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Receptors for Atrial Natriuretic Peptide Are Decreased in the Kidney of Rats with Streptozotocin-induced Diabetes Mellitus

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Abstract

To determine whether decreased renal responsiveness to atrial natriuretic peptide (ANP) in diabetes is mediated by alterations in the renal ANP receptor, ANP receptor density and affinity were measured 17–20 d after streptozotocin injection and compared with values in vehicle-treated controls and streptozotocin-treated rats made euglycemic with insulin. Plasma ANP concentration was significantly greater in hyperglycemic diabetic rats than in control or euglycemic diabetic rats. Both in glomeruli and inner medulla, ANP receptor dissociation constant did not differ among the three study groups, whereas the maximum binding capacity was decreased significantly in hyperglycemic diabetics in comparison with controls and euglycemic diabetics. Glomerular clearance receptors were also decreased significantly in hyperglycemic diabetic rats in comparison with control and euglycemic diabetic rats. To determine whether the decreased number of renal ANP receptors in diabetic rats was associated with a decreased biological response, we measured ANP-dependent cyclic GMP (cGMP) accumulation by isolated glomeruli and inner medullary collecting duct cells in vitro. cGMP accumulation was significantly less in hyperglycemic diabetic rats than in controls or euglycemic diabetic rats both in the presence or absence of the phosphodiesterase inhibitor zaprinast. cGMP phosphodiesterase activity in inner medullary collecting duct cells obtained from control and hyperglycemic diabetic rats did not differ. Thus, the decreased number of biologically active ANP receptors in the kidneys of diabetic rats is accompanied by decreased biological responsiveness in vitro and provides a potential explanation for the reduction in renal sensitivity to ANP in this condition. (J. Clin. Invest. 1995. 95:2451–2457.) Key words: atrial natriuretic peptide • cyclic GMP • diabetes • kidney • receptors

Introduction

Increased exchangeable sodium (1, 2) and extracellular fluid volume (3) occur in diabetic humans and experimental animals, respectively. Whereas the kidney is the principal organ responsible for the chronic regulation of body sodium and fluid composition (4), renal disease per se is an unlikely explanation for these abnormalities since they can occur quite early in the disease, even when glomerular filtration rate is normal and proteinuria is minimal (1, 2). Enhanced renal tubular reabsorption and/or impaired responses to natriuretic influences are more likely explanations.

In both rats (5–7) and humans (8) with insulinopenic diabetes, the glomerular and tubular response of the kidney to atrial natriuretic peptide (ANP) is decreased. The basis for this decreased responsiveness has not been established, but may involve an abnormality of renal ANP receptors. Three distinct natriuretic peptide receptors have been identified (9): ANPR-A and ANPR-B, also called GC-A and GC-B, are both membrane guanyl cyclases that synthesize cyclic GMP (cGMP) as an intracellular second messenger (9, 10). ANPR-C, also known as the ANP clearance receptor, does not signal through activation of guanyl cyclase, but may serve as a clearance receptor for natriuretic peptides; it may also function through intermediate guanine nucleotide regulatory proteins to inhibit adenylyl cyclase (11). While ANPR-A and ANPR-B have rather strict structural requirements for ligand binding (12–15), ANPR-C has less rigorous structural requirements and binds the internally deleted analogues of ANP (c-ANP) (12, 16–18), compounds which do not activate the membrane guanylate cyclase.

ANPR-A and ANPR-C receptors and their mRNAs have been demonstrated in the kidney using in situ autoradiographic techniques (19, 20) and in situ hybridization (21), respectively. This study was designed to investigate the type, number, and affinity of ANP receptors in the kidney of rats with streptozotocin (STZ)-induced diabetes and to assess the biological activity of these receptors by measuring ANP-dependent cGMP accumulation by glomeruli and inner medullary collecting duct cells (IMCD) isolated from these rats.

Methods

Preparation of diabetic rats. Male Sprague-Dawley rats (Bantin and Kingman, Inc., Fremont, CA), ranging in weight from 200 to 250 g, were maintained on standard rat chow and tap water ad libitum with a 12-h light/dark cycle in a quiet environment. In 19 rats, diabetes mellitus was induced by intravenous administration of STZ (Sigma, Chemical Co., St. Louis, MO), 60 mg/kg body wt, dissolved in sodium citrate buffer (0.1 mol/liter, pH 4.5) at a concentration of 20 mg/ml immediately before use. Control rats (n = 11), which were matched for age and weight at the time of STZ administration, received an equal volume of the vehicle. Animals were considered to be diabetic if blood glucose

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1. Abbreviations used in this paper: ANP, atrial natriuretic peptide; Bmax, maximum binding capacity; IMCD, inner medullary collecting duct; PDE, phosphodiesterase; STZ, streptozotocin.
levels were 350 mg/dl or greater 72 h after the injection. Diabetic rats were further divided into two subgroups: rats to which human recombinant insulin (Humulin N; Eli Lilly & Co., Indianapolis, IN) was administered once daily in a fixed dose (2 U/d, n = 11) to prevent ketosis, and rats to which insulin was administered in doses that were adjusted to achieve euglycemia (6–10 U/d, n = 8). Blood glucose levels were measured on a drop of blood obtained by tail vein nicking (Accu-check Bg; Bio-Dynamics, Boehringer-Mannheim, Indianapolis, IN). To determine the 24-h urine volume and the rate of excretion of sodium and cGMP, rats were placed in metabolic cages on the day before killing. Rats were killed by decapitation 17–20 d after the administration of STZ, and trunk blood was collected for determination of plasma glucose and ANP levels. The kidneys were quickly removed, decapsulated, snap-frozen in liquid nitrogen, and stored at –80°C. Plasma ANP was measured by radioimmunoassay (Peninsula Laboratories, Inc., Belmont, CA), after extraction with 0.1% trifluoroacetic acid and 75% methanol.

Receptor binding studies. The density of ANP receptors was assessed by a modification of an in situ receptor binding assay (19) as reported previously (20). Briefly, tissue sections (10 μm) were cut on a cryostat at –15°C, thaw-mounted on poly-l-lysine–coated slides, dried in vacuo for 18 h at –4 to –4°C over silica gel, and stored in sealed Bakelite boxes at –80°C. Immediately before assay, the slides were brought to room temperature. Sections were preincubated for 10 min at room temperature in 100 μl of a buffer (A) containing 30 mM sodium phosphate (pH 7.2), 120 mM sodium chloride, 0.3% bacitracin, and 0.5% receptor grade bovine serum albumin. After preincubation, the buffer was replaced with 100 μl of fresh buffer A containing increasing concentrations (from 25 to 1,500 PM) of [125I]-rat-ANP (1-28) (2,200 μCi/mm) (DuPont/New England Nuclear, Boston, MA), and the sections were placed in a humidified chamber and incubated at room temperature for 15 min. After incubation, the slides were rinsed in an ice-cold buffer containing 30 mM sodium phosphate and 120 mM sodium chloride for 10 s, washed in the same buffer for 5 min, rinsed in deionized water for 10 s, and dried for 2 h in a stream of cool air. For autoradiography, the sections were exposed to LKB-Ulrofilm (Leica Inc., Deerfield, IL) for 3–5 d at room temperature, and films were processed with Kodak D-19 developer and rapid fixer (Eastman Kodak Co., Rochester, NY). Thereafter, the amount of radioligand bound to the tissue sections was determined by placing the slides in a gamma counter. Regional analysis of ANP binding to glomeruli and inner medulla was performed using film autoradiography. Optical density in the different regions was measured by computerized microdensitometry (Image-Analysis-OS2; Imaging Research Inc., St. Catherine, Ontario, Canada).

Non-specific binding was determined on adjacent sections under identical incubation conditions except for the addition of 1 μM unlabeled rat ANP (1-28). (Peninsula Laboratories, Inc.). ANPR-C receptors were identified by addition of 10 μM c-ANP (4-23) (Peninsula Laboratories, Inc.). ANPR-A receptors were calculated by subtraction of ANPR-C receptors from total ANP receptors. The apparent dissociation constant (Kd) and the maximal binding capacity (Bmax) were calculated by Scatchard analysis using the LIGAND program of Munson and Rodbard (22). Because 10 μM c-ANP (4-23) could theoretically occupy some of the ANPR-A receptors, we performed experiments in which IMCD cells obtained from kidneys of normal rats were incubated with 10 μM c-ANP (4-23) and cGMP production was measured (see below). Accumulation of cGMP did not differ significantly in cells incubated with or without cGMP (126±15 vs 119±21 fmol/mg protein per 10 min, respectively), ruling out the possibility of significant c-ANP binding to ANPR-A receptors at this concentration.

Preparation of isolated glomeruli and IMCD cells. Under Inactin® (B&K, Guelen, Germany) anesthesia (100 mg/kg body wt, intraperitoneally), the suprarenal aorta was cross-clamped via the distal aorta, and the kidneys were flushed with cold isotonic saline (5–10 ml). Glomeruli were isolated as described previously (23). In brief, for each preparation the renal cortices were dissected and contained by a pastelike consistency. The homogenate was passed successively through a 106-μm sieve that excluded the tubules and blood vessels and a 75-μm sieve that retained the glomeruli and allowed cells and small debris to pass through. Glomeruli were suspended in ice-cold 20 mM Tris-HCl buffer, pH 7.4, containing 135 mM sodium chloride, 10 mM potassium chloride, 10 mM sodium acetate, and 5 mM glucose (buffer B) and centrifuged at 120 g for 2 min. The supernatant was discarded, and the pellet was resuspended in the same buffer and recentrifuged. By light microscopic examination, the final pellet consisted of nearly pure isolated decapsulated glomeruli with <5% contamination with tubules. No afferent or efferent arterioles were observed.

IMCD cells were isolated as follows (24): the kidneys were perfused with 50–60 ml of saline followed by 5 ml of saline containing 0.2% of collagenase (type II; Sigma Chemical Co.). The inner medullae were excised, finely minced, and incubated in the same solution for 90 min at 37°C. The resulting suspension of inner medullary cells was layered on a 16% Ficoll (Sigma Chemical Co.) solution in noncarbonate Ring- er’s buffer (buffer C) and centrifuged at 2,300 g for 40 min. Cells were subsequently washed through buffer C and then a solution of buffer C containing 7.5% albumin to remove traces of contaminating collagenase.

Determination of cGMP accumulation by glomeruli and IMCD cells. Freshly isolated glomeruli were resuspended in buffer B with 1 mM calcium chloride and IMCD cells in buffer C containing 7.5 mM glucose. 350-μl aliquots containing 0.2–0.5 mg protein/tube were preincubated for 10 min at 37°C in a shaking water bath. Incubations were then carried out for 10 min at 37°C with varying concentrations of theophylline (0 to 10 mM), in the presence or absence of a phosphodiesterase (PDE) inhibitor, zaprinast (1 mM, M&B 22,948; Rhone-Poulenc, Rorer, Dan- genham, United Kingdom). Incubations were terminated by adding 750 μl of ice-cold trichloroacetic acid (TCA; final concentration, 6.6%) and cooling to 4°C. The precipitated protein was sedimented by centrifugation at 4,500 rpm for 15 min at 4°C, and the pellets were dissolved in 0.5 N sodium hydroxide and assayed for protein content by the method of Lowry (25) using bovine serum albumin as a standard. The supernatant fluid was extracted five times with four volumes of water-saturated ethyl ether to remove the TCA before being evaporated to dryness under a stream of air and stored at –70°C until assayed for cGMP content. For the cGMP assay, samples were dissolved in 50 mM sodium acetate buffer, pH 6.2, mixed thoroughly, and 100 μl aliquots acetylated and assayed in duplicate with a commercially available kit according to the manufacturer’s instructions (New England Nuclear). Results are expressed as femtomoles of cGMP generated per milligram of protein per 10 min of incubation.

Preparation of IMCD cell membranes and cytosol. IMCD cell membranes were prepared by homogenizing freshly dispersed cells in 10 mM Tris-HCl (pH 7.4), 10 mM Mes, 1 mM EDTA, 1 mM EGTA, 0.1 μM leupeptin, 0.1 μM pepstatin, 0.1 μM PMSF, 0.25 M sucrose, 5 mM 2-mercaptoethanol, and 0.1% Triton X-100. Cytosol was obtained by homogenizing cells in a buffer containing 5 mM Tris-HCl (pH 7.4), 3 mM MgCl2, and 1 mM EDTA, in an amount equal to 1:3 wt/wt volume. In each case, cells were homogenized in a Potter-type Teflon-glass homogenizer for 10–12 strokes at 4°C; the homogenates were initially centrifuged at 200 g for 5 min and the resultant supernatants were incubated with agitation at 4°C for 60 min followed by centrifugation at 100,000 g for 60 min. The resulting pellets containing membranes were resuspended in 1:10 vol/vol homogenizing buffer, rehomogenized, and again ultracentrifuged, while the supernatants representing the cytosolic fraction were stored on ice. The supernatant from the second high speed centrifugation was added to that previously collected and immediately frozen at –70°C for subsequent determination of PDE activity, and the pellet resuspended in 100 mM Hepes (pH 7.4) and 50 mM NaCl for assay of guanylate cyclase activity.

Determination of guanylate cyclase activity in IMCD cells membranes. Guanylate cyclase activity was assayed by a modification of the method of Cunning (26). Briefly, enzyme activity was determined by incubation of 40–100 μg of membrane protein with 4–5 mM GTP substrate in the presence of 2 mM manganese, 1.2 mM ATP, 1 mM xanthine, 5 mM theophylline, creatine phosphokinase (3 U), and 7.5 mM phosphocreatine. All incubations were performed for 5 min at 37°C and terminated by 12% TCA precipitation. After centrifugation, the supernatants were either extracted, air-dried, and reconstituted in 50 mM sodium acetate (pH 6.2) followed by cGMP measurements in duplicate
by radioimmunoassay. cGMP production was linear as a function of incubation time (1–10 min) and protein concentration (10–100 μg). Protein concentration was determined by the Coomassie protein assay.

Determination of PDE activity in IMCD cells. cGMP PDE activity was assayed according to the method of Torres et al. (27). Cytosolic fractions of IMCD cell homogenates, collected as described (10–15 μg protein/tube), were incubated with 1 μM [3H]cGMP (New England Nuclear) in 200 μl of a buffer containing 10 μM magnesium sulphate, 2 mM EGTA, 0.1% bovine serum albumin, and 50 mM Tris-HCl (pH 7.5). After a 20-min incubation period at 37°C, the reaction was stopped by heating at 95°C for 3 min, and the [3H]-5′-nucleotide products converted to 3H-nucleosides by incubation with an excess of 5′-nucleotidase (Crotalus atrox snake venom; Sigma Chemical Co.). Nucleosides were separated from nucleotides on a QAE-Sephadex column according to the procedure of Wells et al. (28) and column eluates counted for 3H activity in a liquid scintillation counter. Enzyme activity is expressed in picomoles of cGMP hydrolyzed per minute per milligram of protein. With this assay, activity was linear over time and with protein concentration.

Statistical analysis. Data are presented as mean±standard error.

Table I. Characteristics of Control, Hyperglycemic Diabetic, and Euglycemic Diabetic Rats

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>Hyperglycemic</th>
<th>Euglycemic</th>
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<tbody>
<tr>
<td>n</td>
<td>11</td>
<td>7</td>
<td>8</td>
</tr>
<tr>
<td>Body wt (g)</td>
<td>356±12</td>
<td>285±9*</td>
<td>342±8</td>
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<tr>
<td>Kidney wt (g)</td>
<td>2.37±0.07</td>
<td>2.46±0.07</td>
<td>2.40±0.05</td>
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<td>Kidney/body wt ratio (×100)</td>
<td>0.67±0.01</td>
<td>0.86±0.02*</td>
<td>0.70±0.02</td>
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<tr>
<td>Blood glucose (mg/dl)</td>
<td>128±5</td>
<td>486±41*</td>
<td>105±30</td>
</tr>
<tr>
<td>Plasma ANP (pg/ml)</td>
<td>229±20</td>
<td>312±33*</td>
<td>178±22</td>
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<tr>
<td>Urinary vol (ml/24 h)</td>
<td>19.7±1.6</td>
<td>117.0±7.0*</td>
<td>30.8±4.8</td>
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<tr>
<td>Sodium excretion (mM/24 h)</td>
<td>1.96±0.23</td>
<td>2.50±0.29</td>
<td>2.14±0.03</td>
</tr>
<tr>
<td>cGMP excretion (nM/24 h)</td>
<td>41.6±5.2</td>
<td>37.8±6.4</td>
<td>32.5±3.7</td>
</tr>
</tbody>
</table>

Values are means±SEM. Significant difference versus control rats is indicated by *P < 0.01, and †P < 0.05.

Comparisons were done by ANOVA (Statview; BrainPower Inc., California, CA). A P value < 5% was considered to be statistically significant.

Results

Blood glucose, body weight, and plasma ANP. Hyperglycemia occurred within 3 d of administration of STZ and persisted thereafter in the rats that received low-dose insulin therapy (Fig. 1, Table I). Hyperglycemia was completely ameliorated in the rats that received the higher doses of insulin (Fig. 1, Table I). Rats treated with the higher doses of insulin gained weight rapidly, reaching values not significantly different from control at the end of the experimental period, whereas the rate of growth was significantly slower in rats treated with low-dose insulin. Data are mean±SEM of 11 control, 11 low-dose, and 8 high-dose insulin–treated animals.

Figure 1. Blood glucose concentration (top) and body weight (bottom) in control, low-dose, and high-dose insulin–treated diabetic rats. Hyperglycemia occurred within 3 d of administration of STZ and persisted thereafter in the rats that received low-dose insulin therapy. Hyperglycemia was completely ameliorated in the rats that received the higher doses of insulin. Rats treated with the higher doses of insulin gained weight rapidly, reaching values not significantly different from control at the end of the experimental period, whereas the rate of growth was significantly slower in rats treated with low-dose insulin. Data are mean±SEM of 11 control, 11 low-dose, and 8 high-dose insulin–treated animals.
Increasing concentrations of radioligand resulted in an increase in both total and nonspecific binding. Receptor saturation was approached at 1.2 nmol/liter in both control and hyperglycemic diabetic rats (Fig. 3). Scatchard analysis of these data suggested the presence of a single receptor site in each of the three experimental groups (Fig. 4). ANP receptor affinity, expressed as the apparent $K_d$, did not differ among the three study groups, whereas the $B_{max}$ for ANP was significantly decreased in hyperglycemic diabetic rats as compared with controls and normoglycemic diabetic rats. Regional analysis of ANP binding was performed by computerized microdensitometry and showed that ANP receptors in glomeruli and inner medulla in the hyperglycemic diabetic rats. Equilibrium binding data obtained in glomeruli in the presence of c-ANP showed that the number of ANP receptors was significantly lower in the kidney of hyperglycemic diabetic rats in comparison with controls and euglycemic diabetics (Table II).

$\text{cGMP accumulation by glomeruli and IMCD cells and guanylate cyclase activity.}$ ANP induced a dose-dependent increase in glomerular cGMP generation in the three groups (Fig. 5). Both in glomeruli and IMCD cells, cGMP accumulation was lower in hyperglycemic diabetic rats than in controls at the higher concentrations used. Similar differences between hyperglycemic diabetic and control rats were found when glomeruli were incubated in the presence of zaprinast (Fig. 6), indicating that these differences could not be ascribed to an alteration in the activity of cGMP PDE. Guanylate cyclase activity was comparable in membranes obtained from IMCD cells in control rats, with a value of 5.983 ± 5.48 fmol GMP·mg protein$^{-1}$·min$^{-1}$ and hyperglycemic diabetic rats (5.835 ± 8.14 fmol GMP·mg protein$^{-1}$·min$^{-1}$) rats, indicating no effect of diabetes on basal activity of this enzyme.

$\text{PDE enzymatic activity in IMCD cells.}$ Measurement of PDE enzymatic activity in IMCD cytosol showed comparable values in control rats (180 ± 28 fmol·mg protein$^{-1}$·min$^{-1}$) and hyperglycemic diabetic rats (156 ± 23 fmol·mg protein$^{-1}$·min$^{-1}$; NS).

**Discussion**

This study demonstrates that the number of ANP receptors is decreased in the kidney of rats with STZ-induced diabetes mellit-
This abnormality is associated with a decreased production of cGMP by glomeruli and IMCD cells which is not corrected by inhibition of PDE activity. When studied 2 wk after administration of STZ, we found that plasma ANP levels in hyperglycemic diabetic rats were significantly increased in comparison with controls and that insulin therapy sufficient to achieve strict glycemic control normalized plasma ANP levels. In previous studies, plasma ANP levels in diabetic rats were increased as early as 1 wk after STZ injection (30) and persisted at elevated levels for at least 6 wk (7, 31). Plasma ANP levels are also increased in patients with insulin-dependent diabetes mellitus and diabetic nephropathy.

### Table II. Binding Parameters of Biological (ANPR-A) and Clearance (ANPR-C) Receptors for ANP in the Kidney of Control, Hyperglycemic Diabetic, and Euglycemic Diabetic Rats

<table>
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<th>Diabetic</th>
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<tr>
<td></td>
<td>Control</td>
<td>Hyperglycemic</td>
<td>Euglycemic</td>
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<td>3</td>
<td>3</td>
<td>3</td>
</tr>
<tr>
<td>Glomeruli</td>
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<tr>
<td>ANPR-A $K_d$ (nM)</td>
<td>1.2±0.4</td>
<td>1.1±0.2</td>
<td>1.4±0.5</td>
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<tr>
<td>ANPR-A $B_{max}$ (pM/mm$^3$)</td>
<td>23.8±1.9</td>
<td>14.1±3.1$^*$</td>
<td>20.9±3.0</td>
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<tr>
<td>ANPR-C $K_d$ (nM)</td>
<td>0.5±0.3</td>
<td>0.7±0.4</td>
<td>0.6±0.2</td>
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<tr>
<td>ANPR-C $B_{max}$ (pM/mm$^3$)</td>
<td>8.4±1.3</td>
<td>4.7±1.2$^*$</td>
<td>7.2±1.9</td>
</tr>
<tr>
<td>Inner medulla</td>
<td></td>
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<tr>
<td>ANPR-A $K_d$ (nM)</td>
<td>1.3±0.3</td>
<td>1.2±0.4</td>
<td>1.8±0.7</td>
</tr>
<tr>
<td>ANPR-A $B_{max}$ (pM/mm$^3$)</td>
<td>25.3±2.8</td>
<td>13.7±2.7$^*$</td>
<td>19.9±3.5</td>
</tr>
</tbody>
</table>

Values are means±SEM. Significant differences versus control rats are indicated by $^*$ $P < 0.05$, and $^* P < 0.01$. This increase in ANP levels could be a consequence of the plasma volume expansion present in both STZ-diabetic rats (7, 31, 36, 37) and patients with diabetes mellitus (2, 38–41). Despite increased plasma levels of ANP, the response to maneuvers known to stimulate the release of the peptide, such as saline infusion, is impaired (5, 31). This has been interpreted as a consequence of partial exhaustion of the storage pool of the hormone as also demonstrated by morphologic studies in which the abundance of secretory granules is decreased in atrial cardiomyocytes obtained from diabetic rats (31, 42).

Blunted glomerular and tubular responses to volume expansion (5, 6) and exogenous ANP infusion (7) have been reported in rats with STZ-induced diabetes. Similarly, a decreased natriuretic response to endogenously secreted ANP has been shown in type 1 diabetic subjects after head-out water immersion (8). In the present study, we found that the number of both biological and clearance ANP receptors is significantly decreased in the kidneys of STZ-diabetic rats, providing an explanation for the decreased renal response to the hormone. Similarly, Benigni et
Circulating levels of peptide hormones may directly regulate target tissue receptor number, and plasma concentrations may negatively correlate with the density of specific receptors (46). This also applies to circulating ANP, the levels of which determine receptor density in the kidney (47–49). Therefore, homologous down-regulation caused by increased plasma ANP levels is the most obvious explanation of the decreased ANP receptor number in the kidney of diabetic rats. Alternatively, a decreased number of receptors could reflect a primary phenomenon independent of plasma ANP levels. It has been shown that the number of glomerular receptors for other circulating peptides such as angiotensin II (50, 51) and thromboxane A2 (52) are decreased in diabetes mellitus. It is also known that glucose levels regulate the activity of protein kinase C (53, 54) which in turn regulates the number of receptors present on the cell membrane (55, 56). Therefore, it is possible that receptors for different peptides are regulated by plasma glucose levels through changes in the activity of protein kinase C. Further studies will be required to distinguish among these various possibilities.

In summary, our results demonstrate a reduced binding of ANP in both glomeruli and IMCD cells of diabetic rat kidneys. The diminished binding is associated with diminished biological activity in vitro in these target sites of renal ANP action. This abnormality may consequently be the mechanism underlying renal resistance to ANP in diabetic animals and patients.

Acknowledgments

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