Angiotensin II enhances noradrenaline release from sympathetic nerves of the rat prostate via a novel angiotensin receptor: implications for the pathophysiology of benign prostatic hyperplasia

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Abstract

The renin–angiotensin system (RAS) is present in the human prostate and may be activated in benign prostatic hyperplasia (BPH), possibly contributing to the pathophysiology of this disorder by enhancing local sympathetic tone and cell growth. The functional role of the RAS in the prostate, however, is unknown. The present study was undertaken to determine whether angiotensin (Ang) II enhances sympathetic transmission in the prostate. The neuronal stores of the rat prostate were labelled with [³H]noradrenaline (NA). Ang II and Ang I enhanced [³H]NA release in a concentration-dependent manner. The Ang II receptor subtype 1 (AT₁ receptor) antagonist losartan and the AT₂ receptor antagonist PD123319 inhibited this facilitatory effect of Ang II and Ang I, whereas the other AT₂ receptor antagonist, CGP42112, was without effect. Bradykinin also increased [³H]NA release, which was inhibited by the B₂ receptor antagonist Hoe140. The angiotensin–converting enzyme inhibitor captopril inhibited the effect of Ang I, but potentiated that of bradykinin. Interestingly, captopril alone produced an increase in [³H]NA release which was inhibited by Hoe140. Losartan, but not PD123319 or CGP42112, inhibited [¹²⁵I]-Ang II binding in Chinese hamster ovary cells transfected with the AT₁a or AT₁b receptor. In contrast, in cells expressing the AT₂ receptor, PD123319 and CGP42112, but not losartan, inhibited [¹²⁵I]-Ang II binding. In conclusion, Ang II enhances the release of NA from sympathetic nerves of the rat prostate via a novel functional receptor distinct from the cloned AT₁a, AT₁b or AT₂. These data provide direct evidence in support of a functional role for the local RAS in modulating sympathetic transmission in the prostate, which may have important implications for the pathophysiology of BPH.

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Introduction

Benign prostatic hyperplasia/hypertrophy (BPH) is a highly prevalent disorder that represents the most common cause of urinary obstruction in the ageing male population and frequently co-exists with hypertension (Garraway et al. 1991, Maruenda et al. 1999). BPH is characterised by increased cellular proliferation of stromal elements and enhanced sympathetic smooth muscle tone (Isaacs & Coffey 1989, McNee 1990, Madsen & Bruskewitz 1995). In patients with BPH, approximately 50% of total urethral pressure is due to increased sympathetic smooth muscle tone, which can be ameliorated by α₁-adrenoceptor blockers such as prazosin, terazosin or doxazosin (Furuya et al. 1982, Kirby 1989, Madsen & Bruskewitz 1995). The prostate is innervated by sympathetic nerves which, upon stimulation, release the chemical transmitter noradrenaline (NA) and evoke smooth muscle contraction, mediated by α₁-adrenoceptors of the α₁A (α₁D) subtype (Forray et al. 1994, Marshall et al. 1995). Moreover, there is some evidence to suggest that local sympathetic activity may also influence prostate growth (McVary et al. 1994). For example, it has been reported that unilateral sympathectomy of the lower urinary tract reduces ventral prostate weight in rats (McVary et al. 1994). Thus enhanced sympathetic activity is a major factor in the development of BPH, influencing smooth muscle tone and, possibly, growth of the prostate.

The renin–angiotensin system (RAS) is a hormonal cascade that has an important role in regulating blood pressure and cardiovascular homeostasis (Peach 1977, Johnston 1990). Similarly, hyperactivity of the RAS has been implicated in the development of hypertension and other cardiovascular disorders (Nicholls et al. 1998).
Angiotensin II (Ang II) is the principal effector peptide of the RAS and is formed by the sequential cleavage of the precursor macromolecule, angiotensinogen, to the inactive decapeptide Ang I by renin, and then hydrolysed to the active octapeptide Ang II by angiotensin–converting enzyme (ACE) (Johnston 1990). It should be noted, however, that ACE is not a specific enzyme and can degrade bradykinin and a host of other peptides to inactive fragments (Fabiani et al. 2000). In addition to its formation within the circulation, Ang II can also be generated locally in many target tissues such as the kidney, heart, brain and blood vessels, and may therefore also mediate autocrine or paracrine effects (Campbell 1987, Dzau 1988, Johnston 1992).

The effects of Ang II are subserved by at least two distinct receptor subtypes, denoted AT1 and AT2 (Griendling et al. 1996, Unger et al. 1996, Fabiani 1999). In rodents, but not higher species or humans, two further isoforms of the AT1 receptor have been identified and termed AT1a and AT1b (Iwai & Inagami 1992, Yoshida et al. 1998). The characterisation of Ang II receptors into two major classes was promulgated by the development of selective non-peptide antagonists such as losartan and PD123319 (Timmermans et al. 1993). Ang II receptors sensitive to losartan were designated AT1, whereas those sensitive to PD123319 were designated AT2 (Timmermans et al. 1993). Ang II exerts a variety of biological effects that serve to modulate cardiovascular function and structure, including vasoconstriction, stimulation of aldosterone release and promotion of cell growth/hypertrophy, all of which are mediated by the AT1 receptor (Chung et al. 1998, Fabiani 1999). The functional role of the AT2 receptor is less well understood, but may be involved in anti-proliferation, apoptosis, differentiation and, possibly, vasodilatation (Chung et al. 1998, Csikos et al. 1998, Fabiani 1999).

Of particular note, the RAS can interact with the sympathetic nervous system in a stimulatory manner at several different levels of the neuronal network (Saxena 1992). Ang II is well known to facilitate the release of NA from sympathetic nerve terminals in many tissues including the heart, kidney and blood vessels (Story & Ziogas 1987). Moreover, Ang II is able to amplify the post-junctional actions of NA and other excitatory mediators on cardiac and smooth muscle cells (Purdy & Weber 1988). This facilitatory effect on sympathetic transmission represents one of the most potent actions of Ang II, usually requiring concentrations that are much lower than those required to produce direct vasoconstriction or cardiac inotropic and chronotropic effects (Story & Ziogas 1987).

Despite the known role of sympathetic activity in BPH, as described above, the functional role of the RAS in the prostate is unknown. Furthermore, although it was reported several years ago that the biochemical activity of ACE is significantly greater in BPH than in normal prostate or a variety of other tissue homogenates (Yokoyama et al. 1980, Van Sande et al. 1985), no further work has since been undertaken to investigate the physiological or pathophysiological role of the RAS in the prostate. Our group has shown more recently that both the protein and mRNA expression of ACE (Nassis et al. 2000, 2001), in addition to Ang II immunoreactivity (Dinh et al. 2001a), are increased in BPH. Furthermore, we have observed that AT1 receptors predominate in the human prostate and are down-regulated in BPH (Dinh et al. 2001a,b). Taken together, these findings suggest that the local RAS is activated in BPH, which may have pathophysiological consequences.

Nothing is known about the functional interaction of the RAS with sympathetic nerves in the prostate. We hypothesise that a local tissue-based RAS is functionally active and facilitates sympathetic neuroeffector transmission in the prostate, such that hyperactivity of the local RAS contributes to the development of BPH by enhancing local sympathetic activity. In view of the possibility that the RAS may be activated in BPH, the present study was undertaken to determine whether Ang II, the main effector molecule of the RAS, facilitates sympathetic transmission in the prostate. Specifically, we aimed to examine the effects of exogenous and locally generated Ang II on NA release from sympathetic nerves of the rat prostate and endeavoured to characterise the Ang II receptor subtype(s) involved.

Materials and Methods

Rat prostate preparation

Male Sprague–Dawley rats (250–350 g) were killed by decapitation and then exsanguinated. An incision was made in the lower abdominal region and the prostate identified and dissected free. The isolated prostate was transferred to a Petri dish containing pre-warmed physiological salt solution (PSS) and continuously gassed with carbogen (95% O2–5% CO2). Excess fatty tissue was removed and the prostate cut into two even sections. Each piece of prostate tissue was tied at both ends with fine silk threads.

Radiolabelling of noradrenergic transmitter stores with [3H]NA

The procedure used to radiolabel the noradrenergic transmitter stores was adopted from Fabiani & Story (1994, 1996). Prostate tissues were placed in small glass-jacketed organ baths and equilibrated for 10 min in 2 ml PSS, maintained at 37 °C and continuously gassed with carbogen. After this initial equilibration period, 5 μl [3H]NA (30–50 Ci/mmol) was added to each organ bath and the preparations allowed to incubate for 30 min. After incubation, prostate tissues were removed from the organ baths and immersed in a small volume of PSS to rinse any loosely bound radioactivity. Prostate preparations were then
mounted vertically between two platinum electrodes in acrylic flow chambers and superfused with PSS at a rate of 2 ml/min using an ISCO Wiz peristaltic pump (ISCO Inc., Lincoln, NE, USA). In order to further remove loosely bound radioactivity, the tissues were washed for 90 min before experimental procedures were commenced. After the first 30 min of the washout period, the preparations were subjected to a 30 s period of electrical field stimulation with a train of 1 ms monophasic square-wave pulses at a frequency of 5 Hz and a supramaximal voltage of 20 V, delivered by a Grass S88 stimulator (Grass Medical Instruments, Quincy, MA, USA). This brief ‘priming stimulus’ was intended to assist in the removal of any non-specifically bound radioactive material.

Stimulation of intrinsic sympathetic nerves
After the washout period, the intrinsic sympathetic nerves of the prostate preparations were subjected to two 60 s periods of electrical field stimulation (1 ms pulses, 5 Hz, 20 V). The first period of stimulation was given immediately after the 90 min washout period and the second period of stimulation given 30 min later. The effects of Ang II, Ang I and various drugs on the resting and stimulation-induced effluxes of radioactivity were examined by adding the drugs to the PSS superfusing the prostate preparations 15 min before the second period of stimulation. The drugs then remained present for the duration of the experiment.

Determination of resting and stimulation-induced effluxes of radioactivity
The superfusate from the prostate preparations was collected at 3 min intervals by an automated ISCO Retriever IV fraction collector (ISCO Inc.). Each 3 min (6 ml) fraction of superfusate was mixed with 4 ml of scintillant (Ultima Gold, Packard Bioscience BV, Groningen, The Netherlands) and the radioactivity present determined by liquid scintillation counting. External automatic standardisation was utilised to correct for counting efficiency and the data were expressed in disintegrations per minute (d.p.m.).

The resting efflux of radioactivity from the prostate preparations was determined for each of the two periods of stimulation (R1 and R2) from the amount of radioactivity present in the fraction of superfusate collected immediately before stimulation. The stimulation-induced efflux of radioactivity for each of the two periods of stimulation (S1 and S2) was determined by subtracting the resting efflux from the amount of radioactivity present in each of the fractions of superfusate collected from the commencement of stimulation, and summing the differences:

\[
S_1 = (F_1 + F_2 + F_3 + \ldots) - nR_1 \\
S_2 = (F'_1 + F'_2 + F'_3 + \ldots) - nR_2
\]

where F and F’ represent the radioactive content in each fraction collected after the first and second period of stimulation respectively, R1 and R2 represent the resting efflux for the first and second period of stimulation respectively and n represents the number of fractions. In each experiment, the resting and stimulation-induced effluxes for the second period of stimulation were expressed as percentages of the corresponding values for the first period of stimulation (% R2/ R1 and % S2/S1 respectively).

Receptor constructs, expression and binding assay
The cloning of the rat AT1a, receptor and its incorporation into the mammalian expression vector, pRc/CMV, have been described previously (Thekkumkara et al. 1995). The rat AT1b receptor, sub-cloned into pRc/CMV, was kindly provided by Dr T J Thekkumkara (Department of Medicine, University of Colorado, Denver, CO, USA). The rat AT2 cDNA was kindly provided by Dr T J Murphy (Department of Pharmacology, Emory University School of Medicine, Atlanta, GA, USA) and subsequently sub-cloned into pRc/CMV using unique 5’ (HindIII) and 3’ (Xbal) restriction sites.

Chinese hamster ovary (CHO-K1) cells were transfected in 12-well plates with 0·6 µg/well of either AT1a, AT1b or AT2 receptor plasmid DNA using lipofectAMINE (4·8 µl/well), as previously described (Thomas et al. 1998). At 48 h post-transfection, competition radioreceptor-binding assays were performed using 0·05 nM [125I]-Ang II as tracer and increasing concentrations of Ang II, losartan (an AT1-selective ligand), PD123319 (an AT2-selective ligand) or CGP42112 (an AT2-selective ligand). Non-linear regression of the data was achieved using GraphPad Prism (Graphpad Software Inc., San Diego, CA, USA) and Kd and Bmax were determined by the method of Swillens (1992).

Drugs, solutions and radiochemicals
The following drugs were used: angiotensin II (AUSPEP Pty Ltd, Parkville, VIC, Australia; Sigma Chemical Co., St Louis, MO, USA); angiotensin I (AUSPEP Pty Ltd; Sigma Chemical Co.); ß-conotoxin GVIA (AUSPEP Pty Ltd); tetrodotoxin (Sigma Chemical Co.); losartan (Merck & Co Inc., Rathway, NJ, USA), PD123319 (Research Biochemicals International, Natick, MA, USA), captopril (Sigma Chemical Co.), CGP42112 (Novartis Pharma AG, Basel, Switzerland), idazoxan (Research Biochemicals International), atropine (Sigma Chemical Co.), bradykinin (Sigma Chemical Co.) and Hoe140 (Hoechst AG, Frankfurt, Germany).

Stock solutions of indomethacin were made in 2 M Na2CO3. All other stock solutions and intermediate dilutions of drugs were made in distilled water. Final concentrations of drugs were achieved by the appropriate
dilution in PSS, continuously gassed with carbogen and maintained at a temperature of 37 °C.

The PSS had the following composition (mM): NaCl, 118; KCl, 4-7; CaCl2, 2-5; MgSO4, 0-45; NaHCO3, 25; KH2PO4, 1-03 and d-(+)-glucose, 11-1. EDTA (0-067 mM) and ascorbic acid (0-14 mM) were also present to prevent oxidation of NA.

(t)-[Ring 7, 8-3H]-noradrenaline (specific activity 30–50 Ci/mmol) was supplied by Amersham Pharmacia Biotech Pty Ltd (Buckinghamshire, England, UK).

Statistical analysis of results
Results are expressed as mean±s.e.m.; n represents the number of experiments. The levels of statistical significance of differences were determined by one-way analysis of variance (ANOVA) followed by Dunnett’s test or Student–Newman–Keuls (SNK) test, where appropriate. All statistical analyses were performed using the statistical program SigmaStat for Windows (Jandel Scientific Software Inc., San Rafael, CA, USA). In all cases, probability levels less than 0-05 (P<0-05) were taken to indicate statistical significance.

Results
Effects of exogenous and locally-generated Ang II on [3H]NA release from rat prostate
Control experiments Figure 1 shows the mean content of radioactivity (d.p.m.) in consecutive 3 min fractions of superfuse collected from control rat prostates (n=6) loaded with [3H]NA and subjected to two periods of stimulation. The mean resting efflux of radioactivity for the first period of electrical field stimulation (R1) was 1414±134 d.p.m. per 3 min collection (n=6). There was a gradual decrease in the resting efflux between the two periods of stimulation, such that the resting efflux preceding the second period of stimulation, expressed as a percentage of the first (%R2/R1), had a mean value of 80-1±1-0%. Electrical field stimulation evoked an increase in the radioactive efflux from prostate preparations loaded with [3H]NA. The mean stimulation-induced efflux evoked by the first period of stimulation (S1) was 2954±587 d.p.m. (n=6). The mean value of the stimulation-induced efflux for the second period of stimulation, expressed as a percentage of the first (%S2/S1), was 87-0±3-0%.

Characterisation of stimulation-induced efflux To confirm that the source of radioactivity released from the radiolabelled prostate by electrical field stimulation was due to the neuronal exocytotic release of [3H]NA from sympathetic nerves, the effects of tetrodotoxin (neuronal Na+ channel blocker), ω-conotoxin (neuronal N-type Ca2+ channel blocker) and Ca2+-free PSS on stimulation-induced efflux were investigated. Tetrodotoxin (1 µM), ω-conotoxin (0-1 µM) and the removal of extracellular Ca2+ from the PSS, introduced 15 min before the second period of stimulation, virtually abolished the stimulation-induced efflux of radioactivity (Fig. 2; P<0-05). The selective α2-adrenoceptor antagonist idazoxan was used to determine whether transmitter NA release was subject to auto-inhibition mediated by prejunctional α2-adrenoceptors. Idazoxan (0-1 µM), introduced into the PSS 15 min before the second period of stimulation, markedly enhanced stimulation-induced efflux, by approximately 170% above control (Fig. 2; P<0-05).

Effects of Ang II and Ang I As shown in Fig. 3, introduction of exogenous Ang II (0-001–1 µM) into the PSS, 15 min before the second period of stimulation, produced a concentration-dependent increase in the stimulation-induced efflux of [3H]NA, with a maximal effect at 0-1 µM (P<0-05). The precursor molecule Ang I was used to determine whether locally generated Ang II could also modulate transmitter NA release in the prostate. Similar to Ang II, Ang I (0-001–1 µM), introduced 15 min before the second period of stimulation, also enhanced the stimulation-induced efflux in a concentration-dependent manner, but was approximately 10-fold less potent than Ang II, with a maximal effect at 1 µM (Fig. 3; P<0-05). The maximal increase in stimulation-induced efflux produced by both Ang II and Ang I was approximately 40% above control.
Effects of Ang II and Ang I in the presence of losartan The Ang II receptor antagonists losartan, PD123319 and CGP4112 were used to characterise the angiotensin receptor subtype(s) mediating the effects of Ang II and Ang I on stimulation-induced [3H]NA release. When introduced alone 15 min before the second period of stimulation, the AT1 receptor antagonist losartan had no effect on stimulation-induced efflux. However, when introduced 15 min before the second period of stimulation in combination with Ang II (0·1 µM) or Ang I (1 µM), losartan (0·01 and 0·1 µM) significantly inhibited the increase in stimulation-induced efflux produced by Ang II and Ang I, in a concentration-dependent manner (Fig. 4; P<0·05).

Effects of Ang II and Ang I in the presence of PD123319 When introduced alone 15 min before the second period of stimulation, the AT1 receptor antagonist PD123319 had no effect on the stimulation-induced efflux. However, as seen with losartan, when introduced in combination with Ang II (0·1 µM) or Ang I (1 µM) 15 min before the second period of stimulation, PD123319 (0·01 and 0·1 µM) also inhibited the increase in stimulation-induced efflux produced by Ang II and Ang I, in a concentration-dependent manner (Fig. 5; P<0·05).

Effects of Ang II and Ang I in the presence of CGP42112 When introduced alone 15 min before the second period of stimulation, the AT2 receptor ligand CGP42112, like PD123319 or losartan, had no effect on the stimulation-induced efflux. However, in contrast to PD123319 or losartan, CGP42112 (0·01 and 0·1 µM), introduced in combination with Ang II (0·1 µM) or Ang I (1 µM), did not alter the facilitatory effect of Ang II or Ang I on stimulation-induced efflux (Fig. 6; P>0·05).

Effects of Ang I in the presence of captopril The ACE inhibitor captopril was used to determine whether the effects of Ang I on stimulation-induced efflux were due to generation of Ang II via an ACE-dependent pathway. When introduced alone 15 min before the second period of stimulation, captopril (3 µM) produced a small but significant increase in stimulation-induced efflux (Fig. 7; P<0·05). However, when introduced in combination with Ang I (1 µM) 15 min before the second period...
of stimulation, captopril (3 µM) markedly inhibited the facilitation of stimulation-induced efflux by Ang I (Fig. 7; P<0·05).

**Effects of bradykinin in the absence and presence of Hoe140 or captopril** The potential effects of bradykinin on transmitter NA release in the prostate were also investigated. Bradykinin (1 µM), introduced 15 min before the second period of stimulation, produced a significant increase in the stimulation-induced efflux, approximately 35% above control (Fig. 7; P<0·05). The selective B2 receptor antagonist Hoe140 was used to characterise the bradykinin receptor subtype involved in mediating the effects of bradykinin on stimulation-induced efflux. When introduced alone 15 min before the second period of stimulation, Hoe140 had no effect on the stimulation-induced efflux. However, when introduced in combination with bradykinin (1 µM) 15 min before the second period of stimulation, Hoe140 (0·3 and 1 µM) significantly inhibited the facilitatory effect of bradykinin (1 µM) on the stimulation-induced efflux (Fig. 7; P<0·05).

The ACE inhibitor captopril was used again to determine whether exogenously applied bradykinin is subject to degradation by ACE in the prostate. As shown in Fig. 7, in the presence of captopril (3 µM), introduced 15 min before the second period of stimulation, bradykinin further increased the stimulation-induced efflux, to approximately 50% above control and about 15% greater than that achieved with bradykinin alone (P<0·05). Interestingly, when introduced in combination with captopril (3 mM) 15 min before the second period of stimulation, Hoe140 (1 µM) inhibited the increase in stimulation-induced efflux produced by captopril (3 µM) alone (Fig. 7; P<0·05).
Effects of drugs on resting effluxes Addition of all the various drugs tested had little or no effect on the mean values of resting efflux (% R2/R1) (data not shown).

Competition displacement of [125I]-Ang II binding from the cloned rat AT1a, AT1b or AT2 receptor in CHO-K1 cells

In order to characterise further the receptor subtype mediating the effects of Ang II on NA release, binding studies were also performed in CHO cells expressing each of the cloned rat AT1a, AT1b and AT2 receptor. Figure 8 shows competition displacement curves of [125I]-Ang II binding to the cloned rat AT1a, AT1b and AT2 receptor expressed in CHO-K1 cells (three experiments performed in triplicate). The AT1a receptor displayed a Kd of 1.8±0.3 nM for Ang II and a receptor density (Bmax) of 1066±356 fmol/mg protein. The Kd and Bmax for the AT1b receptor were 4.9±1.4 and 571±160 fmol/mg protein respectively, whereas those for the AT2 receptor were 3.3±0.7 nM and 1254±301 fmol/mg protein. Displacement of [125I]-Ang II by losartan was observed for the AT1a and AT1b receptors, but not the AT2 receptor, whereas PD123319 and CGP 42112 displaced bound [125I]-Ang II from the AT2 receptor, but not the AT1a or AT1b receptor.

Discussion

Recent evidence from our laboratory suggests that the local RAS is activated in BPH. It is well documented that Ang II, the bioactive product of the RAS, facilitates sympathetic transmission in many cardiovascular organs by enhancing the release of the chemical transmitter NA from sympathetic nerve terminals. However, nothing is known about the interaction of the RAS with sympathetic neurotransmission in the prostate, which may possibly be implicated in the pathophysiology of BPH. Therefore, the
present study was undertaken to determine whether exogenous and locally generated Ang II enhances transmitter NA release in the rat prostate, in an effort to ascribe a functional role of the local RAS in modulating sympathetic activity in the prostate.

Radiotracer techniques have been used widely and extensively to monitor transmitter NA release in many sympathetically innervated tissues. The noradrenergic transmitter stores of the rat prostate were labelled with $[^3H]$NA, as described previously (Fabiani & Story 1994, 1996). Electrical field stimulation evoked an increase in the efflux of radioactivity from the radiolabelled prostate preparations. The neuronal Na$^+$ channel blocker tetrodotoxin, the neuronal N-type Ca$^{2+}$ channel blocker $\gamma$-conotoxin, and the removal of extracellular Ca$^{2+}$ each abolished the stimulation-induced efflux of radioactivity.

This confirms that the stimulation-induced efflux is caused by the neuronal exocytotic release of $[^3H]$NA by a mechanism dependent on the influx of Ca$^{2+}$ through N-type Ca$^{2+}$ channels. It is well established that sympathetic nerve terminals are endowed with $\alpha_2$-adrenoceptors that subserve auto-inhibition of transmitter NA release (Starke 1987, Rand et al. 1990, Fuder & Muscholl 1996).

In the present study, the $\alpha_2$-adrenoceptor antagonist idazoxan markedly enhanced stimulation-induced $[^3H]$NA release in the rat prostate, which is consistent with blockade of prejunctional inhibitory $\alpha_2$-adrenoceptors on sympathetic nerve terminals, and similar to that observed in other tissues, such as rat atria (Loiacono et al. 1985, Story et al. 1985), human atria (Abadie et al. 1996), rat kidney (Rump et al. 1990), rat mesenteric artery (Fabiani & Story 1996) and rabbit ear artery (Loiacono et al. 1985). These initial findings suggest that the stimulation-induced efflux from the radiolabelled rat prostate is caused by the evoked exocytotic release of $[^3H]$NA from sympathetic nerves and is subject to auto-inhibition mediated by prejunctional $\alpha_2$-adrenoceptors. Thus the stimulation-induced efflux may be taken as an index of transmitter NA release from sympathetic nerves of the rat prostate.

It is generally accepted that one of the most important actions of Ang II on sympathetic neurotransmission is the prejunctional facilitation of transmitter NA release (Story & Ziogas 1987, Saxena 1992). In the present study, Ang II, over a concentration range of 0.001–1 µM, enhanced the stimulation-induced efflux of $[^3H]$NA in a concentration-dependent manner, with a maximal effect at 0.1 µM, of approximately 35% above control. The facilitatory effect of Ang II on stimulation-induced $[^3H]$NA release in the rat prostate is consistent with findings in other tissues such as the rat caudal artery (Cox et al. 1995, 1996a,b), rat and human kidney (Rump et al. 1990, 1995), guinea-pig (Brasch et al. 1993) and human atria (Rump et al. 1994, Abadie et al. 1996).

Many tissues, including blood vessels, heart, kidney, adrenals and brain, are capable of generating Ang II locally.
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In the present study, the precursor molecule Ang I was therefore used to determine whether locally generated Ang II could modulate transmitter NA release in the prostate. Similar to Ang II, the inactive precursor Ang I also enhanced the [3H]NA release in a concentration-dependent fashion but was approximately 10-fold less potent than Ang II. The facilitatory effect of Ang I on [3H]NA release in the rat prostate was markedly inhibited by the ACE inhibitor captopril, indicating that the effects of Ang I are due to local conversion to Ang II via an ACE-dependent pathway. This finding suggests that a tissue-based RAS is indeed present and functionally active in the rat prostate and capable of generating Ang II locally. Facilitation of [3H]NA release by Ang I in the rat prostate is consistent with other studies in guinea-pig atria (Ziogas et al. 1984, Brasch et al. 1993), rat kidney (Böke & Malik 1983, Rump et al. 1990), human kidney (Rump et al. 1995), and rat vena cava (Ziogas & Story 1991), which was reportedly caused by locally generated Ang II.

We also endeavoured to characterise the Ang II receptor subtype(s) mediating the actions of Ang II on sympathetic transmission in the rat prostate. Previous studies suggest that, in a number of tissues from different species such as guinea-pig atria (Brasch et al. 1993), rat atria (Gironacci et al. 1994), rat trachea (Boicos et al. 1996), human kidney (Rump et al. 1995) and various mouse tissues (Cox et al. 1999), the facilitatory effect of Ang II on transmitter NA release is mediated by AT1 receptors, as they were inhibited by losartan and unaffected by PD123319. However, in the present study, the increase in [3H]NA release by both Ang II and Ang I in the rat prostate was significantly inhibited, in a concentration-dependent manner, by the selective AT1 receptor antagonist losartan in addition to the AT2 receptor antagonist PD 123319, whereas the other AT2 receptor antagonist, CGP42112, had no effect. These findings suggest that the receptor mediating the effects of exogenous and locally generated Ang II on NA release in the rat prostate involves a receptor that is sensitive to both losartan and PD123319. Similar findings have also been reported by Cox et al. (1995, 1996a,b) who showed that the facilitation of [3H]NA release by Ang II in the rat caudal artery was also inhibited by both losartan and PD123319.

The other AT2 receptor antagonist, CGP42112, did not alter the facilitatory effect of Ang II and Ang I on [3H]NA release, which suggests that AT2 receptors may not be involved. Cloning studies have revealed the existence of at least two subtypes of the AT1 receptor in rodents (rats and mice), but not higher species or humans, which have been designated AT1a and AT1b (Iwai & Inagami 1992, Yoshida et al. 1992). Interestingly, it has been reported in rat mesangial cells that two distinct AT1 receptor binding sites exist, which are coupled to activation of phospholipase C-mediated intracellular Ca2+ mobilisation and inhibition of adenylate cyclase, and sensitive to inhibition by losartan and PD123319 but not CGP42112 (Ermsberger et al. 1992, Zhou et al. 1993, Madhun et al. 1993). The AT1 binding site that displayed a greater affinity for losartan was denoted AT1A, and the other binding site, which displayed a greater affinity for PD123319, was denoted AT1B (here, subscript A and B denote the pharmacologically characterised receptors, as opposed to the cloned receptors denoted by subscript a and b).

To delineate further the Ang II receptor subtype involved in mediating the effects of Ang II on transmitter NA release in the rat prostate, Ang II binding studies were performed in CHO cells expressing each of the cloned rat Ang II receptors (AT1a, AT1b and AT2) to establish their sensitivity to inhibition by losartan, PD123319 and CGP42112. As expected, losartan inhibited Ang II binding in a concentration-dependent manner in CHO cells transfected with either the AT1A or AT1B receptor, whereas PD123319 and CGP42112 had little or no effect. Conversely, PD123319 and CGP42112 inhibited Ang II binding in CHO cells expressing the AT2 receptor and losartan had little or no effect. As the functional receptor mediating the effects of Ang II on NA release in the rat prostate was sensitive to inhibition by both losartan and PD123319 but not CGP42112, this receptor subtype does not appear to correspond to the cloned AT1A, AT1B or AT2 as determined by binding studies in CHO cells. We therefore propose that the functional receptor mediating the effects of Ang II on sympathetic transmission in the rat prostate may be similar to the functional AT1B Receptor as described above (Ermsberger et al. 1992, Zhou et al. 1993, Madhun et al. 1993, Cox et al. 1995, 1996a,b), which is pharmacologically distinct from the cloned rat AT1B receptor.

We have observed in other studies the overwhelming presence of typical AT1 receptors in both the human (Dinh et al. 2001a,b) and rat prostate (unpublished observations). Given that several isoforms of the AT1 receptor exist in the rat but not in the human, coupled with the apparent absence of AT2 receptors in the rat prostate, the functional AT1 receptor mediating the effects of Ang II on NA release in the rat prostate would appear to represent a novel subtype of the AT1 receptor that is uniquely sensitive to both losartan and PD123319 but not to CGP42112. This novel prejunctional AT1 receptor mediating the effects of Ang II on NA release in the rat prostate represents only a fraction of the total AT receptor pool in the rat prostate. Furthermore, given that the human prostate expresses almost exclusively AT1 receptors but not AT2 receptors, and that humans are not known to express any isoforms of the AT1 receptor, it seems highly likely that the potential effects of Ang II on NA release in the human prostate would also be mediated by the AT1 receptor, which is still consistent with the findings in the rat prostate. Thus the characterisation of this functional AT1B receptor subtype mediating the effects of Ang II.
on NA release in the rat prostate represents a unique phenomenon in the rat and might not be pertinent in the human prostate, as humans are not known to contain any isoforms of the AT1 receptor. Unfortunately, because of the unavailability of fresh and functionally viable human prostate tissues, it is not possible also to conduct NA release studies in the human prostate in order to compare directly the effects of Ang II on NA release in the rat prostate, and the receptor subtype involved, with that in the human prostate. Nonetheless, the findings of the present study provide direct evidence that Ang II facilitates NA release in the rat prostate, which may have direct implications for the pathophysiology of BPH in humans (see below).

Very recently, it was reported that AT1 and bradykinin B2 receptors form stable heterodimers in vascular smooth muscle cells and that this association dramatically affects the signalling and regulation of the AT1 receptor (AbdAlla et al. 2000). Whether AT1 and B2 receptors dimerise and cross-modulate in the prostate remains to be determined, but such a possibility may also be responsible for the changes in AT1 receptor pharmacology and functionality observed.

ACE has broad substrate specificity which, in addition to converting Ang I to Ang II, also degrades bradykinin to inactive peptide fragments (Fabiani et al. 2000). In the present study, captopril itself produced a significant increase in [3H]NA release. The facilitatory effect of captopril on transmitter NA release in the rat prostate may potentially be the result of accumulation of bradykinin as a consequence of impaired bradykinin degradation after ACE inhibition. Indeed, the observed increase in [3H]NA release in the rat prostate was inhibited by the bradykinin B2 antagonist Hoe140, suggesting that this facilitatory effect of captopril was mediated by bradykinin. Bradykinin has been shown to enhance transmitter NA release in a variety of tissues, including rat kidney (Böke & Malik 1983) and human kidney (Rump et al. 1995), rat and mouse vas deferens (Llona et al. 1991), rat atria (Chulak et al. 1995) and human atria (Rump et al. 1997), and rat hypothalamus (Tsuda et al. 1993). Consistent with these observations, bradykinin enhanced [3H]NA release in the rat prostate to an extent similar to that produced by Ang II or Ang I, suggesting that bradykinin enhances sympathetic transmission in the prostate. Moreover, the effect of bradykinin on [3H]NA release was further enhanced after ACE inhibition with captopril, suggesting that exogenous bradykinin is subject to degradation by prostatic ACE. This is consistent with the findings of other studies in which the facilitatory effect of bradykinin on NA release was revealed after ACE inhibition with captopril in the human kidney and atria (Rump et al. 1995, 1997). These effects of bradykinin on [3H]NA release in the rat prostate were significantly inhibited by Hoe140, suggesting that facilitation of transmitter NA release by bradykinin in the rat prostate is mediated by B2 receptors. This is also in accordance with other studies in which bradykinin was found to increase NA release in the rat heart (Kurz et al. 1997) and mouse heart (Chulak et al. 1998), pithed rat and PC12 cells (Dendorfer & Dominiak 1995, Dendorfer et al. 1996) via B2 receptors – an effect that was markedly antagonised by Hoe140.

These findings suggest that ACE may have an important neuromodulatory role on sympathetic transmission in the prostate by regulating the synthesis of Ang II and the metabolism of bradykinin, both of which have demonstrable effects on transmitter NA release in the rat prostate. Therefore, although ACE inhibition may suppress local Ang II–mediated effects on sympathetic transmission in the prostate, this may potentially be compensated by the effects mediated by bradykinin accumulation. Further to this, bradykinin has been shown to induce contraction of the canine prostate, which can be potentiated further by ACE inhibition (Steidle et al. 1990). Thus, if the RAS has an obligatory role in the pathophysiology of BPH, then suppression of the RAS with AT1 receptor blockers, rather than ACE inhibition, may offer potential benefits by virtue of the fact that the former do not interfere with bradykinin metabolism.

The findings of the present study provide direct evidence that Ang II enhances NA release from sympathetic nerves of the rat prostate. These data establish a novel functional role for the RAS in the modulation of sympathetic transmission in the prostate, which may have important implications for the understanding of the pathophysiology of BPH. Increased local sympathetic activity is a characteristic feature of BPH and represents a target for drug treatment with α1-adrenoceptor blockers. Recent findings from our laboratory suggest that the local RAS is activated in BPH. Specifically, we have shown that the expression of ACE mRNA and protein (Nassis et al. 2000, 2001) and Ang II peptide (Dinh et al. 2001a) is significantly increased in BPH compared with the normal prostate. Furthermore, we have demonstrated that AT1 receptors predominate in the human prostate and are down-regulated in BPH, which may be due to receptor internalisation as a result of receptor hyper-stimulation by increased tissue concentrations of Ang II (Dinh et al. 2001a,b). It is possible, therefore, that hyperactivity of the local RAS resulting in increased tissue concentrations of Ang II may represent an important factor in the pathophysiology of BPH by enhancing local sympathetic activity in the prostate. Facilitation of NA release from sympathetic nerves by Ang II would consequently result in hyper-stimulation of α1-adrenoceptors, causing contraction of prostatic smooth muscle and urethral compression, with subsequent resistance to urinary outflow.

In conclusion, exogenous and locally generated Ang II facilitates the release of NA from sympathetic nerves of the rat prostate by a prejunctional mechanism. The receptor subtype mediating the effects of Ang II on sympathetic transmission in the rat prostate is unclear, but may involve...
a novel functional Ang II receptor distinct from the cloned AT$_1$-b, AT$_1$h or AT$_2$. These novel data provide direct evidence in support of a functional role for the local RAS in modulating sympathetic activity in the prostate, which may have important implications for the pathophysiology of BPH.

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