In vivo bradykinin B2 receptor activation reduces renal fibrosis

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Angiotensin-converting enzyme (ACE) inhibitors reduce the progression of various fibrotic renal diseases both in humans and in animal models. Unilateral ureteral obstruction (UUO) is an animal model of accelerated renal tubulointerstitial fibrosis that is attenuated by ACE inhibition. Although ACE inhibitors increase bradykinin concentrations in addition to their effect on angiotensin II formation, the role of bradykinin in renal fibrosis has not been studied. We show here that genetic ablation (B2−/− mice) or pharmacological blockade of the bradykinin B2 receptor increases UUO-induced interstitial fibrosis in mice, whereas transgenic rats expressing increased endogenous bradykinin show reduced UUO-induced interstitial fibrosis. The increased interstitial fibrosis in B2−/− mice was accompanied by a decreased activity of plasminogen activators (PAs) and metalloproteinase-2 (MMP-2), enzymes involved in ECM degradation, suggesting that the protective effects of bradykinin involve activation of a B2 receptor/PA/MMP-2 cascade. This ability of bradykinin to increase PA activity was confirmed in primary culture proximal tubular cells. Thus, in both mice and rats, bradykinin B2 receptor activation reduces renal tubulointerstitial fibrosis in vivo, most likely by increasing ECM degradation.


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Nonstandard abbreviations used: unilateral ureteral obstruction (UUO); angiotensin-converting enzyme (ACE); angiotensin type I (AT1); plasminogen activator (PA); urokinase-type PA (tPA); tissue-type PA (uPA); periodic acid-Schiff (PAS); proliferating cell nuclear antigen (PCNA); α-smooth muscle actin (α-SMC); Krebs Henseleit buffer (KHB); monocyte chemoattractant peptide-1 (MCP-1); PA inhibitor type 1 (PAI-1); N-methyl-D-aspartate (NMDA).
protein–coupled B2 receptor in renal tubulointerstitial fibrosis have been reported. That is the reason why we studied the functional role of the bradykinin B2 receptor in the development of tubulointerstitial fibrosis caused by UUO.

We report here that renal tubulointerstitial fibrosis induced by UUO is significantly higher in B2 receptor knockout (B2−/−) mice than in wild-type (B2+/+) mice. Confirming this observation, UUO-induced tubulointerstitial fibrosis was reduced in transgenic rats overexpressing the tissue kallikrein gene (expressing increased endogenous bradykinin). It was found that B2−/− mice have a lower plasminogen activator (PA) activity associated with a decrease in metalloproteinase (MMP-2) activity, which can explain the increased UUO-induced tubulointerstitial fibrosis in B2−/− mice. The ability of bradykinin to increase PA activity was confirmed in primary culture renal proximal tubular cells. The present study demonstrates, we believe for the first time in vivo, that bradykinin B2 receptor activation plays a protective role in renal tubulointerstitial fibrosis.

**Methods**

**Animals.** B2−/− mice were generously provided by F. Hess and T. MacNeil (Merck & Co. Inc., Rahway, New Jersey, USA) (14). B2−/− mice were originally on a mixed genetic background (J129sv × C57Bl/6j). We have backcrossed (ten times) the B2−/− mice to C57Bl/6j and have therefore used C57Bl/6j as control mice. Microsatellite analyses confirmed the C57Bl/6j genetic background of the mice by backcrossing generated B2−/− mice (Nucleis, Angers, France). Furthermore, the mice are housed in a pathogen-free environment. Transgenic rats overexpressing the human tissue kallikrein gene, TGR(hKLK1), and their transgene negative littermates were generated as described previously (15).

**Experimental protocols.** Male mice or rats at the age of 8 weeks were used in these experiments, ten mice or five rats in each group. The unilateral ureteral ligation was performed as follows: under oxygen-isoflurane anesthesia, through a longitudinal, left abdominal incision, the ureter was exposed and ligated with a 6/0 nylon thread at the uretero-pelvic junction. In sham operations, the ureter was exposed but not ligated and repositioned. Mice and rats were maintained on a standard mouse or rat chow and tap water. All treatments were initiated 1 day before obstruction and continued throughout the time of obstruction (3 hours to 14 days). B2 receptor antagonist HOE-140 (500 µg/kg; HOECHST) was injected daily by one subcutaneous injection. At the end of the different protocols, mice or rats were sacrificed, and the kidneys were removed and sectioned at 3 µm and subjected to the following analyses. Once deparaffinized and rehydrated, the kidney sections were stained with Sirius red and periodic acid-Schiff (PAS) to evaluate interstitial collagen deposition and tubular atrophy, respectively. Immunohistochemical studies for the detection of fibronectin, collagen type I, III, IV, macrophages, the proliferating cell nuclear antigen (PCNA), and α-smooth muscle actin (α-SMC) for the detection of myofibroblasts were performed as follows. Deparaffinized sections were incubated with 3% hydrogen peroxide to block the endogenous peroxidase activity. The sections were then incubated for 1 hour at room temperature in a humidified atmosphere with the following primary Ab's: goat polyclonal anti-human fibronectin (SC-6953; Santa Cruz Biotechnology Inc., Santa Cruz, California, USA), rabbit polyclonal anti-rat collagen type I (AB755; Chemicon International, Temecula, California, USA), rabbit anti-human collagen III and IV (T59105R and T59106R, respectively; Interchim, Montluçon, France), the rat mAb to mouse F4/80 (RM2900; Caltag Laboratories Inc., Burlingame, California, USA) for macrophage detection, and the monoclonal mouse anti-human α-SMC (DAKO EPOS method, U7033; DAKO S.A., Trappes, France). For fibronectin and collagen I, the sections were sequentially processed with the goat (fibronectin) or rabbit (collagen I) ABC Staining System (Santa Cruz Biotechnology Inc.), which uses 3,3′-diaminobenzidine as chromogen. For the visualization of collagen III and IV and F4/80, the DAKO Envision System was used (DAKO S.A.). Cell proliferation was assessed by detection of PCNA using the Zymed PCNA staining kit (Zymed Laboratories Inc., San Francisco, California, USA). Sections were counterstained with hematoxylin. For all samples, negative controls for the immunohistochemical procedures included substitution of the primary Ab with nonimmune sera. The presence of myofibroblasts was analyzed on PCNA adjacent sections to be able to compare both localizations. Glomeruli and vasculature were excluded from the evaluation in the present study.

**Histomorphometric analysis.** Analyses were performed by an operator unaware of the origin of each kidney section. Under a light microscope (Nikon Eclipse 600) at ×200 magnification, 20 nonoverlapping fields (to obtain around 90% of the kidney section) per kidney section were captured with a numeric camera (Cool SNAP, RS Photometrics, Tucson, Arizona, USA) connected to the microscope. Quantitative analysis of the pictures was performed by using the Adobe Photoshop 5.5 software, which allows counting of the pixels stained specifically (red for Sirius red and brown for the immunohistochemical studies). The results are given in percentage of specific colored pixels per total number of pixels of the 20 fields, representing thus the surface occupied by the analyzed marker. For PCNA and α-SMC detection, marked cells were counted.

**Immunohistochemistry.** Kidneys were fixed in Carnoy’s solution, dehydrated, embedded in paraffin, and then sectioned at 3 µm and subjected to the following analyses. Once deparaffinized and rehydrated, the kidney sections were stained with Sirius red and periodic acid-Schiff (PAS) to evaluate interstitial collagen deposition and tubular atrophy, respectively. Immunohistochemical studies for the detection of fibronectin, collagen type I, III, IV, macrophages, the proliferating cell nuclear antigen (PCNA), and α-smooth muscle actin (α-SMC) for the detection of myofibroblasts were performed as follows. Deparaffinized sections were incubated with 3% hydrogen peroxide to block the endogenous peroxidase activity. The sections were then incubated for 1 hour at room temperature in a humidified atmosphere with the following primary Ab’s: goat polyclonal anti-human fibronectin (SC-6953; Santa Cruz Biotechnology Inc., Santa Cruz, California, USA), rabbit polyclonal anti-rat collagen type I (AB755; Chemicon International, Temecula, California, USA), rabbit anti-human collagen III and IV (T59105R and T59106R, respectively; Interchim, Montluçon, France), the rat mAb to mouse F4/80 (RM2900; Caltag Laboratories Inc., Burlingame, California, USA) for macrophage detection, and the monoclonal mouse anti-human α-SMC (DAKO EPOS method, U7033; DAKO S.A., Trappes, France). For fibronectin and collagen I, the sections were sequentially processed with the goat (fibronectin) or rabbit (collagen I) ABC Staining System (Santa Cruz Biotechnology Inc.), which uses 3,3′-diaminobenzidine as chromogen. For the visualization of collagen III and IV and F4/80, the DAKO Envision System was used (DAKO S.A.). Cell proliferation was assessed by detection of PCNA using the Zymed PCNA staining kit (Zymed Laboratories Inc., San Francisco, California, USA). Sections were counterstained with hematoxylin. For all samples, negative controls for the immunohistochemical procedures included substitution of the primary Ab with nonimmune sera. The presence of myofibroblasts was analyzed on PCNA adjacent sections to be able to compare both localizations. Glomeruli and vasculature were excluded from the evaluation in the present study.

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RNA preparation and Northern blot analysis. Total RNA from snap-frozen kidney sections was extracted using the RNaseasy Midi kit (QIAGEN Inc., Valencia, California, USA). Glyoxal and dimethyl sulfoxide denatured RNA (20 µg) was separated on a phosphate-buffered 1% agarose gel and transferred under vacuum to a blotting device on a positively charged polyamide membrane (Schleicher & Schuell Inc., Keene, New Hampshire, USA). The cDNA probes (between 200 and 350 bp) were obtained by RT-PCR, verified by sequencing, and radiolabeled with [α-32P]dATP using the Strip-EZ DNA Kit (Ambion Inc., Austin, Texas, USA). Hybridization was performed in Ultrahyb buffer (Ambion Inc.). Densitometric analysis was performed as described (16).

Primary culture of mouse renal proximal tubular cells. Mouse proximal tubular cells were obtained by an extension of a method described previously (17). Briefly, after removal of the kidneys, the kidney cortex was minced in HBSS at 4°C and incubated two times for 20 minutes at 37°C with 0.2 mg/ml collagenase solution supplemented by 1 mg/ml BSA. After centrifugation (5 minutes at 135 g), the pellet was washed two times with HBSS and two times with HBSS containing 1% BSA. To obtain homogenous populations of nephron segments, the mixture of tubules was resuspended in 42% Percoll made isotonic with concentrated (10%) Krebs Henseleit buffer (KHB), pH 7.3 — 1.18 M NaCl, 47 mM KCl, 100 mM HEPES, 200 mM cyclic acid, 1.26 mM MgSO4, 11.4 mM KH2PO4, and 50 mM glucose — and centrifuged (34,900 g for 30 minutes at 4°C). The F4 layer, containing the proximal tubules, was suspended in culture medium (DMEM/HAM F12) supplemented with 5% FCS, 10 µg/ml insulin, 5 mM transferrin, 10 ng/ml EGF, 0.1 µM dexamethasone, and 5 µg/ml triiodothyronin, and plated at a concentration of 1.6 mg of protein per 100 mm² of a Petri dish coated with collagen. The proximal tubular origin and epithelial cell type was confirmed by positive alkaline phosphatase activity and cytokeratin staining (data not shown). Western blot analysis and calcium mobilization experiments on proximal tubular cells were performed as described previously (18). The B2 receptor Ab (mouse mAb B40820) used in the Western blot experiment (dilution 1/500) was from Transduction Laboratories (Lexington, Kentucky, USA).

Kidney protein extraction for enzyme activity measurements and zymography. For PA activity determination, one-sixth of a kidney is rapidly transferred into a 1.5-ml reaction tube containing 150 µl of ice-cold buffer A (13 mM sodium citrate, 0.33 M sodium acetate, pH 3.9) and manually homogenized on ice for 10 seconds using a potter’s device (1.5 ml Piston Pellet; Treff AG, Degersheim, Switzerland), followed by centrifugation for 5 minutes at 12,000 g at 4°C. Supernatant (100 µl) is then added to 5 µl of 20% acetic acid, gently mixed, and stored at −80°C until further use. For MMP-2 and MMP-9 zymography, one-sixth of a kidney is transferred into 150 µl of ice-cold buffer (10 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1 mM EDTA, 1 mM EGTA) in a 1.5-ml reaction tube and treated as described for PA activity, but without addition of 5 µl of 20% acetic acid. Protein concentrations were determined by the Bio-Rad Protein Assay (Bio-Rad Laboratories, Ivry sur Seine, France).

PA activity and zymography. PA activity present in cell supernatants and kidney extracts was determined using a chromogenic assay (spectrolyse/fibrin; Biopool International, Umea, Sweden). This kit contains a specific EPA stimulator, Desafib-x. Samples were diluted in 0.9% of NaCl before analysis. MMP-2 and -9 activities were determined using zymography. Cell supernatant (20 µl) or kidney extract (25 µg) with nonreducing SDS sample buffer was loaded onto 8–10% SDS-polyacrylamide gels containing 1 mg/ml gelatin. After electrophoresis, gels were soaked two times for 20 minutes in 2.5% (vol/vol) of Triton X-100 at room temperature and incubated overnight at 37°C in activation buffer (10 mM Tris-HCl, pH 7.5, 1.25% Triton X-100 [vol/vol], 5 mM CaCl2, and 1 µM ZnCl2). The following day, gels were stained for 3 hours with 0.25% (wt/vol) Coomassie brilliant blue in a mixture of methanol/acetic acid/water (50/10/40 vol/vol) and destained in the same mixture without Coomassie brilliant blue.

Statistical analysis. All the data are expressed as mean plus or minus SD. Statistical analyses were performed using SPSS software (SPSS Science, Chicago, Illinois, USA). ANOVA with post hoc Tukey alpha test was performed for comparison between the different groups. P values less than 0.05 were considered statistically significant.

Results

UUO-induced tubulointerstitial fibrosis is significantly higher in B2−/− mice. Interstitial collagen deposition was studied histomorphometrically in Sirius red–stained renal sections of B2−/− and B2+/+ mice as an index of the fibrotic response to UUO.

Interstitial fibrosis increased progressively from day 1 to day 14 and became significantly different between the two mouse strains 3 days after obstruction (Figure 1a). At day 5, the difference between B2−/− and B2+/+ mice was the largest (Figure 1b), with an interstitial collagen expression that occupied 35% ± 4% and 17% ± 2% of the kidney sections, respectively. Since this difference was smaller at day 14, and extreme tissue damage was observed at day 14, all further experiments were carried out 5 days after obstruction. No significant difference between both mice strains in tubulointerstitial fibrosis of sham-operated and the contralateral kidney was observed (data not shown). PAS and hematoxylin-stained kidney sections showed an increase in tubular damage characterized by tubular dilation, atrophy, and hypercellularity in the obstructed kidneys, but without differences between B2−/− and B2+/+ mice (not shown).
Further evidence of B2 receptor–mediated reduction of tubulointerstitial fibrosis. To verify that the effect observed in B2−/− mice was effectively due to the absence of B2 receptor activation, we have performed UUO either on transgenic rats exhibiting increased endogenous bradykinin expression caused by overexpression of the human tissue kallikrein gene TGR(hKLK1) (15) or on B2+/+ mice treated with a specific kinin-B2 receptor antagonist (HOE-140).

TGR(hKLK1) rats displayed significantly less UUO-induced tubulointerstitial fibrosis than their non-transgenic littermates (Figure 1c), which was abolished by HOE-140 treatment. Furthermore, HOE-140 also increased UUO-induced interstitial fibrosis in B2+/+ mice (Figure 1d). B2−/− mice, the transgenic rat model, and pharmacological treatment thus strongly suggest that in both mice and rats, bradykinin B2 receptor activation reduces renal fibrosis.

Identification of ECM compounds. We next determined, using immunostaining, expression of ECM compounds known to be increased after UUO (Figure 2). After 5 days of UUO, the expression of fibronectin and collagen I, III, and IV was increased in both mouse strains. However, only collagen type III and IV expression was significantly higher in the B2−/− mice (Figure 2, b and c, respectively).

How does bradykinin modify tubulointerstitial fibrosis? Analysis of mRNA expression and enzyme activity. Current evidence supports the idea that fibrosis of several organs, including the kidney, is a consequence of the disturbance of the balance between ECM synthesis and degradation.

Using Northern blotting, we first studied mRNA expression in kidneys of both B2+/+ and B2−/− mice treated with HOE-140. (Figure 3). B2+/+ mice treated with HOE-140 displayed significantly less UUO-induced interstitial fibrosis than their non-TGR(hKLK1) counterparts, and B2−/− mice treated with HOE-140. (Figure 3b). Collagen expression in kidneys from 5 day–obstructed B2−/− mice treated with saline only or with saline containing HOE-140 at 500 µg/kg/d; n = 10 per group. Data expressed as the mean ± SD percentage of the interstitial area stained; *P < 0.05 compared with sham-operated rats; †P < 0.05 compared with HOE-140–treated B2+/+ mice without difference between B2−/− and B2+/+ mice treated with saline only or with saline containing HOE-140. (Figure 3).

The two main systems involved in ECM degradation are the matrix metalloproteinases and the plasminogen activation system, which are closely interlinked (19). We thus investigated these degrading systems and found that UUO clearly increased mRNA expression of matrix metalloproteinases MMP-2 and MMP-9 and their respective inhibitors TIMP-1 and TIMP-2 (Figure 4), as well as the mRNA expression of tPA, uPA, and the PA inhibitor type 1 (PAI-1) (Figure 4a). However, no difference in mRNA expression was found between B2+/+ and B2−/− mice.

Since it has been reported that bradykinin is a potent stimulus for tPA secretion (20), we have measured total PA activity in sham-operated and obstructed kidneys of both mice strains. In preliminary experiments we have observed that the specific stimulator of tPA) increased PA activity by only three times (not shown), while it is described to increase tPA activity by 300 times. In a control reaction in which we added purified tPA to kidney extracts, Desafib-X increased by 240 ± 15 times PA activity, showing that the lack of increase in PA activity in kidney extracts is probably caused by a low tPA concentration. Non-Desafib-X–stimulated PA activity is thus a mixture of tPA and uPA activity. All reported activities in this study are non-Desafib-X–stimulated total PA activities. UUO induced an increase in PA activity in kidneys of both

Figure 1
Quantitative analysis of UUO–induced renal interstitial collagen expression in B2−/− knockout mice, kallikrein transgenic rats, and B2 receptor antagonist–treated wild-type mice. (a) Collagen expression in obstructed kidneys from B2−/− and B2+/+ mice after different periods of ureteral obstruction (3 hours to 14 days). Data are expressed as the mean ± SD percentage of the interstitial area stained; n = 10; *P < 0.05 compared with respective B2+/+ (same time of obstruction). (b) Representative photographs of Sirius red–stained renal sections of 5 day–obstructed and sham-operated B2+/+ and B2−/− mice. Original magnification ×400. (c) Collagen expression in kidneys from 5 day–obstructed and sham-operated kallikrein transgenic and control (Sprague Dawley) rats. One group was treated with HOE-140. n = 5 per group. *P < 0.05 compared with sham-operated rats; †P < 0.05 compared with UUO kallikrein transgenic rats without HOE-140. (d) Collagen expression in kidneys from 5 day–obstructed B2−/− mice treated with saline only or with saline containing HOE-140 at 500 µg/kg/d; n = 10 per group. Data expressed as the mean ± SD percentage of the interstitial area stained; *P < 0.05 compared with without HOE-140.
B2−/− and B2+/+ mice. However, there was a significantly lower PA activity in the obstructed kidneys of B2−/− mice (Figure 4b), whereas no difference between the two mouse strains in PA activity was detected in the sham-operated kidneys.

This decrease in PA activity in obstructed kidneys from B2−/− mice was paralleled with a lower MMP-2 activity in kidneys of B2−/− mice as assessed by zymography (Figure 4c); no difference between B2+/+ and B2−/− mice was observed for MMP-9 (data not shown).

Since the previous in vivo experiments were based on the difference between the presence or absence of B2 receptors, we have used primary cultures of renal proximal tubular cells obtained from kidneys of B2+/+ mice to verify if bradykinin via B2 receptor activation was able to stimulate PA activity (Figure 5a). Interestingly, we were not able to detect bradykinin-induced PA activity in tubular cells cultured in standard culture medium, whereas lovastatin or TGF-β (positive controls) were able to stimulate PA activity. However, pretreatment of the tubular cells with proinflammatory factors (TNF-α and IFN-γ) resulted in a bradykinin-induced PA activity, which was blocked by the specific B2 antagonist (HOE-140). The increased bradykinin-induced PA activity after pretreatment of cells by the proinflammatory factors was not due to increased bradykinin B2 receptor expression (Western blot, Figure 5b) or coupling (calcium mobilization, Figure 5c). Pretreatment with a nitric oxide inhibitor did not block the bradykinin-induced PA activity (Figure 5a). Bradykinin treatment of proximal renal tubular cells obtained from B2−/− mice did not display increased PA activity under the conditions described above (not shown).

Taken together, these data strongly suggest that bradykinin is reducing UUO-induced tubulointerstitial fibrosis by increasing the activity of enzymes involved in ECM degradation and by decreasing collagen IV mRNA expression.

**Macroglage infiltration is decreased in B2−/− mice.** One of the early events in the development of tubulointerstitial fibrosis is tubulointerstitial inflammation and, more precisely, interstitial macrophage infiltration. In the sham-operated kidney and in the contralateral unobstructed kidney (not shown), a low level of macrophages was present in the tubulointerstitium, but no significant difference was found between both mouse strains. As expected, UUO induced a significant increase in interstitial macrophage infiltration, and, surprisingly, this macrophage infiltration was found to be significantly lower in the obstructed kidney of B2−/− mice compared with B2+/+ mice (Figure 6a).

**Cell proliferation is not modified in B2−/− mice.** Cell proliferation also being a hallmark of obstructive nephropathy, we evaluated renal cell proliferation by immunostaining for PCNA. A very small number of PCNA-positive cells were found in the sham-operated kidney and in the contralateral unobstructed kidney (not shown), but UUO clearly increased both tubular and interstitial PCNA staining in the obstructed kidneys of both mice groups, however, without difference between B2−/− and B2+/+ mice (Figure 6b). Using an Ab that recognizes α-SMC, and thus myofibroblasts, on PCNA adjacent sections (Figure 6c), it was found that the majority of proliferating cells identified with PCNA were myofibroblasts, with no difference between the two mouse strains.
Discussion

It is known that bradykinin is involved in the control of a wide range of physiological functions, including cardiovascular homeostasis (regulation of systemic blood pressure and organ blood flow), water and electrolyte transport, and pain-transmitting mechanisms (21). Besides these well-known physiological effects, bradykinin is involved in inflammation, and increasing clinical evidence suggests that bradykinin could play a role in the beneficial effect of ACE inhibitors in diabetic nephropathy and cardiovascular pathologies (22–24).

In the present study we describe, we believe for the first time and in vivo, by using both genetically engineered animals and pharmacological tools, a bradykinin B2 receptor–dependent renal antifibrotic effect.

**The bradykinin B2 receptor and ECM accumulation.** We have shown that B2-receptor activation in vivo reduced UUO-induced tubulointerstitial fibrosis. Renal fibrosis was more severe in B2+/− mice treated with a B2 receptor antagonist. In contrast, transgenic rats overexpressing kallikrein (thus expressing increased bradykinin levels) displayed less UUO-induced tubulointerstitial fibrosis, an effect that was blocked by B2 receptor antagonist administration. These observations are consistent with the antifibrotic effect of kallikrein overexpression on cardiac fibrosis (15) and with the observation that bradykinin levels are reduced in the obstructed rat kidney (25). By using immunohistochemistry, it was observed that the differences in tubulointerstitial fibrosis observed between B2+/− and B2−/− mice were mainly caused by differences in collagen III and IV expression. Collagen IV was the ECM protein that made up the largest part of the kidney sections compared with the other collagens. This is consistent with previous studies reporting a preferential accumulation of collagen IV in this obstructive nephropathy model (7).

ECM accumulation in pathological states is believed to result from an imbalance between both synthesis and degradation, with increased synthesis and decreased degradation leading to accumulation. The increased levels of ECM proteins observed in obstructed kidneys are consistent with previous studies reporting the accumulation of collagen IV in this obstructive nephropathy model (7).

**The renal plasminogen system and ECM degradation.** We have shown that both tPA and uPA, the proteolytic enzymes involved in ECM degradation, were decreased in obstructed kidneys. The decrease in tPA and uPA activity was accompanied by an increase in PAI-1, the inhibitor of tPA and uPA, suggesting that the decrease in tPA and uPA activity was due to the increase in PAI-1. The activation of MMP-2, a gelatinase involved in ECM degradation, was decreased in obstructed kidneys. The decrease in MMP-2 activity was accompanied by an increase in GAPDH, the reference gene for normalization.

**The role of kallikrein in ECM degradation.** We have shown that kallikrein, the enzyme responsible for the conversion of bradykinin to kinin, was increased in obstructed kidneys. The increase in kallikrein activity was accompanied by a decrease in PAI-1, the inhibitor of tPA and uPA. The decrease in PAI-1 activity was accompanied by an increase in MMP-2, the gelatinase involved in ECM degradation. The increase in kallikrein activity was accompanied by a decrease in MMP-2 activity, suggesting that kallikrein is involved in the degradation of ECM.

**Conclusion.** We have shown that bradykinin B2 receptor activation in vivo reduced UUO-induced tubulointerstitial fibrosis. The decrease in fibrosis was accompanied by an increase in tPA and uPA activity and a decrease in PAI-1 and MMP-2 activity. These observations suggest that bradykinin B2 receptor activation in vivo is a potential therapeutic target for the treatment of obstructive nephropathy.
and degradation (19). We analyzed how bradykinin B2 receptor activation can reduce ECM expression by studying both ECM synthesis and degradation.

First, mRNA expression of 14 proteins that have been shown to be involved/increased in this model (2) was studied. We observed in the obstructed kidney a net increase in mRNA expression of fibronectin, collagen I, III, and IV, TGF-β, MCP-1, osteopontin, MMP-2 and -9, TIMP-1 and -2, as well as the plasminogen activation system (PAI-1, tPA, and uPA). However, no significant difference between B2−/− and B2+/+ mice was observed except for collagen IV mRNA expression that was significantly higher in obstructed kidneys of the B2−/− mice. The latter is consistent with the observed increased collagen IV protein expression. In vitro, bradykinin treatment either increased or decreased ECM production, depending on the cell type used. Bradykinin increased extracellular collagen synthesis by rat vascular smooth muscle cells (increased collagen I mRNA expression and gene-reporter activity, increased TIMP-1 protein expression, and increased TGF-β protein expression) (26). On the other hand, in cultured cardiac fibroblasts, bradykinin decreased ECM production (collagen I, III, and fibronectin mRNA expression) by a mechanism probably involving nitric oxide production (27). Besides our in vivo study, no data are available on the effect of bradykinin on collagen IV (mRNA or protein) expression.

Second, we observed that B2 receptor activation increased PA activity. PA activity was significantly reduced in obstructed kidneys of B2−/− mice, and in primary culture, proximal tubular cells obtained from B2−/− mice pretreated with inflammatory mediators, bradykinin via B2 receptor activation increased PA activity. In our system, PA activity is composed of both tPA and uPA activity. Although, to our knowledge, nothing is known about the effect of bradykinin on uPA activity, bradykinin was found to be one of the most potent stimuli of tPA release in the circulation in both rodents (20) and human vasculature (28). In this context it is important to stress that we determined PA activity and not tPA or uPA release as described above. How bradykinin modifies this PA activity and release is not known. In our study the decreased PA activity in B2−/− mice was not due to increased PAI-1 mRNA expression, since we observed the same increase in both mouse strains, which is con-

Figure 5
(a) Effect of bradykinin (24 hours at 0.1µM) on PA activity from primary culture proximal tubular cells pretreated (24 hours) with TNF-α (150 U/ml) and IFN-γ (500 U/ml) (boxed part of the figure) or non-pretreated. TGF-β, 10 ng/ml; HOE-140, 0.1 µM; N-nitro-l-arginine methyl ester (l-NAME), 0.5 mM; lovastatine, 5 µM. (b) Western blot of B2 receptor (B2R) expression. (c) Increase in intracellular calcium ([Ca 2+]i) following bradykinin (0.1 µM) stimulation (time of stimulation 5 minutes). Values are mean ± SEM of five different experiments. *P < 0.05 compared with control cells. Representative blots or traces are shown.

Figure 6
Immunohistochemical analysis of interstitial macrophage- (a), PCNA- (b), and α-SMC–positive (c) cell expression in kidney sections from obstructed (5 day) and sham-operated kidneys of B2−/− mice. Data are expressed as the mean ± SD; n = 10; *P < 0.05 compared with respective sham-operated mice. ‡P < 0.05 compared with B2+/+ UUO mice.
sistent with an earlier report that bradykinin induces tPA but not PAI-1 release in human vasculature (28). In human vasculature bradykinin-induced tPA release was nitric oxide and cyclooxygenase independent (29). We found in primary culture proximal tubular cells from B2−/− mice that bradykinin-induced PA activity was nitric oxide independent. Another possibility for PA activation/secretion by bradykinin might be a direct interaction between the B2 receptor and the PAs. This was shown recently for the NMDA (N-methyl-D-aspartate) receptor, which was found to directly interact with tPA. NMDA treatment induced tPA release from the NMDA receptor, which resulted in tPA-dependent cleavage of the NR1 subunit of the NMDA receptor, thereby increasing the receptor’s activity (30). Such a direct interaction with the B2 receptor might explain the rapid tPA release upon bradykinin treatment in humans (29).

Third, decreased PA activity in B2−/− mice coincided with decreased MMP-2 activity. Collagen IV is the preferential substrate for MMP-2 (31). Plasmin can transform metalloproteinases from their latent to their active forms. Furthermore, uPA seems to be able to activate MMP-2 by activation of a membrane-bound metalloproteinase (19). The paralleled decrease in PA and MMP-2 activity related to increased ECM expression thus suggests a role for the B2 receptor/PA activity/MMP-2 activity cascade in ECM degradation.

Finally, the increased collagen IV mRNA expression can also be the result of the decreased PA activity in B2−/− mice since this will result in increased fibrin deposition that stimulates collagen accumulation (19).

**Bradykinin and macrophage infiltration.** Ureteral obstruction is primarily a nonimmune stimulus, but results in an interstitial immune infiltrate by the production of tubular-derived chemotactic factors as early as 4 hours after the onset of obstruction (32, 33). This infiltrate (mainly macrophages and lymphocytes) releases cytokines and chemokines, which can induce collagen synthesis by fibroblasts and cause further injury of tubules (2). Surprisingly, kidneys of UUO B2−/− mice exhibited a significantly lower number of interstitial macrophages than B2+/− mice. This reduced number of macrophages in the tubulointerstitium of obstructed kidneys of B2−/− compared with B2+/− mice, accompanied by increased tubulointerstitial ECM deposition, seems to contradict the role of this early event in the development of renal fibrosis after ureteral obstruction. It is, however, consistent with the well-described proinflammatory actions of bradykinin, mainly characterized by its vasodilatory effects and leukocyte accumulation (34, 35). In general, decreased interstitial macrophage infiltration is correlated to reduced tubulointerstitial fibrosis in the UUO model. There are, however, quite a few exceptions in the UUO model in which interstitial macrophage infiltration is not related to or has different effects on the degree of tubulointerstitial fibrosis. A reduction of around 50% of infiltrating macrophages in the UUO model resulted in a reduction of 10% and 25% of tubulointerstitial fibrosis in AT1 receptor-deficient mice (36) and treatment with a chemokine receptor antagonist (37), respectively, showing that the degree of macrophage infiltration was not correlated to the same degree of fibrosis. In another study it was shown that an AT2 receptor antagonist reduced interstitial fibrosis without an effect on interstitial macrophage infiltration (38). Furthermore, in our study reduced interstitial macrophage infiltration was not related to reduced activated fibroblast infiltration, which was observed in studies where reduced macrophage infiltration is accompanied by reduced interstitial fibrosis (37, 39).

The absence of a positive correlation between decreased macrophage infiltration and decreased tubulointerstitial fibrosis in our study might thus have at least two origins: (a) the absence of differences in activated fibroblast infiltration between B2−/− and B2+/− mice, and (b) the reduction of about 35% of infiltrating macrophages in B2−/− mice, which does not compensate the reduced PA activity in this strain, thus resulting in increased tubulointerstitial fibrosis compared with B2+/− mice.

Taken together, the results of this study show that bradykinin B2 receptor activation in vivo reduces UUO-induced tubulointerstitial fibrosis most likely by increasing the activity of a PA/MMP-2 cascade. These observations favor a potential role of bradykinin in the antifibrotic effects of ACE inhibitors.

15. Silva, J.A., et al. 2000. Reduced cardiac hypertrophy and altered blood...


