10 min and APIII at the same concentration of the preincubation was added simultaneously with AVP.

This intracellular release of Ca2+ occurs in the absence of extracellular Ca2+ and is blocked by dantrolene, a known inhibitor of [Ca2+] transient mobilization (26, 30, 31). The IP2-induced Ca2+ release can be monitored by the rapid rise in cytosolic free Ca2+ as assessed by fluorescent dyes and the rapid cellular efflux of 45Ca2+. Both cytosolic Ca2+ and 45Ca2+ efflux measurements were used in the present study to examine whether APIII alters this initial phase of contraction induced by AVP.

APIII attenuated in a dose-response manner both the AVP-induced rise in cytosolic free Ca2+ and 45Ca2+ efflux. Quin 2 and fura 2 fluorescent dyes were used to assess cytosolic free Ca2+ in the present study and the results were similar. Other investigators have also shown in glomerular mesangial cells that results were similar when [Ca2+] was assessed by these same indicators (32). No effect of APIII alone could be demonstrated on either cytosolic free Ca2+ or 45Ca2+ efflux, results similar to those observed by other authors (33, 34). These in vitro data are compatible with the in vivo observations that the vasodilatory effect of the atrial peptides is most easily demonstrable in the presence of vasoconstrictor hormones (12). Also, ANF has been reported to interfere with Ca2+ mobilization and contraction by norepinephrine (34, 35), histamine (34), and AII (33), although for AII the evidence is not consistent (34, 36, 37).

The effect of APIII to decrease both the AVP-induced rise in cytosolic free Ca2+ and 45Ca2+ efflux suggested an inhibitory effect on either IP3 generation or the effect of IP3 to increase

Figure 7. APIII-stimulated cGMP production in aortic smooth muscle cells. Each column represents the mean of three to four determinations done in triplicate in separate cell cultures in the presence of 0.5 mM MIX.

Figure 8. Effect of 8-bromo cGMP on AVP-stimulated [Ca2+]. Each trace has been redrawn and represents a typical time course of [Ca2+], with the mean peak value (n = 5).
Ca\(^{2+}\) release from the sarcoplasmic reticulum. Even though we did not find a decrease in IP₃ production by AVP in cells pretreated with ANF, the observed decrease in IP₃, a metabolite of IP₃, suggests that an earlier decrease of IP₃ could have occurred. The issue of the role of ANF on inositol phosphates metabolism is still controversial. Experiments in glomerular mesangial cells did not detect an effect of APII to alter AVP-induced IP₃ generation (37). Although these latter results were obtained in mesangial, not aortic smooth muscle cells, and with APII not APIII, they suggest that atrial peptides may act by a mechanism distal to IP₃ production, e.g., the endoplasmic reticulum or the contractile apparatus. In this regard, it is interesting to note that cGMP, the purported secondary messenger of atrial peptides, has been proposed to alter sarcoplasmic Ca\(^{2+}\) release (38, 39) and to modify the kinetics of membrane Ca\(^{2+}\) transport (40). However, there are other reports showing that ANF (41) or its putative second messenger cGMP (42) may cause a decrease in inositol phosphatide breakdown. This could account for the reduction in Ca\(^{2+}\) release by AVP and other hormones.

Further studies were therefore performed in the present investigation to examine whether cGMP might be involved in these effects whereby APIII alters AVP-induced Ca\(^{2+}\) kinetics in rat aortic vascular smooth muscle. As in previous studies (15, 16), the present results support the findings that atrial peptides stimulate cGMP but not cAMP in vascular smooth muscle. The effect of APIII to stimulate cGMP was demonstrated to be dose-dependent and the 10⁻⁸ M dose of APIII that inhibited AVP-induced Ca\(^{2+}\) kinetics and cell contraction also increased cGMP. This observation therefore was compatible with the possibility that cGMP might be involved in the observed interactions of ANF and AVP on cellular Ca\(^{2+}\) kinetics.

**Figure 9.** \(^{45}\)Ca\(^{2+}\) efflux. Effect of 8-bromo cGMP on AVP-induced \(^{45}\)Ca\(^{2+}\) efflux and inhibition of AVP-induced efflux by APIII in the presence of methylene blue (four to nine experiments per group).

**Figure 10.** 8-bromo cGMP mimics the effect of APIII on AVP-induced \(^{45}\)Ca\(^{2+}\) uptake (crosshatched bar). Methylene blue does not affect inhibition of AVP-induced \(^{45}\)Ca\(^{2+}\) uptake by APIII (solid bar). Each bar represents the mean±SEM of four to six determinations done in duplicate in separate cell cultures.
Table I. Effect of ANF on AVP-mediated Inositol Phosphate Production

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<td>IP₁</td>
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n = 8.
* P < 0.05 respective to the same AVP concentration in the absence of ANF.

The effect of the stable cGMP analogue, 8bcGMP, on AVP-induced ⁴⁵Ca²⁺ efflux and cytosolic free Ca²⁺ was therefore examined. 8bcGMP decreased AVP-induced ⁴⁵Ca²⁺ efflux and cytosolic free Ca²⁺ concentration and cell shape change. The effective doses of 8bcGMP were, however, quite large, i.e., 1 and 5 mM, thus a pharmacological rather than physiological effect of the agent cannot be excluded. The concentrations of the 8bcGMP at the intracellular target sites may, however, have been in the normal range for cGMP.

Whereas some investigators have suggested that soluble guanylate cyclase may be responsible for the effect of atrial peptides to stimulate cGMP (43, 44), other results suggest that particulate guanylate cyclase is involved (16, 44). In the present studies, the absence of an effect of methylene blue, an inhibitor of soluble guanylate cyclase, suggests that, if cGMP modulates the interaction between APIII and AVP, it is the particulate rather than soluble guanylate cyclase which is involved. However, the possibility of the existence of non-cGMP-mediated effects of AVP must be taken into account, as suggested by recent reports (46).

In the present study APIII was also shown to diminish the effect of AVP to increase ⁴⁵Ca²⁺ uptake in a dose-response manner. Thus, while 10⁻⁸ M APIII blocked the cellular uptake induced by 10⁻⁸ M AVP, 10⁻⁹ M APIII did not block this effect of the same dose of AVP. Such reduction in ⁴⁵Ca²⁺ uptake may account in part for the decreased [Ca²⁺], peak observed with the fluorescent indicators. Taylor and Meisneri (47) reported that APIII inhibits norepinephrine-induced ⁴⁵Ca²⁺ uptake by aortic rings. Furthermore, these investigators found that APIII is much less effective in inhibiting potassium-induced ⁴⁵Ca²⁺ uptake thus suggesting that atrial peptides may interfere primarily with receptor-operated rather than voltage-dependent channels.

The apparent discrepancy between the concentrations of ANF that inhibit cell shape change and those that reduced Ca²⁺ mobilization is suggestive of an ANF action beyond the Ca²⁺-releasing mechanisms. Similar results have been recently reported by Takuwa and Rasmussen (34), who found a strong inhibitory effect of ANF on the AVP-dependent rat aortic contraction, in the absence of a decrease in AVP-stimulated Ca²⁺ transient. As a control in the present study, the effect of ANF on AVP-induced [Ca²⁺] was examined and these previous results (33) were confirmed. Furthermore, the partial reduction by ANF of the AVP-mediated [Ca²⁺] increase was unlikely to explain totally the inhibition of cell shape change with ANF. Specifically, the [Ca²⁺] transient induced by AVP in the presence of ANF, although reduced, was still in the range which is associated with cell activation.

In summary, ANF may interfere with the initial phase of AVP-induced vascular contraction by diminishing cellular Ca²⁺ mobilization, perhaps by impairing the effect of IP₃ on the endoplasmic reticulum. The present results, however, also incriminate another effect(s) of ANF to block vascular contraction caused by AVP. This latter effect of ANF appears to be different than the initial Ca²⁺ mobilization by IP₃, and may involve protein kinase activation and diminished cellular Ca²⁺ influx.

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