Microvascular barrier dysfunction plays a major role in the pathophysiology of acute kidney injury (AKI). Angiopoietin-1, the natural agonist ligand for the endothelial-specific Tie2 receptor, is a non-redundant endothelial survival and vascular stabilization factor. Here we evaluate the efficacy of a polyethylene glycol-clustered Tie2 agonist peptide, vasculotide (VT), to protect against endothelial-cell activation with subsequent microvascular dysfunction in a murine model of ischemic AKI. Renal ischemia reperfusion injury (IRI) was induced by clamping of the renal arteries for 35 minutes. Mice were treated with VT or PEGylated cysteine before IRI. Sham-operated animals served as time-matched controls. Treatment with VT significantly reduced transcapillary albumin flux and renal tissue edema after IRI. The protective effects of VT were associated with activation of Tie2 and stabilization of its downstream effector, VE-cadherin in renal vasculature. VT abolished the decline in renal tissue blood flow, attenuated the increase of serum creatinine and blood urea nitrogen after IRI, improved recovery of renal function and markedly reduced mortality compared to PEG [HR 0.14 (95% CI 0.05–0.78) \( P < 0.05 \)]. VT is inexpensive to produce, chemically stable and unrelated to any Tie2 ligands. Thus, VT may represent a novel therapy to prevent AKI in patients.
that protects the vasculature of adult mice from plasma leakage induced by vascular endothelial growth factor (VEGF) and other inflammatory stimuli. In contrast, release of Angiopoietin-2 (Angpt2) from endothelial Weibel-Pallade bodies disrupts the constitutive Angpt1-Tie2 signaling by preventing Angpt1 from binding to the receptor, thereby promoting inflammation and permeability.

Recent studies in animals and humans have demonstrated that renal ischemia induces a transient imbalance between renal Angpt1 and -2 expression in favor of Angpt2. During reperfusion renal vascular endothelium acutely releases pre-stored Angpt2. Given the absence of redundant systems to bypass the function of Tie2, it was speculated that exogenous repletion of Angpt1 rescues Tie2 signaling and thus effectively abolishes microvascular leakage in ischemia-reperfusion injury (IRI). Indeed, overexpression of an engineered Angpt1 variant in mice attenuates renal IRI and pretreatment of donor hearts with this Angpt1 variant or with an inhibitor of Angpt2 improves IRI after heart transplantation in rats. Thus, pharmaceutical agents that activate the Tie2 receptor pathway leading to vascular stabilization and protection against microvascular dysfunction in AKI would be highly desirable. In this study we tested the effects of the novel Tie2-agonist tetrameric peptide, Vasculotide (VT), in a murine model of renal IRI.

**Results**

**Vasculotide activates Tie2 signalling and protects vascular barrier in ischemic kidneys.** Figure 1 shows the VT (Supplementary Fig. S1) induced changes of Tie2 expression and phosphorylation in kidney homogenates 24 h after IRI. Mice received 200 ng VT (or molar equivalent of PEGylated cysteine (PEG)) intraperitoneally at -16 h and -1 h prior to IRI or sham surgery, respectively. VT pretreatment almost completely prevented Tie2 de-phosphorylation (Fig. 1A). To resolve the impact of the changes in Tie2 activation we next surveyed vascular leakage in the kidney of these mice. The anti-leakage effect of VT on the kidney after IRI was already macroscopically visible (Fig. 1B). IRI caused a significant increase in transcapillary leakage, which was significantly reduced by VT pretreatment (Fig. 1C and D). Immunofluorescence for VE-Cadherin and the endothelial marker CD31 at 24 h after IRI or sham operation revealed preserved endothelial integrity in VT-pretreated kidneys (Fig. 1E).

--

Figure 1. Effects of Vasculotide on Tie2 signaling, capillary leakage and endothelial integrity. Mice (n = 8 per group) were pre-treated with 200 ng of Vasculotide (VT) or pegylated cystein (PEG) i.p. at -16 h, -1 h prior to renal ischemia reperfusion injury (IRI) or sham surgery. (A) Immunoblots for phosphoTie-2Y992 (pTie2) and total Tie2 (Tie2) in kidney homogenates obtained 24 h after renal IRI. VT: Vasculotide; PEG: PEG-Cys (control); β-Tub: β-Tubulin. (B) Macroscopic aspect of Evans blue dye in kidneys, (C) Evans blue dye content and (D) Wet-to-dry-ratio in kidney homogenates 24 h after IRI or sham operation. OD: optical density. (E) Immunofluorescence for VE-Cadherin and the endothelial marker CD31 at 24 h after IRI or sham operation. VECad: VE-Cadherin. Data are means ± SEM. *P < 0.05; **P < 0.01 vs. IRI + PEG-Cys. +P < 0.05; ++P < 0.01; +++P < 0.001 vs. sham.
Vasculotide improves kidney function and reduces mortality after renal IRI. Bilateral IRI resulted in a significant increase in serum creatinine and blood urea nitrogen (BUN) levels in PEG-treated controls. Treatment with VT was sufficient to attenuate ischemic AKI, as evidenced by significantly lower serum creatinine and BUN levels in VT-treated mice compared to controls (P < 0.05), especially at later time points (day 3 and day 7 after IRI) (Fig. 5A,B and Supplementary Fig. S3).

Lastly, we performed survival experiments, the benchmark of any successful treatment regimen. Mice received an additional dose of 200 ng VT (or molar equivalent PEG) i.p. 24 h post to sham or surgery in addition to standard VT-pretreatment. As a result, only 38% of mice pre-treated with PEG survived, compared to 92% in the VT group, respectively. When compared to the PEG group, pre-treatment with VT was significantly associated with better outcome [Hazard Ratio 0.14 (95% confidence interval 0.05–0.78)] (P < 0.05) (Fig. 5C).

Discussion

Although AKI is a frequent complication in the intensive care unit (ICU) that is associated with increased mortality and morbidity, no specific treatments are available. Current understanding of the pathophysiology of AKI...
emphasizes the complex interplay between inflammation and ischemia-reperfusion leading to microcirculatory dysfunction and localized hypoperfusion and ischemia of the kidney. Here we used a reliable model of AKI to study the impact of VT on kidney function and mortality. We show that VT counteracts microvascular barrier dysfunction of the renal microcirculation in renal IRI. VT activated Tie2 and its major downstream pathway VE-cadherin in the kidney, prevented IRI-induced renal vascular leakage and congestion, improved renal function and recovery, and significantly attenuated mortality after renal IRI. These effects were independent of changes in systemic hemodynamics and tubular injury. Together, the present data suggest that Tie2 activation by VT is sufficient to preserve endothelial barrier function and to confer protection against ischemic AKI.

We and others have shown that Tie2 protein abundance, Tie2 activation, and downstream signaling are suppressed in murine lungs and kidneys during endotoxic and hemorrhagic shock. Recently David et al. demonstrated that VT is able to overcome the decreased Tie2 protein abundance during endotoxemia by increased activation of the remaining Tie2 molecules. Tie2 expression, however, has not been addressed in renal IRI. We found marked loss of phosphorylated Tie2 in kidneys subjected to IRI. As in septic lungs, VT was able to increase Tie2 phosphorylation despite loss of total Tie-2 protein, thereby preserving net flux through this signaling pathway. In a proof-of-principle study, Jung et al. demonstrated that adenoviral overexpression of cartilage oligomeric matrix protein-angiopoietin-1 (COMP-Angpt1), an engineered variant of native Angpt1 with higher activity, attenuates renal IRI in mice. Although the current study basically corroborates their findings, some contrasting results deserve further discussion. First, overexpression of COMP-Angpt1 ameliorated tubular function.

Figure 2. Vasculotide improves renal perfusion. Mice were pre-treated with 200 ng of VT or PEG i.p. at −16 h and −1 h prior to renal IRI (n = 7 per group) or sham surgery (n = 4 per group). Change of cortical (A) and medullary (B) perfusion parameters (contrast enhanced ultrasound) 1 h after renal IRI. Every mouse was examined immediately before and 1 h after IRI or sham operation. Renal vascular resistance 1 h after IRI was calculated as follows: RVR = MAP/RBF. VT: Vasculotide; PEG: PEG-Cys (control); WiR: wash in rate; TTP: time to peak; RBF: renal blood flow; RVR: renal vascular resistance. Data are expressed as means ± SEM. *P < 0.05 vs. IRI + PEG-Cys. **P < 0.05; ***P < 0.01 vs. Sham.

Table 2. Systemic hemodynamics before and 1 h after renal IRI or sham operation. bpm: beats per minute; MAP: mean arterial pressure. Data are presented as means ± SD.

|                        | sham   | PEG IRI | VT IRI |
|------------------------|--------|---------|--------|
| MAP pre surgery [mmHg] | 106 ± 1| 107 ± 8 | 104 ± 8|
| MAP post surgery [mmHg]| 111 ± 6| 107 ± 8 | 104 ± 7|
| HR pre surgery [bpm]   | 383 ± 14| 390 ± 10| 392 ± 16|
| HR post surgery [bpm]  | 384 ± 17| 394 ± 15| 399 ± 12|

injury, whereas VT seemed to have no significant tubulo-protective effect, as shown by histology and urinary NGAL. Given that tubuli do not express Tie2, the latter finding is not surprising. However, an explanation for this discrepancy could be that COMP-Angpt1 is capable of mediating some protective effects independently of Tie2, possibly through its integrin-binding sequence QRHREGDS that VT lacks. Such an integrin-β1 dependent suppression of hypoxia-induced apoptosis by Angpt1 has already been established in cardiomyocytes, which lack Tie2 as well. Another possibility to account for the presumable distinctions between COMP-Angpt1 and VT is that higher severity of AKI (22 vs. 35 min clamp time in the current study) may preclude detection of subtle tubular protection conferred by VT. In line with this notion, Jung et al. detected only small histopathologic distinctions despite rather huge differences in serum creatinine values. In fact we too observed a trend towards less tubular damage in VT-treated mice. According to the steeper decline of creatinine and BUN at day 3 and 7, it is conceivable that this difference might become significant at later time points.

Second, in contrast to the aforementioned study by Jung et al. we clamped not only the renal artery, but the whole renal pedicle, which has been shown to reduce renal blood flow even further. They used standard invasive laser-Doppler flowmetry to assess renal blood flow in kidneys after IRI. CEUS yielded results similar to laser-Doppler flowmetry in murine kidneys treated with increasing doses of the vasoconstrictor Endothelin-1, supporting the validity of this non-invasive technique. We performed repeated CEUS measurements in a single mouse (pre- and post-surgery) in exactly the same position of animal, transducer and recording target region. Together these differences resulted in a stronger decline in medullary blood flow (−30 vs. −60%) shortly after reperfusion in our model. We further noted a marked decline in cortical blood flow (−0% vs −50%), owing to the higher severity of our model – which was tailored to result in a mortality of >50% in the control group. It is thus remarkable, that VT completely prevented IRI-induced changes in both, medullary and cortical perfusion parameters. This appeared to be mediated independent of changes in systemic hemodynamics, which argues against improved renal blood flow as mere consequence of better blood pressure. Herrler et al. recently uncovered that edema-related pressure elevation in the ischemic kidney can last for several days and accounts for impaired renal perfusion in AKI. They conclusively demonstrated that surgical pressure relief – via perforation of the renal capsule – prevented any reduction in renal blood flow after renal IRI. We therefore speculate that the improvement in renal perfusion seen with VT treatment is mainly due to reduced capillary leakage and subsequent formation of renal tissue edema after IRI. Mechanistically, Tie2 stimulation counteracts (hyper-)permeability through multi-level effects on intracellular signaling, cytoskeleton and junction-related molecules. In accordance with previous in vitro and in vivo studies we observed stabilization of the particularly important adherens-junction protein VE-cadherin, a well-defined downstream target of Tie2, conferred by VT.

We also analyzed survival after ischemic AKI. Probability of death from ischemic AKI was 86% lower in the VT-pretreated group [HR 0.14 (95% CI 0.05–0.78)]. Recent evidence suggests, that not only primary ischemic damage to the kidney, but also secondary, so-called remote organ injury, may affect outcome in AKI. Given that Tie2 mRNA and protein are most abundant in the pulmonary vasculature, it is conceivable that the lung is not only uniquely dependent on Tie2 signaling to maintain endothelial integrity, but that it may also constitute a target for deleterious Angpt2 released from the ischemic kidney during reperfusion. At least pulmonary vascular-specific Tie2 gene expression has already been shown to decline in response to renal IRI. It is thus conceivable, that part of the survival benefit in our model depends on a remote-organ “benefit” due to Tie2-stimulation by VT.

Our data suggest that selective protection of microvascular barrier function is sufficient to improve renal recovery and outcome in renal IRI. A Tie2 agonist drug may therefore have great potential as a vascular barrier-protective agent in human AKI. The potential clinical use of recombinant Angpt1, the endogenous ligand for Tie2, is limited by its short protein half-time, an inherent predisposition to protein aggregation, poor solubility of the protein, and the costly production. Due to these limitations the recombinant form of human Angpt1, and actually its improved variant COMP-Angpt1, represent considerable challenges for use in AKI patients. In contrast, VT was invented based on phage display experiments in which over 1 billion unique peptides were screened to identify those capable of binding Tie2 with high affinity but outside of the shared Angpt1/2 binding pocket. Vasculotide bears no sequence homology to Angpt1, a fact that negates the potential of an individual to develop
autoantibodies to endogenous Angpt1 or components of the chimeric protein, COMP-Angpt1. Preliminary work also demonstrates that VT neither activates, nor inhibits cellular receptors other than Tie2 (Supplementary Fig. 4, Supplementary Table 1, 2). In previous work, we have observed that a single dose of VT (200 ng per mouse i.p.) can sustain Tie2 phosphorylation in kidneys for >24 h\(^23\). Pretreatment with the same dose protected vascular barrier function in mouse models of acute skin radiation damage\(^4\) and septic multiple organ failure\(^23\), while a slightly higher dose (500 ng per mouse) was used in models of endotoxemic lung injury\(^2\) and lethal influenza\(^4\). Given that loss of Tie2 signalling (via loss of Tie2 and/or binding of released Angpt2) seems to be an early event in AKI, it seems to be exceedingly important that VT is given as soon as possible in order to maximize protective signaling through the remaining pool of Tie2 receptors. In line with this notion, a VT-rescue approach (i.e. VT first administered 2 h after reperfusion) still resulted in improved survival compared to PEG [HR 0.25 (95% CI

Figure 4. Vasculotide does not prevent acute tubular injury. Mice (n = 8 per group) were pre-treated with 200 ng of VT or PEG i.p. at −16 h, −1 h prior to renal IRI or sham surgery. (A) Representative H&E stained kidney sections, (B) mean tubular injury score and (C) distribution of tubular injury severity (PEG-IRI vs VT-IRI: Chi-square test P = 0.3) between the groups 24 h after IRI or sham surgery. (D) Levels of urinary NGAL 24 h after renal IRI. Data are means ± SEM. +++ P < 0.001 vs. sham.
but differences between groups were modest (Supplementary Fig. S5). However, because of the immediate and strong decline of renal blood flow (i.e. functional dropout of impaired capillaries) in IRI, any drug delivered through the bloodstream at 2 h after the insult may not have full access to its (pharmacological) target (as with pre-treatment). Therefore, post-treatment testing in this specific severe model of renal IRI represents a formidable treatment challenge. From a clinical perspective, endothelium-targeted (pre-)treatment with VT could represent a feasible approach to preventing AKI in the setting of radio contrast administration, renal transplantation and major surgery in high-risk patients (including coronary artery bypass graft). Further studies are needed to identify optimal dosing regimens, improve understanding of prophylactic vs. therapeutic (i.e. post injury) VT administration and to examine the long term impact of (specifically) improving the vascular component of AKI. As mentioned above, VT did not prevent tubular damage, which is a potential disadvantage over recombinant Angpt-1 and COMP-Angpt1. In future studies we will investigate VT in a less severe, unilateral ischemia model and compare VT and recombinant Angpt-1 in cultured tubular epithelial cells exposed to hypoxia and inflammatory stimuli.

In summary, we provide proof of principle in support of the efficacious use of PEGylated VT, a drug-like Tie2 receptor agonist, to activate Tie2 in vivo, counteract microvascular endothelial barrier dysfunction, improve renal recovery, and reduce mortality in ischemic AKI. Further studies are needed to pave the road for clinical application of this therapy concept.

**Figure 5. Vasculotide improves kidney function and reduces mortality after renal ischemia reperfusion injury.** Serum levels of (A) creatinine and (B) blood urea nitrogen (BUN) at day 1, 3 and 7 after renal IRI (n = 8 per group) or sham surgery (n = 4 per group). (C) survival after pretreatment with 200 ng VT or PEG – 16 h and – 1 h before renal IRI as well as + 24 h after surgery (n = 13 per group) or sham operation (n = 4). Data are expressed as means ± SEM. *P < 0.05; **P < 0.01 vs. IRI + PEG-Cys. ***P < 0.05; ****P < 0.01; *****P < 0.001 vs. sham. HR: Hazard Ratio, CI: Confidence Interval.
Methods
Vasculotide. Tournaire R. et al. previously described the discovery of a short synthetic peptide (HHHRHSF) that binds with high affinity to the extracellular portion of the Tie2 receptor but lacks the capacity to displace either Angpt1 or Angpt2. Using this peptide clustered as a tetramer by way of avidin/biotin, Van Slyke et al. demonstrated that Tie2 could be activated in a manner analogous to Angpt1. Subsequently, this proof of principle compound, termed Vasculotide, was reengineered into a more pharmaceutically amenable preparation which excludes the avidin/biotin complex in favor of a 4-armed, polyethylene glycol (average molecular weight 10 kDa) scaffold (Supplementary Fig. S1). A control version of VT was similarly prepared by conjugating 10 kDa PEG (VT backbone) quenched with cysteine on each arm. Synthesis, purification and validation of VT are described in detail elsewhere.

In vivo animal studies. All procedures were approved by the local committee for care and use of laboratory animals (Landesamt für Natur, Umwelt- und Verbraucherschutz Nordrhein-Westfalen – LANUV) and were performed according to international guidelines on animal experimentation. Eight- to 10-wk-old male C57BL6/J mice (20–25 g) were obtained from Charles River (The Charles River Laboratories; Sulzfeld, Germany). Mice were maintained on mouse chow and tap water ad libitum in a temperature-controlled chamber at 24 °C with a 12:12-h light-dark cycle. Renal IRI in mice was induced by bilateral clamping of the renal pedicles. In brief, mice were anesthetized using isoflurane (induction 3%, maintenance 1.5%, oxygen flow 3 L/minute) and a ventral midline abdominal incision was made. Both renal pedicles were clamped with microvascular clips for 35 min after that reperfusion was induced by removing the clips. The incision was then closed in layers using 4–0 surgical sutures. Sham animals underwent the same procedure except for clamping of the renal pedicles. Urine and tissues were harvested 24 hours after renal IRI.

Evans Blue Permeability Assay and wet/dry ratio. 24 h after surgery, mice were anesthetized and 1% w/v Evans blue (100 μl) was injected intravenously. Evans blue dye avidly binds to serum albumin and therefore can be used as a tracer for macromolecule flux across the microvasculature. 30 min after application of Evans blue, a blood sample was collected and kidneys were perfused with 10 ml PBS for 2 min via cardiac puncture. Then the kidneys were removed and homogenized in 1.5 ml formamide, followed by incubation at 70 °C for 24 h. The concentration of Evans Blue dye in appropriate dilutions of serum and cleared kidney homogenates was measured spectrophotometrically at 620 nm. The following formula was used to correct the optical densities for contamination with heme pigments: $E_{620} \text{(corrected)} = E_{620} \text{(raw)} - (E_{405} \text{(raw)} \times 0.014)$.

For wet/dry ratio, 24 h after IRI or sham operation, kidneys were harvested and weighed (wet weight). After incubation at 70 °C for 24 h, the organs were weighed again (dry weight). Wet/dry ratio is defined as wet weight/dry weight.

Examination of renal perfusion by CEUS. CEUS examination was performed immediately before and 1 h after surgery, thus every animal served as its own healthy baseline control. In brief, mice were fixed in supine position on a heat plate to prevent hypothermia. Heart rate, body temperature and respiratory rate were monitored throughout the whole examination. Both kidneys were visualized and the most accessible position was chosen to perform the study. A tripod was used to fix the ultrasound transducer (MS 250 Nonlinear Contrast Imaging transducer, VisualSonics Inc, Toronto, Canada) in position and to minimize intra-examination variability. Image depth, focus, gain and frame rate were optimized at the beginning of each study and held constant during the examinations. The contrast agent used in this study was MicroMarker™ (VisualSonics Inc, Toronto, Canada), a non-targeted contrast agent, which contains phospholipid-shelled microbubbles 2–3 μm in size, encapsulating a C3F15/N2 gas mixture. A bolus of 2 μl was injected via the tail vein using a syringe pump to ensure a constant infusion rate and pressure. Starting the contrast agent bolus, an image sequence of the bolus perfusion was acquired (18 MHz, 32 dB). After that a destruction-replenishment sequence was performed. Destruction of the microbubbles was performed by applying five ultrasound pulses (flash) of high acoustic power and then the refilling sequence was acquired. To avoid breathing and other movement artefacts, all sequences were acquired using motion correction (sequences during breathing were skipped).

Data analysis of CEUS measurements. The Vevo 2100 ultrasound imaging system (VisualSonics Inc, Toronto, Canada) was used to acquire all images. All sequences were exported and analyzed with the VevoCQ Contrast Quantification Software (VisualSonics Inc, Toronto, Canada), an application designed for quantifying perfusion in small animal models by CEUS imaging. It provides quantitative measurements by computing different perfusion parameters by means of a dedicated curve-fitting algorithm applied on contrast-uptake kinetics (time intensity curves).

After loading the sequences, 4 microvascular regions of interest (ROI) - 2 in the cortex and 2 in the medulla - were captured. Absence of large vessels in ROIs was verified by lack of bubbles after microbubble destruction (flash). The ROIs of the pre-surgery (baseline) examination were copied and pasted in the corresponding post-surgery sequences to ensure that exactly the same regions were examined in one animal pre- and post-surgery. The software then determined for each ROI the peak enhancement (PE), WiR and TTP from the bolus perfusion sequences and the relative blood volume (rBV) and mean transit time (mTT) from the destruction-replenishment sequences (Supplementary Fig. S2). PE is a measure of relative blood volume, while TTP is an absolute time measurement from the beginning of the bolus to the peak of contrast enhancement. WiR is defined as the maximum slope of the curve fit function which comprises both amplitude and time and is another measure of relative blood flow. The rBV is a measure of the blood volume from destruction to whole refilling and mTT is the time which is needed until half of rBV is reached (Table 1). The RBF was calculated as the rBV/mTT ratio. TTP and mTT are measured in seconds while all other parameters are given in arbitrary units.
In order to minimize inter-animal variability the mean value of the 2 ROIs from each cortex and medulla was calculated and expressed as percentage change of the post-surgery examination compared to the individual baseline study. Relative renal vascular resistance (rRVR) was calculated as follows: rRVR = MAP/rRBF (given in a.u.).

**Measurement of blood pressure and heart rate.** Immediately before the first ultrasound examination, average blood pressure and heart rate were measured via tail cuff in each animal (5 measurements) using the mouse blood pressure system MRBP-M01 (ITC Life science, Woodland Hills, CA, USA). The procedure was repeated after surgery immediately before the second ultrasound examination.

**Clinical Chemistry.** Serum creatinine and BUN were measured on an Advia 1800 automated analyzer system (Siemens Healthcare Diagnostics, Eschborn). Creatinine was measured based on the enzymatic reaction of Tanganelli et al. (1982). The method was validated by regular analyses of reference sera supplied by the national German INSTAND proficiency testing program and the international quality assurance program of the US Centers for Disease Control and Prevention. The enzymatic creatinine method is traceable to an HPLC candidate reference method, which uses reference materials from the National Institute of Standards and Technology. Urinary levels of NGAL were detected using a specific ELISA kit (BioPorto Diagnostics, Hellrup, Denmark) according to the manufacturer’s instructions.

**Immunofluorescence.** Immunofluorescence (IF) was performed on ice cold acetone-fixed cryosections (6 μm) using the following primary antibodies: polyclonal rabbit anti-mouse VE-cadherin Ab (Santa Cruz, Biotechnology, Dallas, Texas, USA), monoclonal rat anti-mouse PECAM-1 (CD31) Ab (BD Biosciences, Franklin Lakes, New Jersey, USA). Nonspecific binding sites were blocked with 10% normal donkey serum (Jackson Immunoresearch Laboratories, PA, USA) for 30 minutes. Thereafter sections were incubated with the primary antibody (1:200) for 1 h. Both incubations were performed in a humid chamber at room temperature. Slides were washed and the secondary antibodies (goat anti-rabbit Cy3 (Dianova, Hamburg, Germany) and donkey anti-rat alpha fluor 488 (Invitrogen, Carlsbad, CA, USA)) were applied overnight at 4°C (1:1000). For negative controls the staining procedure was performed as described without the primary antibodies. Data were acquired on a Zeiss Axioimager (20x/0.45 objective) equipped with a Hamamatsu ORCA–ER camera. Images were analyzed using Velocity software (Improvision, Forchheim, Germany).

**Western Blot analysis.** For the Western Blot, kidneys of the mice were stored at −80°C and were directly collected in 1 x Laemmlli (4% SDS, 5% 2-mercaptoethanol, 10% glycerol, 0.002% bromophenol blue, 0.0625 M Tris-HCl (pH 6.8) in a tissue grinder. After homogenization of the tissue the samples were heated for 5 min at 95°C. The samples were then pushed through a 20-gauge-needle and equal quantities of cell lysates were separated on SDS-PAGE gels. Proteins were transferred to a polyvinylidenfluoride (PVDF) membrane (Merck Millipore, Darmstadt, Germany) and incubated for 30 min at 37°C in blocking buffer (5% bovine serum albumin (BSA) dissolved in tris-buffered saline (TBS) containing 0.05% Tween-20 (TBS-T). β-Tubulin served as loading control. Primary antibodies (anti-Tie2 (C-20), Santa Cruz Biotechnology, Dallas, Texas, USA; anti-phospho-Tie2 (Y992), R&D Systems, Minneapolis, MN, USA) were incubated (1:1000 in 5% BSA dissolved in TBS-T) for 1 h at room temperature. Binding of primary antibodies was detected using horseradish peroxidase conjugated secondary antibodies (Dianova, Hamburg, Germany) with a chemiluminescence detection reagent (Roche, Rotkreuz, Switzerland) exposed to a medical X-ray film (Fujifilm, Tokio, Japan).

**Histology.** Paraffin-embedded kidney sections were stained with hematoxylin-eosin. Tubular injury was scored by estimating the percentage of tubules in the cortex or the outer medulla that showed epithelial necrosis or had luminal necrotic debris and tubular dilatation as follows: 0 = none; 1 = < 5%; 2 = 5–25%; 3 = 25–75%; and 4 = > 75%.

**Transmission Electron Microscopy.** Kidney tissue samples were fixed for 12 h at 4°C in 3% cacodylate-buffered glutaraldehyde (pH 7.35) and then transferred into 5% sucrose for electron microscopy. Post fixation was performed with 1% osmium tetroxide and 50 mm potassium ferricyanide. Specimens were then washed with distilled water, dehydrated in graded alcohols, and embedded in araldite resin. The ultrathin sections (50 nm) were cut and placed on copper grids. The sections were stained with 5% uranyl acetate and 0.2% lead citrate and 4°C (1:1000). For negative controls the staining procedure was performed as described without the primary antibodies. Data were acquired on a Zeiss AxioImager (20x/0.45 objective) equipped with a Hamamatsu ORCA–ER camera. Images were analyzed using Velocity software (Improvision, Forchheim, Germany).

**Statistical analysis.** Data are presented as means ± standard error of mean (SEM). Differences between groups were compared by one-way analysis of variance (ANOVA) with Tukey as post hoc test for multiple comparisons. Kaplan-Meier plots were used to illustrate survival between treatment groups and statistical assessment was performed by the log-rank test. Animals still alive at 14 days after IRI were censored at day 14. All tests were two-sided and significance was accepted at p < 0.05. GraphPad Prism Version 5.02 (GraphPad Prism Software Inc, San Diego, California, USA) was used for data analysis and figure preparation.

**References**

1. Uchino, S. et al. Acute renal failure in critically ill patients: a multinational, multicenter study. JAMA 294, 813–818 (2005).
2. Lopes, J. A. et al. Acute kidney injury in intensive care unit patients: a comparison between the RIFLE and the Acute Kidney Injury Network classifications. Crit Care 12, R110 (2008).
3. Sutton, T. A. Alteration of microvascular permeability in acute kidney injury. Microvasc Res 77, 4–7 (2009).
4. Bonventre, J. V. & Yang, L. Cellular pathophysiology of ischemic acute kidney injury. J Clin Invest 121, 4201–4221 (2011).
5. Molitoris, B. A. Therapeutic translation in acute kidney injury: the epithelial/endothelial axis. J Clin Invest 124, 2355–2363 (2014).
6. Sutton, T. A. et al. Injury of the renal microvascular endothelium alters barrier function after ischemia. *Am J Physiol Renal Physiol* **285**, F191–198 (2003).

7. Herrler, T. et al. The intrinsic renal compartment syndrome: new perspectives in kidney transplantation. *Transplantation* **89**, 40–46 (2010).

8. Sato, T. N. et al. Distinct roles of the receptor tyrosine kinases Tie-1 and Tie-2 in blood vessel formation. *Nature* **376**, 70–74 (1995).

9. Davis, S. et al. Isolation of angiopoietin-1, a ligand for the TIE2 receptor, by secretion-trap expression cloning. *Cell* **87**, 1161–1169 (1996).

10. Grams, M. E. et al. Requisite role of angiopoietin-1, a ligand for the TIE2 receptor, during embryonic angiogenesis. *Cell* **87**, 1171–1180 (1996).

11. Thurston, G. et al. Angiopoietin-1 protects the adult vasculature against plasma leakage. *Nat Med* **6**, 460–463 (2000).

12. Gamble, J. R. et al. Angiopoietin-1 is an antipermeability and anti-inflammatory agent in vitro and targets cell junctions. *Circ Res* **87**, 603–607 (2000).

13. Fiedler, U. et al. Angiopoietin-1 and angiopoietin-2 share the same binding domains in the Tie-2 receptor involving the first Ig-like loop and the epidermal growth factor-like repeats. *J Biol Chem* **278**, 1721–1727 (2003).

14. Fiedler, U. et al. The Tie-2 ligand angiopoietin-1 is stored in and rapidly released upon stimulation from endothelial cell Weibel-Palade bodies. *Blood* **103**, 4150–4156 (2004).

15. Parikh, S. M. et al. Excess circulating angiopoietin-2 may contribute to pulmonary vascular leak in sepsis in humans. *PLoS Med* **3**, e46 (2006).

16. Roviezzo, F. et al. Angiopoietin-2 causes inflammation in vivo by promoting vascular leakage. *J Pharmacol Exp Ther* **314**, 738–744 (2005).

17. Benest, A. V. et al. Angiopoietin-2 is critical for cytokine-induced vascular leakage. *PLoS One* **8**, e70459 (2013).

18. Kharroun, M. et al. Renal ischemia-reperfusion induces a dysbalance of angiopoietins, accompanied by proliferation of pericytes and fibrosis. *Am J Physiol Renal Physiol* **305**, F901–910 (2013).

19. de Vries, D. K. et al. Renal ischemia-reperfusion release of angiopoietin-2 from human grafts of living and deceased donors. *Transplantation* **96**, 282–289 (2013).

20. Jung, Y. J. et al. Peritubular capillary preservation with COMP-angiopoietin-1 decreases ischemia-reperfusion-induced acute kidney injury. *Am J Physiol Renal Physiol* **297**, F952–960 (2009).

21. Syrjala, S. O. et al. Donor Heart Treatment With COMP-Ang1 Limits Ischemia-Reperfusion Injury and Rejection of Cardiac Allografts. *Am J Transplant* **15**, 2075–2084 (2015).

22. Syrjala, S. O. et al. Angiopoietin-2 inhibition prevents transplant ischemia-reperfusion injury and chronic rejection in rat cardiac allografts. *Am J Transplant* **14**, 1096–1108 (2014).

23. Kumpers, P. et al. The synthetic tie2 agonist peptide vasculotide protects against vascular leakage and reduces mortality in murine abdominal sepsis. *Crit Care* **15**, R261 (2011).

24. Van Slyke, P. et al. Acceleration of diabetic wound healing by an angiopoietin peptide mimetic. *Tissue Eng Part A* **15**, 1269–1280 (2009).

25. Deshpande, N., Needle, A. & Willmann, J. K. Molecular ultrasound imaging: current status and future directions. *Clin Radiol* **65**, 567–581 (2010).

26. Needle, A. et al. Nonlinear contrast imaging with an array-based micro-ultrasound system. *Ultrasound Med Biol* **36**, 2097–2106 (2010).

27. Haase, M. et al. Accuracy of neutrophil gelatinase-associated lipocalin (NGAL) in diagnosis and prognosis in acute kidney injury: a systematic review and meta-analysis. *Am J Kidney Dis* **54**, 1021–1024 (2009).

28. Prowle, J. R. et al. Combination of biomarkers for diagnosis of acute kidney injury after cardiopulmonary bypass. *Ren Fail* **37**, 408–416 (2015).

29. Bellomo, R., Kellum, J. A. & Ronco, C. Acute kidney injury. *Lancet* **380**, 756–766 (2012).

30. David, S. et al. Acute administration of recombinant Angiopoietin-1 ameliorates multiple-organ dysfunction syndrome and improves survival in murine sepsis. *Cytokine* **55**, 251–259 (2011).

31. van Meurs, M. et al. Shock-induced stress induces loss of microvascular endothelial Tie2 in the kidney which is not associated with reduced glomerular barrier function. *Am J Physiol Renal Physiol* **297**, F272–281 (2009).

32. David, S. et al. Effects of a synthetic PEGylated Tie-2 agonist peptide on endotoxemic lung injury and mortality. *Am J Physiol Lung Cell Mol Physiol* **300**, L851–862 (2011).

33. Yuan, H. T., Suri, C., Yancopoulous, G. D. & Woof, A. S. Expression of angiopoietin-1, angiopoietin-2 and the Tie-2 receptor tyrosine kinase during mouse kidney maturation. *J Am Soc Nephrol* **10**, 1722–1736 (1999).

34. Lee, S. W. et al. Angiopoietin-1 protects heart against ischemia/reperfusion injury through VE-cadherin dephosphorylation and myocardial integrin-beta/ERK/caspase-9 phosphatase phosphorylation. *Mol Med* **17**, 1095–1106 (2011).

35. Li, X. et al. Acute renal venous obstruction is more detrimental to the kidney than arterial occlusion: implication for murine models of acute kidney injury. *Am J Physiol Renal Physiol* **302**, F519–525 (2012).

36. Sullivan, J. C. et al. Novel use of ultrasound to examine regional blood flow in the mouse kidney. *Am J Physiol Renal Physiol* **297**, F228–235 (2009).

37. Augustin, H. G., Koh, G. Y., Thurston, G. & Altalaloo, K. Control of vascular morphogenesis and homeostasis through the angiopoietin-Tie system. *Nat Rev Mol Cell Biol* **10**, 165–177 (2009).

38. Milam, K. E. & Parikh, S. M. The angiopoietin-Tie signaling axis in the vascular leakage of systemic inflammation. *Tissue Barriers* **3**, e95708 (2015).

39. van Meurs, M. et al. Bench-to-bedside review: Angiopoietin signalling in critical illness - a future target? *Crit Care* **13**, 207 (2009).

40. Grams, M. E. & Rabb, H. The distant organ effects of acute kidney injury. *Kidney Int* **81**, 942–948 (2012).

41. Feltus, C. E., Hassoun, H. T., Lie, M. L., Cheadle, C. & Rabb, H. Pulmonary endothelial cell activation during experimental acute kidney injury. *Shock* **36**, 170–176 (2011).

42. Cho, C. H. et al. Designed angiopoietin-1 variant, COMP-Ang1, protects against radiation-induced endothelial cell apoptosis. *Proc Natl Acad Sci USA* **101**, 5553–5558 (2004).

43. Tournaire, R. et al. A short synthetic peptide inhibits signal transduction, migration and angiogenesis mediated by Tie2 receptor. *EMBO Rep* **5**, 262–267 (2004).

44. Korpela, E. et al. Vasculotide, an Angiopoietin-1 mimetic, reduces acute skin ionizing radiation damage in a preclinical mouse model. *BMC Cancer* **14**, 614 (2014).

45. Sugiyama, M. G. et al. The Tie2-agonist Vasculotide rescues mice from influenza virus infection. *Sci Rep* **5**, 11030 (2015).

**Acknowledgements**

We thank Richard Holtmeier, Dirk Wennmann and Rita Schröter (Münster, Germany) for extraordinary technical assistance. This work was supported by an institutional research grant to ER from Medical Faculty of the Westfälische Wilhelms-University Münster. DJD is a Canada Research Chair in Angiogenesis and Lymphangiogenesis Signaling. This study was funded in part by the Heart and Stroke Foundation of Canada (GRANT No. NA7027, to PVS & DJD). We acknowledge support by Innovative Medizinische Forschung (IMF, Grant No. RU221207) of the University of Münster and by the Dr. Werner Jackstädt-Stiftung. We acknowledge support by Deutsche Forschungsgemeinschaft and Open Access Publication Fund of University of Münster.
Author Contributions
E.R. performed the experiments, analyzed the data, prepared the figures and wrote the manuscript. J.S. performed the ultrasound studies and reviewed the manuscript. P.V.S. and D.D. generated the PEGylated Tie2 mimetic Vasculotide and contributed to the manuscript. T.S., K.B. and T.G. performed electron-, immune- and light microscopic studies and reviewed the manuscript. S.R. and H.P. contributed to the design of the study and reviewed the manuscript. P.K. had the initial idea, supervised the project, analyzed the data, prepared figures and contributed to the manuscript.

Additional Information
Supplementary information accompanies this paper at http://www.nature.com/srep

Competing financial interests: DD and PVS are listed as inventors on three patents related to VT filed by the Sunnybrook Research Institute.

How to cite this article: Rübig, E. et al. The Synthetic Tie2 Agonist Peptide Vasculotide Protects Renal Vascular Barrier Function In Experimental Acute Kidney Injury. Sci. Rep. 6, 22111; doi: 10.1038/srep22111 (2016).

This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/