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Chapter 6

Renal and urinary levels of endothelial protein C receptor correlate with acute renal allograft rejection

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Abstract

The Endothelial Protein C Receptor (EPCR) is expressed on leukocytes, on endothelium of large blood vessels and to a lesser extent on capillaries. Membrane bound EPCR plays an important role in the activation of protein C, which has anticoagulant, anti-inflammatory and cytoprotective effects. After cleavage by a protease, EPCR is also found as a soluble protein. Acute rejection of kidney allografts can be divided in T-cell-mediated rejection (TCMR) and antibody-mediated (ABMR) rejection. The latter is characterized by strong activation of coagulation. Currently no reliable non-invasive biomarkers are available to monitor rejection. Renal biopsies were available from 81 renal transplant patients (33 without rejection, 26 TCMR and 22 ABMR), we had access to mRNA material, matched plasma and urine samples for a portion of this cohort. Renal EPCR expression was assessed by RT-PCR and immunostaining. Plasma and urine sEPCR levels were measured by ELISA. ABMR patients showed higher levels of EPCR mRNA than TCMR patients. EPCR expression on glomeruli was significantly more elevated in ABMR patients than in TCMR or control patients. In the peritubular capillaries, EPCR expression was higher in ABMR patients than in control patients. EPCR expression was higher in tubules and arteries of rejection patients than in control patients. Plasma sEPCR levels did not differ. Urine sEPCR levels were more elevated in the ABMR group than in patients with TCMR or without rejection. ROC analysis demonstrated that urinary sEPCR is appropriate to discriminate between ABMR patients and TCMR or control patients. We conclude that urinary sEPCR could be a novel non-invasive biomarker of antibody mediated rejection in renal transplantation.
Introduction

The Endothelial Protein C Receptor (EPCR) is a type 1 transmembrane glycoprotein, which belongs to the CD1 receptor family (Fukudome and Esmon, 1995). EPCR is primarily expressed on monocytes (Galligan et al., 2001), neutrophils (Sturn et al., 2003), the endothelium of large blood vessels and to a lesser extent on capillaries (Laszik et al., 1997). It binds to the Gla domain of Protein C (PC) (Regan et al., 1997) resulting in a 20-fold increase of the PC activation rate (Taylor et al., 2001). Once activated, Active Protein C (APC) can be released in the circulation or can stay bounded to EPCR. Circulating APC plays an important role as an anticoagulant by proteolytically degrading the coagulation factors Va and VIIIa (Walker and Fay, 1992), which are important co-factors in the intrinsic and common pathways of the coagulation cascade. EPCR bound APC signals through the G protein-coupled Protease-Activated Receptor 1 (PAR-1) (Riewald et al., 2002) and exerts several anti-inflammatory and cytoprotective effects such as inhibition of the release of inflammatory mediators (Brueckmann et al., 2004; Grey et al., 1994; White et al., 2000; Yuksel et al., 2002), down-regulation of the expression of adhesion molecules (Joyce et al., 2001), inhibition of neutrophil and eosinophil migration (Feistritzer et al., 2003; Sturn et al., 2003), anti-apoptotic activities (Joyce et al., 2001; Mosnier and Griffin, 2003) and the protection of endothelial barrier function (Feistritzer and Riewald, 2005; Finigan et al., 2005).

The soluble form of EPCR (sEPCR), resulting from the cleavage of the extracellular domain of the membrane bound EPCR (mEPCR) by a metalloprotease (Xu et al., 2000), can decrease the activation of PC by competing with mEPCR for PC (Xu et al., 2000). sEPCR also inhibits APC anticoagulant activity by blocking the interaction with negatively charged membranes (Liaw et al., 2000), an interaction that is necessary for effective inactivation of coagulation factors Va and VIIIa.

Currently the role of the EPCR/APC complex in renal transplantation is unknown; however APC has been extensively studied in inflammation settings and in sepsis. For example, Gupta et al. showed increased renal injury in rats with acquired PC deficiency in a polymicrobial sepsis model (Gupta et al., 2007) and Keller et al. discovered that treatment with APC attenuates inflammation and preserves renal function during sepsis in rats (Keller et al., 2011).
There are two types of acute allograft rejection that can occur either separately or together: T-cell-mediated rejection (TCMR) and acute antibody-mediated rejection (ABMR). TCMR is the most common form of acute allograft rejection, caused by effector T-cells that infiltrate and proliferate in the graft (-draining lymph nodes) leading to graft rejection (Nankivell and Alexander, 2010). ABMR is caused by donor-specific antibodies and is characterised by histological changes such as leukocyte infiltration in the glomeruli and peritubular capillaries (PTC), tubular necrosis, congestion of PTC, infiltration of granulocytes, endothelial cell damage and finally fibrinoid arterial necrosis (Colvin, 2007). Damaged endothelial cells release injury molecules such as cytokines, chemokines, von Willebrand factor and P-selectin, which can induce leukocyte adhesion and activation of the complement and coagulation cascade (Nankivell and Alexander, 2010). As a result of activation of the complement cascade during ABMR currently one of the most reliable surrogate markers for ABMR is C4d positivity of PTC (Colvin, 2007), which requires having access to biopsy material, involving an invasive procedure for the patient. In the clinical setting it is important to distinguish between patients with TCMR and ABMR, because the treatments of these two types of rejection are different (Nankivell and Alexander, 2010).

In the current study we investigate the EPCR expression pattern in kidney transplants on both mRNA and protein level; and correlate plasma and urine sEPCR levels upon acute renal allograft rejection. We describe how urinary sEPCR can distinguish between ABMR and TCMR.

**Materials & Methods**

**Patients**

Eighty-one patients who underwent kidney transplantation between 1994 and 2008 were retrospectively selected from the patient population of the Academic Medical Center at the University of Amsterdam. Patients were selected based on pathological diagnosis. Renal biopsies that fulfilled the minimal criteria for diagnostic assessment (7 glomeruli and at least 1 artery) according to the Banff 1997 criteria were available from all patients (Racusen et al., 1999). All biopsies were stained for Haematoxylin-Eosin, Periodic Acid Schiff Diastase and Methenamine Silver (Jones) and scored.
following the Banff 2007 guidelines (Solez et al., 2008). Patients were divided in 3 groups, according to the biopsy diagnosis. The TCMR group consisted of 26 patients with interstitial infiltration, tubulitis or intimal arteritis (Racusen et al., 1999). Twenty patients with acute-tubulus-necrosis-like inflammation, capillary and/or inflammation and/or thrombosis, arteritis and C4d tubular deposition were assigned to the ABMR group (Racusen et al., 1999). In addition, two patients with signs of ABMR-associated microvascular inflammation (Sis et al., 2012) with negative C4d staining were assigned to the ABMR group. Finally, the control group consisted of 33 patients without signs of rejection. Kidney material of the control group came from protocol biopsies and showed no signs of inflammation or rejection. Time-matched mRNA, derived from frozen transplant biopsies, was available for 14 patients in the control group, 13 in the TCMR group and 11 in the ABMR group as well as matched serum samples for 21 patients in the control group, 22 in the TCMR group and 9 in the ABMR group. Matched urine samples (taken from 24 hours urine samples) were available for 22 patients in the control group, 21 in the TCMR group and 8 in the ABMR group. Circulating donor specific antibodies are not systematically measured in kidney transplant patients in our institution. Consequently, according to the Banff criteria (Solez et al., 2008), we considered a patient belonging to the ABMR group when showing a diffuse C4d staining of the peritubular capillaries on frozen section. If frozen sections were not available, we performed C4d staining on paraffin sections, patients with more than 50% of positive peritubular capillaries were considered as positive for ABMR. Seven patients who showed signs of both TCMR and ABMR were included in the ABMR group. All biological material was collected for previous studies for which written informed consent was given by all patients for their information to be stored in the hospital database and to be used for research. This research project used left-over biological material, anonymised and delinked from patient records, and as such was not subject to any requirement for ethical review or approval.

**RNA extraction and processing for real-time PCR**

mRNA from complete kidney tissue was extracted from frozen renal biopsies cut into 25 mm thick sections using a Microm HM500 cryostat (Adamas Instruments BV) and collected in an Eppendorf tube containing TRIzol (Invitrogen, Breda, The Netherlands). After 5 minutes incubation at room temperature RNA was extracted
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using chloroform. cDNA was synthesised using a standard procedure. Real-time reverse-transcriptase polymerase chain reaction (RT-PCR) was performed on a Lightcycler® 480 Real-Time PCR System using Lightcycler® 480 SYBR Green I Master (Roche Applied Science, Mijdrecht, the Netherlands). Specific primers were designed (synthesized by Eurogentec, Liège, Belgium) for human PROCR (EPCR) (forward ACCTTGGCCTTTCCCTCTGAC, reverse CTCCCATTACAGCCACTTC). Results were analyzed using LinRegPCR 12.4 software (Heart Failure Research Center, Academic Medical Center, Amsterdam, the Netherlands). PROCR gene expression was normalized against two different housekeeping genes: HPRT1 (hypoxanthine phosphoribosyltransferase 1) and ACTBL2 (beta-actin). Comparable results were obtained with the two housekeeping genes. We decided to present the ratios between Ct values of PROCR and HPRT1 (forward TTGTTGGATATGCCCTTGACT, reverse CCGCTGTCTTTTAGCTTG).

Immunostaining

C4d staining was performed on frozen sections if available using a mouse anti-human C4d antibody (AbD Serotec, Dusseldorf, Germany, ref. 2222-8004). Alternatively, C4d staining was performed on paraffin sections using a rabbit anti-human C4d antibody (Cell Marque, Rocklin, USA, ref. 404A-14), as described previously (Scheepstra et al., 2008). Paraffin sections of kidney biopsies were immunostained for EPCR. Antigen retrieval with Tris EDTA pH 9 (20 minutes at 121°C) was performed for optimal staining. 4 mm thick sections were incubated for 60 hours at 4°C with goat anti human EPCR monoclonal antibody (452 ng/mL, kind gift of Dr. C. Esmon, Oklahoma Medical Research Foundation). Primary antibody binding was detected with a peroxydase kit (30 minutes incubation at room temperature, Powervisoin poly HRP-Anti-mouse IgG, Immunologic, Duiven, Netherlands). Staining was developed with Ultra DAB (Immunologic, Duiven, Netherlands). On EPCR stained sections, the intensity of immunostaining was scored semiquantitatively on a scale from 0–3 (respectively absent, weak, moderate or strong) following the method of Faust and colleagues (Faust et al., 2001). Intensity of staining was evaluated in five kidney substructures: glomeruli, peritubular capillaries, arteries, veins and tubules. All sections were coded and scored by two blinded investigators. For each section a mean score was calculated from at least three high power fields. C4d staining was evaluated following the Banff 2007 recommendations (Solez et al., 2008) by a blinded pathologist.
Enzyme-linked immunosorbent assay (ELISA)
Twenty mL aliquots of urine and plasma were pre-treated with 10 mL 1N HCl. After 10 minutes of incubation at room temperature acidified samples were neutralized with 9 mL 1N NaOH. Prior to the assay samples were diluted with Calibrator Diluent (RD5–24 provided with the ELISA kit) to reach a total dilution factor of 15.6 for urine samples and 39 for plasma samples. Soluble EPCR concentrations in urine and plasma were measured using Human EPCR Quantikine kit (R&D System, Abingdon, UK) according to the manufacturer’s protocol. According to the data sheet the mean minimal detectable doses is 0.064 ng/mL. Optical densities were measured using a microplate reader set to 450 nm and were corrected with a wavelength of 570 nm. A standard curve was created using the trial version of MasterPlexH ReaderFit software (Hitachi Solution America Ltd.), capable of generating a four parameter logistic curve fit. The urinary concentration of sEPCR was corrected for the urine dilution by dividing the urinary concentration of sEPCR by the urinary creatinine concentration (urine creatinine concentration did not differ between the groups, data not shown).

Glomerular filtration rate (GFR)
The GFR was estimated with use of the CKD-EPI formula (Levey et al., 2009).

Statistical analysis
All data sets were tested for their distribution prior to analyses. Data are expressed as median and range unless stated otherwise. Wilcoxon-Mann Whitney test, Kruskal Wallis, Spearman’s correlation test, the multivariate analysis and Receiver Operating Characteristic (ROC) analyse were performed using SPSS 19 software (IBM Corporation, Stomer NY USA) and the R computing environment (www.r-project.org). Overall a two-tailed P-value of <0.05 was considered significant.

Results
Demographic and clinical characteristics of the patients
Table 1 shows the demographic and clinical characteristics of the patients. The three groups differed in some aspects. The patients in the ABMR group are slightly younger than those in the control group (P < 0.05). As expected, the median serum
creatinine concentration was higher and the estimated glomerular filtration rate (GFR) was lower in patients undergoing renal allograft rejection compared to patients without rejection (P < 0.001). More HLA mismatches were present in patients with ABMR compared with control (P < 0.001) and TCMR groups (P < 0.05) (Table 1). In our hospital, it was not standard care to evaluate donor-specific antibodies at time of rejection. However, panel reactive antibody (PRA) measurements were performed before transplantation, at the time of transplantation, at time of biopsy and after the biopsy. PRA levels did not change significantly over time within the 3 patient groups (multivariate analysis, Figure 1). At time of biopsy however, the ABMR group showed significantly higher levels of PRA compared to control patients (P < 0.001) and patients undergoing T-cell-mediated rejection (P < 0.05, Figure 1 and Table 1).

All patients received immunosuppressive treatment consisting of CD25mAb (induction), corticosteroids, mycophenolate and a calcineurin inhibitor. Acute cellular rejections were treated with pulse doses of methylprednisolone 500 mg iv
for 6 days. Antibody mediated rejections were treated with plasmapheresis for 7 days and rabbitATG (rATG). The starting dose of rATG was 5 mg/kg, 3 to 5 gifts were administered over 14 days. Dosages were titrated based on the total lymphocyte count after each administration (> 300*10^9/L: dose 5 mg/kg; > 200*10^9/L but < 300*10^9/L: dose 3 mg/kg; > 150*10^9/L but < 200*10^9/L: dose 2 mg/kg; < 150*10^9/L: no administration). Table 2 shows the Banff scores of the patients. TCMR patients had more tubulitis and mononuclear cell interstitial inflammation than ABMR patients (P < 0.05). 86% of the biopsies diagnosed as ABMR were C4d positive (P < 0.001 versus TCMR and control).
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Figure 2 | EPCR mRNA expression in kidney transplant. EPCR mRNA levels were measured with qPCR on whole kidney biopsies. Results are shown as ratio between EPCR and HPRT Ct. Antibody-mediated (ABMR) rejection patients showed higher levels of EPCR mRNA than T cell-mediated rejection (TCMR) patients. Results are shown as median, interquartile range and range. ** P < 0.001 (Mann-Whitney Test).

Table 2 | Banff characteristics of the included patients

<table>
<thead>
<tr>
<th>Banff parameters</th>
<th>Control group</th>
<th>ABMR group</th>
<th>TCMR group</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Acute scores</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tubulitis (t)</td>
<td>0 (0 - 1)</td>
<td>1 (0 - 3)</td>
<td>1 (1 - 3)</td>
</tr>
<tr>
<td>Interstitial inflammation (i)</td>
<td>0 (0 - 1)</td>
<td>1 (0 - 3)</td>
<td>2 (1 - 3)</td>
</tr>
<tr>
<td>Glomerulitis (g)</td>
<td>0 (0 - 2)</td>
<td>1 (0 - 3)</td>
<td>1 (0 - 3)</td>
</tr>
<tr>
<td>Peritubular capillaritis (ptc)</td>
<td>0 (0 - 2)</td>
<td>0 (0 - 2)</td>
<td>0 (0 - 2)</td>
</tr>
<tr>
<td>Intimal arteritis (v)</td>
<td>0 (0 - 0)</td>
<td>0 (0 - 3)</td>
<td>0 (0 - 2)</td>
</tr>
<tr>
<td><strong>Chronic scores</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Arteriolar hyalinosis (ah)</td>
<td>0 (0 - 2)</td>
<td>0 (0 - 2)</td>
<td>0 (0 - 2)</td>
</tr>
<tr>
<td>Vascular intima thickening (cv)</td>
<td>0 (0 - 1)</td>
<td>1 (0 - 2)</td>
<td>0 (0 - 3)</td>
</tr>
<tr>
<td>Transplant glomerulopathy (cg)</td>
<td>0 (0 - 2)</td>
<td>0 (0 - 3)</td>
<td>0 (0 - 1)</td>
</tr>
<tr>
<td>Mesangial matrix increase (mm)</td>
<td>0 (0 - 1)</td>
<td>0 (0 - 3)</td>
<td>0 (0 - 3)</td>
</tr>
<tr>
<td>Interstitial fibrosis (ci)</td>
<td>0 (0 - 2)</td>
<td>0 (0 - 2)</td>
<td>0 (0 - 3)</td>
</tr>
<tr>
<td>Tubular atrophy (ct)</td>
<td>1 (0 - 2)</td>
<td>1 (0 - 2)</td>
<td>1 (0 - 3)</td>
</tr>
<tr>
<td><strong>Complement activation</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>N° C4d positive</td>
<td>0 (0%)</td>
<td>19 (89%)</td>
<td>0 (0%)</td>
</tr>
</tbody>
</table>

Data are shown as median (range). 1, 2 P < 0.05), 3, 4 P < 0.001 (Mann Whitney test).
EPCR mRNA levels in kidney transplant biopsies
As shown in Figure 2, PROCR/HPRT1 (EPCR) mRNA ratios were significantly higher in patients with ABMR (0.49 [0.29 – 0.64]) compared to those in patients with TCMR (0.26 [0.23 – 0.45], P < 0.001) but not compared with control patients (0.33 [0.20 – 0.68], P > 0.05).

**Figure 3 | EPCR expression patterns in transplant biopsies.** Representative immunostainings of kidney biopsies for EPCR in glomeruli (x32 magnification), peritubular capillaries (x64), arteries (x64), veins (x64) and tubules (x64). Arteries were always positive; therefore no picture with a score of 0 is shown. For tubules no score of 3 was assigned.
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**EPCR protein expression in kidney transplant**

In order to visualize the expression pattern of EPCR in transplant biopsies, we performed immunostainings. Figure 3 shows representative EPCR staining patterns in kidney transplant biopsies. Intensity of staining was evaluated in five kidney substructures: glomeruli, peritubular capillaries, arteries, veins and tubules on a semi quantitative scale from 0 to 3. In general, arteries were more intensely stained than other substructures. Therefore no score of 0 was assigned for arteries. Intensity of staining was generally weaker in tubules than in other kidney compartments, no staining intensity corresponding to a score of 3 was assigned for tubules.

EPCR expression was significantly higher in patients with ABMR compared to patients without rejection or with TCMR in glomeruli (P < 0.001 and P < 0.05, respectively, Figure 4A). In capillaries ABMR patients showed higher EPCR expression only compared to control (P < 0.05, Figure 4B). In arteries we observed higher EPCR expression in patients with ABMR and TCMR compared to patients without rejection (P < 0.05, Figure 4C). The same expression pattern was observed in tubules (P < 0.05, Figure 4D). EPCR scores of the venous endothelium did not show any differences between ABMR, TCMR and control (data not shown).

**Plasmatic and urinary sEPCR concentration**

Plasma and urine levels of sEPCR were determined by ELISA. The urinary concentration of sEPCR was corrected for dilution. We confirmed the presence of intact sEPCR protein, and not degradation products by western blot analysis (data not shown). Plasma sEPCR levels were not significantly different between ABMR patients (599 ng/mL [67 - 1355]), control patients (623 ng/mL [418 – 1102], p > 0.05) or patients with TCMR (508 ng/mL [381 – 945], P > 0.05) (Figure 5A). Conversely, the urine levels of sEPCR were significantly higher in patients with ABMR (29 ng/mmol creatinine [9 – 137]) than in either patients with TCMR (12 ng/mmol creatinine [5 – 50], P < 0.05) or without rejection (13 ng/mmol creatinine [3 – 30], P < 0.01, Figure 5B). The urine creatinine concentration did not differ between the groups (P = 0.6). We found no correlation between protein EPCR scores in the kidney and sEPCR concentration in plasma or urine (data not shown).
Association of clinical parameters and Banff scores with EPCR

Recently, Sis et al. (Sis et al., 2012) proposed that the cumulative Banff score for glomerulitis and peritubular capillaritis (g + ptc) associates with antibody-mediated inflammation of the microcirculatory circuit, irrespective of C4d positivity. Indeed, C4d-negative ABMR is an increasingly recognized entity. We investigated whether little (g + ptc ≤ 3) or severe (g + ptc > 3) microcirculatory inflammation related to higher levels of urine sEPCR or staining intensity of EPCR in these renal structures. Glomerular EPCR (P = 0.009) and the composite of glomerular and peritubular
capillary EPCR scores (P = 0.009) significantly associated with severe microcirculatory inflammation (g + ptc > 3). Higher scores for EPCR on the peritubular capillaries tended to relate to higher microcirculatory inflammation as well (P = 0.09). Not for each biopsy sample, a matching urine sample was available. Therefore we imputed the missing values with multivariate bootstrap methods. Complete case analysis showed a trend for higher urine sEPCR levels at the time of biopsy when severe microcirculatory inflammation was present (P = 0.06) (Figure 6).

Receiver operating characteristics (ROC) analysis
In order to evaluate the possible usefulness of urinary sEPCR as a non-invasive biomarker for ABMR, a ROC calculation was performed. The area under the ROC curve was used to summarize the discriminative ability of the test, the closer to 1 the better. As shown in Figure 7, the sEPCR concentration in urine can be used to discriminate ABMR patients from TCMR patients (area under the curve of 0.875, P < 0.01) and patients without rejection (area under the curve of 0.8785, P < 0.01). The cut-off value with the highest combined sensitivity and specificity for discriminating between ABMR patients and control patients was at 21.6 ng/mmol creatinine (75% sensitivity, 80% specificity). The optimum sEPCR concentration for discriminating between ABMR and TCMR patients was 22.1 ng/mmol creatinine (75% sensitivity, 82% specificity). The dotted line represents the line of equal sensitivity and specificity.
Discussion

This study is the first to describe the levels of membrane bound and soluble EPCR during acute renal allograft rejection in a transplant patient cohort.

We observed higher EPCR mRNA levels in ABMR patients compared with TCMR patients. On the protein level, we observed in glomeruli more EPCR expression in ABMR patients compared with patients without rejection or with TCMR. In the peritubular capillaries EPCR expression was higher in ABMR patients than in control patients. Immunostaining also revealed a higher expression of EPCR in arteries and tubules from patients with ABMR and TCMR compared with patients...
without rejection. Finally we found increased concentrations of sEPCR in the urine of ABMR patients compared with patients with TCMR or without rejection. Although validation is needed, these findings together with the ROC analysis indicate that urinary sEPCR could be used as a non-invasive biomarker for antibody mediated kidney rejection.

There are some limitations to our study. Since it is a retrospective study some patients could not be studied for all the different tests. Furthermore the ABMR group contains a relatively small number of patients with urine sample available. The reason for this is the low incidence of ABMR within the renal transplant patient population in our institution. Nevertheless even with a population of 8 samples from the ABMR group and 21 from the TCMR group the ROC analysis achieves 80% power to detect a difference of 0.422 between the area under the curve under the

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**Figure 7 | ROC analyses, graphical representation.** Receiver operating characteristic (ROC) analysis of urine sEPCR concentration for the prediction of ABMR in kidney transplantation. ROC analysis demonstrated that urinary sEPCR is appropriate to discriminate between ABMR patients and TCMR or control patients. In grey: ABMR vs. Control, in black: ABMR vs. TCMR.
null hypothesis of 0.500 and an AUC under the alternative hypothesis of 0.875 using a two-sided z-test at a significance level of 0.05.

Another limitation of our study may be the lack of data on donor specific antibodies (DSA). ABMR diagnosis can be complex in the clinical setting, often requiring DSA. Indeed, C4d-negative ABMR is an increasingly recognized phenomenon (Sis et al., 2012). DSA are not routinely determined in our institute. Instead, we have incorporated the less specific PRA data in our study, showing that ABMR patients have higher percentages of PRA than TCMR patients at time of biopsy, i.e. at time of the acute rejection episode.

It is known from the literature that the expression of EPCR is not only limited on endothelial cells. On the mRNA level EPCR is transcribed in HUVEC (Gao et al., 2009), in a human alveolar epithelium cell line (Wang et al., 2007), in rat lung (Lu et al., 2009), pancreatic tissue (Ping et al., 2010), in gastric epithelial cells (Nakamura et al., 2005) and in immortalized human proximal tubular epithelial cell line (Bae et al., 2010). In this study we show for the first time EPCR mRNA in kidney allograft tissue. On the protein level EPCR is known to be expressed on HUVEC (Gao et al., 2009), endothelium in the heart, the lung, the kidney, the skin and on other organs (Laszik et al., 1997) but also on epithelial cells such as a prostatic cancer cell line (Menschikowski et al., 2011), human alveolar epithelium (Wang et al., 2007), rat lung (Lu et al., 2009), pancreatic tissue (Ping et al., 2010) and gastric epithelial cells (Nakamura et al., 2005). In normal renal tissue, EPCR is expressed on vein and arterial endothelium (Laszik et al., 1997). Although the expression of EPCR by tubular epithelial cells has been described in a cell line of artificially immortalized kidney cells (Bae et al., 2010), our study is the first to actually demonstrate EPCR expression by tubular epithelial cells in human tissue.

The roles of EPCR and protein C have been extensively studied in sepsis setting. EPCR and APC form a complex with PAR-1 and initiate biological effects such as anticoagulant, anti-inflammatory, anti-apoptotic and cytoprotective activities in vitro and in vivo (Brueckmann et al., 2004; Feistritzer and Riewald, 2005; Feistritzer et al., 2003; Finigan et al., 2005; Grey et al., 1994; Joyce et al., 2001; Mosnier and
Griffin, 2003; Riewald et al., 2002; Sturn et al., 2003; White et al., 2000; Yuksel et al., 2002). The APC/EPCR/PAR-1 axis inhibits the release of inflammatory mediators (Brueckmann et al., 2004; Grey et al., 1994; White et al., 2000; Yuksel et al., 2002), down-regulates the expression of adhesion molecules (Joyce et al., 2001), inhibits neutrophil and eosinophil migration (Feistritzer et al., 2003; Sturn et al., 2003) and exerts protective activities against infection. APC-EPCR-PAR1 interaction has been shown to play a protective role in several conditions such as in a sepsis model using e. coli (Taylor et al., 2000) and was shown to have anti-apoptotic activities (Joyce et al., 2001; Mosnier and Griffin, 2003) and protect endothelial barrier function (Feistritzer and Riewald, 2005; Finigan et al., 2005). Importantly, Song et al. showed an up-regulation of EPCR expression in the kidney of mice injected with LPS (Song et al., 2009).

These findings clearly indicate that EPCR is part of a protective pathway. Considering this we assume that the higher EPCR expression during acute kidney rejection could be part of a protective mechanism for the graft. EPCR could exert its protective activities on several levels.

Firstly, higher EPCR expression could contribute to enhanced APC formation and therefore more activation of PAR-1 and its associated protective properties. This could suggest that an actively regulated protection mechanism involving the EPCR/ APC/PAR-1 signaling cascade is taking place in the setting of acute allograft rejection. Secondly, it is known that ABMR is characterized by activation of the coagulation cascade, resulting in elevated levels of FVIIa (Nankivell and Alexander, 2010), which can bind to EPCR and activates its associated protective activities (Sen et al., 2011).

EPCR not only exists as a membrane bound receptor but also as a soluble protein. sEPCR has a comparable affinity for APC as mEPCR (Liaw et al., 2000). sEPCR can compete with mEPCR for APC, resulting in a decreased activation of PC (Xu et al., 2000). sEPCR also inhibits APC anticoagulant activity by blocking the interaction with negatively charged membranes (Liaw et al., 2000), a necessary interaction for an effective inactivation of coagulation factors Va and VIIIa. Currently, very little is known about sEPCR in transplantation. The only study on sEPCR in transplantation
was performed by Keven et al., who measured plasma sEPCR concentration before and after kidney transplantation in patients without rejection (Keven et al., 2010). Three months after transplantation the serum sEPCR concentration was significantly lower than before the transplant procedure. From this, Keven et al. concluded that the elevated pre-transplant sEPCR levels reflected endothelial damage, due to the hemodialysis regimen before transplantation (Keven et al., 2010). Elevated levels of sEPCR have been described in vasculitis and SLE, which has fostered the hypothesis that sEPCR may be a biomarker for endothelial dysfunction (Sesin et al., 2005).

Considering this, it is not surprising to observe higher sEPCR production during rejection, knowing that allograft rejection is associated with endothelial damage and dysfunction especially in antibody-mediated rejection (Nankivell and Alexander, 2010). Until now sEPCR concentration was only measured in plasma, in this study we are the first to show sEPCR presence in the urine of patients. The origin of the urinary sEPCR is unclear. The relatively low molecular weight of sEPCR [about 43 kDa (Kurosawa et al., 1997)] makes its filtration through the glomeruli theoretically possible. Therefore sEPCR could be derived either from filtrated blood or from shed tubular mEPCR. Shedding of monocytes or neutrophils bound EPCR might also be an alternative source of urinary sEPCR.

Renal transplantation is the most suitable therapy for end stage kidney disease. Despite recent progress in anti-rejection therapy, approximately 23% of transplanted patients undergo an episode of acute rejection within the first year post-transplant (National Institutes of Health, 2010). For all patients undergoing a rejection episode it is crucial to have the ability to correctly diagnose the presence of rejection. Therefore having access to a convenient biomarker could be of interest for the clinical practice. Currently, sampling of a kidney biopsy and its histological examination is the gold standard for diagnosing both TCMR and ABMR, causing discomfort for the patient.

Therefore we propose that urinary sEPCR could be of interest for diagnosing acute kidney antibody-mediated rejection as proven by the ROC analysis, which demonstrates that urinary sEPCR might be suitable to make the distinction between
ABMR patients and non-rejecting patients (AUC=0.875, P = 0.002) and importantly also between ABMR and TCMR patients (AUC=0.875, P = 0.003). Although validation will be needed in a larger patient cohort, we conclude that urinary sEPCR could be a suitable candidate to diagnose antibody mediated renal rejection in the clinical setting, in a non-invasive way.