Oxidant sensor in the cGMP-binding pocket of PKGIα regulates nitroxyl-mediated kinase activity

Sonia Donzelli1,2,10, Mara Goetz2,1, Kjestine Schmidt3, Markus Wolters4, Konstantina Stathopoulou1,2, Simon Diering1,2, Oleksandra Prysyazhna5, Volkan Polat1,2, Jenna Scotcher5, Christian Dees4, Harihara Subramaniam7,2, Elke Butt6, Alisa Kamynina5, Sophie Schobesberger1,2, S. Bruce King8, Viacheslav O. Nikolaev7,2, Cor de Wit1, Lars I. Leichert9, Robert Feil4, Philip Eaton5 & Friederike Cuello1,2

Despite the mechanisms for endogenous nitroxyl (HNO) production and action being incompletely understood, pharmacological donors show broad therapeutic promise and are in clinical trials. Mass spectrometry and site-directed mutagenesis showed that chemically distinct HNO donors 1-nitrosocyclohexyl acetate or Angeli’s salt induced disulfides within cGMP-dependent protein kinase I-alpha (PKGIα), an interdisulfide between Cys42 of the two identical subunits of the kinase and a previously unobserved intradisulfide between Cys117 and Cys195 in the high affinity cGMP-binding site. Kinase activity was monitored in cells transfected with wildtype (WT), Cys42Ser or Cys117/195Ser PKGIα that cannot form the inter- or intradisulfide, respectively. HNO enhanced WT kinase activity, an effect significantly attenuated in inter- or intradisulfide-deficient PKGIα. To investigate whether the intradisulfide modulates cGMP binding, real-time imaging was performed in vascular smooth muscle cells expressing a FRET-biosensor comprising the cGMP-binding sites of PKGIα. HNO induced FRET changes similar to those elicited by an increase of cGMP, suggesting that intradisulfide formation is associated with activation of PKGIα. Intradisulfide formation in PKGIα correlated with enhanced HNO-mediated vasorelaxation in mesenteric arteries in vitro and arteriolar dilation in vivo in mice. HNO induces intradisulfide formation in PKGIα, inducing the same effect as cGMP binding, namely kinase activation and thus vasorelaxation.

Nitroxyl (HNO) is the one-electron-reduced and protonated sibling of nitric oxide. The extent to which HNO is produced endogenously and the mechanisms of its generation are still under intense investigation1,2. Despite this, pharmacological donors that release HNO have emerged as promising therapeutic reagents, with clinical trials ongoing3–5. The beneficial actions attributed to experimental therapeutic HNO release are multifaceted, such as vasorelaxation of human conduit and resistance arteries and lowering of blood pressure6,7, relief from chronic neuropathic pain8, anti-tumour effects9 and positive cardiac inotropy and lusitropy10,11. Unlike nitric oxide or
some routine clinically-utilised vasodilatory nitrates, HNO is not susceptible to vascular tolerance, is resistant to scavenging by reactive oxygen species, and maintains its effects also under pathophysiological conditions. Several mechanisms for HNO-mediated vasorelaxation have been suggested, including activation of soluble guanylyl cyclase to promote classical cGMP-dependent signaling, opening of voltage and calcium-dependent potassium channels and release of calcitonin gene-related peptide. However, the sGC inhibitor 1H-(1,2,4) oxadiazolo-(4,3-a)quinoxalin-1-one (ODQ) only partially attenuated HNO-mediated vasorelaxation in aortic rings, and in vivo administration of HNO reduced systemic blood pressure without increasing plasma cGMP levels. Therefore, the exact mechanisms of how HNO induces vasorelaxation and what targets in the vasculature mediate this are incompletely understood and require further investigations.

cGMP-dependent protein kinase (PKG) is a homodimeric serine/threonine kinase involved in numerous processes, including regulation of smooth muscle tone and vasorelaxation. The isoform I-alpha (PKG1α) is expressed in the vascular system, it consists of an N-terminal leucine zipper, an autoinhibitory domain, a low and a high affinity cGMP-binding domain and a C-terminal catalytic domain. Classical PKG1α activation occurs upon cGMP binding to regulatory sites, with the release of autoinhibition as a consequence. In addition to α activation occurs and a high affinity cGMP-binding domain and a C-terminal catalytic domain. Classical PKG1α induce substantive catalytic activation compared to cGMP-dependent activation of the kinase. Given that Cys42 of this thiol may be more important for modulating such interactions, as suggested previously, consistent with this regulatory role, Cys42Ser PKG1α knock-in (KI) mice, which cannot form the interprotein disulfide, are hypertensive, illustrating a physiological role for oxidants in blood pressure regulation.

In addition to Cys42, PKG1α possesses other cysteines that might play a role in regulating kinase activity. Among them, Cys117 and Cys195, localised in the high affinity cGMP-binding site have been shown to form an intradisulfide in the crystal structure of PKG1α. This intradisulfide in the second messenger-binding site represents a more rational explanation for modulating kinase activity in response to oxidants than the interdisulfide in the leucine zipper domain. However, whether oxidation of these cysteines plays a role in regulation of PKG1α activity is, to date, unknown.

Materials and Methods

Anti-PKG antibody was from Enzo Life Sciences (JADI-KAP-PK005-D, Lörrach, Germany). NCA was synthesised by S. Bruce King or by Axon Medchem (Groningen, The Netherlands). AS (Na,N₃O₃) was obtained from Cayman Europe (Tallinn, Estonia). Acetylcholine (ACh) was from Sigma-Aldrich (Taufkirchen, Germany) and NOxICAT kit from AB Sciex (Concord, ON, Canada). Ni-NTA agarose beads were from Qiagen (Hilden, Germany). PKG1α from bovine lung (obtained from Süddeutsches GmbH, Würzburg, Germany) was purified as previously described in accordance to the European TierSchlV No.1099/2009. γ³²P-ATP was purchased from GE Healthcare (Freiburg, Germany) and [H]cGMP from American Radiolabeled Chemicals (USA). Glass microfiber filters were purchased from Whatman (Maidstone, UK). Complete protease inhibitor cocktail was from Roche (Berlin, Germany). Human embryonic kidney cells (HEK-293) were purchased from ATCC (293-HEK-293(ATCC©CRL-1573TM); Wesel, Germany).

All methods were carried out in accordance with the relevant guidelines and regulations. Experiments were approved by the relevant institutional or licensing committees as indicated in the relevant section.

Transfection.

HEK-293 cells were used for transfection experiments. Cells were routinely screened for mycoplasm contaminations. Cells were transfected with empty pcDNA3 as a control or pcDNA3 containing the cDNA for human WT, Cys42Ser, Cys117Ser, Cys195Ser, Cys117/195Ser or Cys42/117/195Ser PKG1α (2 μg DNA/well) using Turbofect (Turbofect, Thermo Scientific, Venlo, Limburg, The Netherlands). After 24 hours, cells were treated with NCA (100 µmol/L, 30 min), AS (500 µmol/L, 15 min) or vehicle (1% DMSO or 100 µmol/L decomposed NCA for NCA; 10 mmol/L NaOH, 50 µmol/L nitrite (NO₂⁻), 500 µmol/L decomposed AS for AS). Western immunoblot analysis was performed to assess oxidative PKG1α modifications or in vitro kinase assays to assess PKG1α activity.

Western immunoblot analysis.

Snap frozen isolated mesenteric vessels (after 100 µmol/L NCA), cremaster muscles (after 50 µmol/L NCA or AS) or HEK-293 cell homogenates (after 100 µmol/L NCA or 500 µmol/L AS) in non-reducing (containing in mmol/L: Tris-HCl 187.5 pH 6.8, SDS 6% (w/v), glycerol 30% (v/v), bromophenol blue 0.03% (w/v), maleimide 100) or reducing (9% (w/v) 2-mercaptoethanol) Laemmli sample buffer were subjected to western immunoblot analysis as described previously.

NOxICAT LC-MS/MS analysis.

Modified cysteines in PKG1α by HNO were identified by a thiol-trapping technique using isotope-coded affinity-tag chemistry (NOxICAT) as described previously. Briefly, non-tagged recombinant human PKG1α was pre-treated with DTT (10 mmol/L, 30 min) in an anaerobic chamber, the DTT was removed by buffer exchange via passing the sample through a PD MiniTrapTM G-25 column (28-9180-07) from GE Healthcare. After buffer exchange, samples were exposed to NCA (25 µmol/L, 15 min), AS (25 µmol/L, 15 min) or vehicle for 15 min at RT (Supplementary Figure 1A,B). The reaction was terminated by addition of ice-cold acetone and proteins precipitated overnight at −20°C. The precipitate was resuspended in a mixture of 80 µL denaturing alkylator buffer (DAB; containing in mmol/L: urea 6000, Tris-HCl 200 pH 8, EDTA 10, SDS 0.5% (w/v)) containing 20 µL acetonitrile and 1 vial ICAT reagent “light” from the cleavable ICAT methods development kit (AB Sciex). Reduced cysteines were labeled under denaturing conditions at 37°C in a thermomixer at 1300 rpm for 2 hours. After a second acetone precipitation, oxidised cysteines were reduced for 30 min in 80 µL DAB containing 2 mmol/L TCEP. Subsequently, 1 vial cleavable ICAT reagent “heavy”, resuspended in
20 µL acetonitrile, was added and previously oxidised cysteines were labeled at 37 °C at 1300 rpm for 2 hours. The reaction was stopped by acetone precipitation. The pellet was resuspended in 80 µL denaturing buffer from the cleavable ICAT reagent kit. 20 µL acetonitrile was added to the protein digest with 100 µL aqueous trypsin resuspension from the ICAT reagent kit. ICAT labeled peptides were purified by cation exchanger chromatography and affinity chromatography. Light- and heavy-ICAT-labeled peptides were analysed and quantified by reverse phase nano-liquid chromatography and detected by MS/MS with Fourier transform mass spectrometry in an LTQ Orbitrap instrument (Thermo Fisher Scientific, Waltham, MA). Peptides were identified by SwissProt database and quantified by MaxQuant.

**FRET-experiments in primary vascular smooth muscle cells.** The isolation of primary mouse vascular smooth muscle cells (VSMCs) from cGi500 transgenic mice was approved by the Regierungspräsidium Tübingen in compliance with the humane care and use of laboratory animals. For in-cell FRET measurements, primary VSMCs from R26-CAG-cGi500(L1) mice were used. These cells express the FRET-based biosensor cGi500 (cGMP indicator with an EC50 of 500 mmol/L) [38]. VSMCs were continuously superfused with intracellular-like medium (ICM; containing in mmol/L: HEPES 6.7 pH 7.3, KCl 125, NaCl 19, EGTA 1, CaCl2 0.3) at room temperature. Cells were permeabilised by superfusion with β-escin (100 µmol/L in ICM for 80 sec) and subsequently exposed to ICM supplemented with increasing concentrations of cGMP (0.1, 1, 10 µmol/L; 2 min each). The individual fluorescence of CFP and YFP was recorded simultaneously using a DualView DV2 beam-splitter (Photometrics) and saved as individual TIFF images. After each cGMP application, baseline recovery of the fluorescence signals was achieved before the next drug application. After the initial series of cGMP applications, VSMCs were exposed to 0.02% DMSO (in ICM) or 100 µmol/L NCA (in ICM containing 0.02% DMSO) for 30 min followed by superfusion with ICM supplemented with ifosfamide concentrations of cGMP (0.1, 1, 10 µmol/L). Acquired images were analysed using Fiji [20] and Microsoft Excel to correct for background fluorescence and calculate the CFP/YFP ratio (R) traces, which reflect FRET changes. FRET responses were measured as amplitudes over baseline. The individual FRET changes (ΔR) were then normalised to the FRET change induced by the first application of 10 µmol/L cGMP (ΔR0) and are displayed as ΔR/ΔR0. Further details on measurement and evaluation of FRET signals in VSMCs are described elsewhere [37, 40].

**In vitro cGMP-binding assay.** cGMP-binding affinity to purified bovine PKGIα was assessed by measuring (3H)cGMP binding in the presence of increasing concentrations of unlabeled cGMP. The experiment was performed under ambient air. PKGIα (600 fmol/reaction) was diluted in cGMP-binding buffer (containing in mmol/L: HEPES 6.7 pH 7.4, Mg(CH3COO)2 5, NaH2PO4 2H2O, 3, KCl 130, IBMX 0.1, EGTA 0.1, EDTA 0.1). PKGIα was either reduced with DTT (100 mmol/L; 10 min) or oxidised with NCA (100 µmol/L; 30 min) followed by incubation with (3H)cGMP in the presence of increasing concentrations of unlabeled cGMP (0, 10, 30, 100, 300, 1000 nmol/L) for 1 h. Reactions were terminated by adding cold saturated ammonium sulfate buffer (pH 8.3) and vacuum-filtered through glass microfiber filters (Q 25 mm). The filter papers were washed three times with 2 mL ice-cold ammonium sulfate buffer and air-dried. Papers were put into scintillation vials and suspended in 2 mL 2% SDS (w/v), shaken vigorously and incubated for 1 h at room temperature. Afterwards, 10 µl aqueous scintillation liquid (Rotiszint® Eco plus from Carl Roth) was added and incubated for 10 min prior to scintillation counting.

**In vitro PKG activity assay.** Protein kinase activity was investigated using two different assay systems. On the one hand, activity of purified bovine PKGIα was assessed by Glasside (Calbiochem) phosphorylation in the presence of radiolabeled γ32P-ATP (GE Healthcare). The experiment was performed under ambient air. PKGIα (600 fmol/reaction) was initially reduced with DTT (100 mmol/L; 10 min) or oxidised with NCA (100 µmol/L; 15 min) and in vitro kinase assays performed in assay buffer (containing in mmol/L: Tris 30 pH 7.4, ATP 0.1, MgCl2 15, Glascide 0.1), in the absence of presence of 300 nmol/L cGMP. Reactions were terminated with phosphoric acid (25 mmol/L) before spotting onto P81 phosphocellulose squares (Whatman). After air-drying, the filter papers were washed with phosphoric acid (75 mmol/L; 4 × 2 min), ethanol (1 × 15 sec), air-dried and transferred into scintillation vials. Vials were subjected to Cerenkov counting. Six independent experiments were carried out. PKGIα activity was expressed as pmol phosphate incorporated into PKGIα substrate per minute. On the other hand, activity of human WT, Cys42Ser, Cys117/195Ser, or Cys42/117/195Ser PKGIα was assessed by phosphorylation of recombinantly expressed His6-tagged cardiac myosin-binding protein C (amino acid residues 153–450) [24], which was previously described as a PKGIα substrate [25], in the presence of radiolabeled γ32P-ATP (GE Healthcare). HEK-293 cells were transfected as described before and exposed to AS (500 µmol/L; 15 min), NCA (100 µmol/L; 30 min) or vehicle (NaOH for AS; DMSO for NCA). Cell homogenates were prepared in lysis buffer (containing in mmol/L: Tris 20 pH 7.4, NaCl 150, EDTA 1, EGTA 1, NaF 2, Complete protease inhibitor cocktail) and were diluted 1:1 in assay buffer (containing in mmol/L: Tris 30 pH 7.4, MgCl2 15) with ATP (30 µmol/L spiked with γ32P-ATP (GE Healthcare)). The in vitro kinase reaction was started by addition of His6-tagged C1–M–C2 (500 pmol/reaction) prebound to Ni-NTA agarose beads equilibrated in assay buffer. The reaction was carried out for 30 min at 30 °C and 1300 rpm. Samples were centrifuged at 4 °C for 1 min at 1000 xg, supernatant discarded and the agarose beads resuspended in 75 µL 3x reducing Laemmli sample buffer, heated for 5 min at 75 °C and proteins resolved by 10% SDS-PAGE. Gels were stained with colloidal coomassie to assure equal substrate content between samples, destained in 20% (v/v) methanol, incubated briefly in 20% (v/v) glycerol in water and vacuum-dried. Experiments were analysed by autoradiography and densitometry was performed with GelQuant. NET software provided by biochemlabsolutions.com.

**Myography.** All animal protocols were approved by the local King’s College London UK Ethical Review Process Committee and by the UK Government Home Office (Animals Scientific Procedures Group) and the
study was conducted in accordance with the Home Office Guidance on the Operation of the Animals. Third-order mesenteric arteries from 12-week-old male C57BL/6 mice were mounted for isometric tension recordings in a pressure myograph (Danish Myo Technology), stretched to the optimal pre-tension conditions (using DMT Normalisation Module), bathed in Krebs solution maintained at 37°C and gassed with 95% CO₂: 5% O₂ (containing in mmol/L: NaCl 119, KCl 4.7, KH₂PO₄ 1.2, NaHCO₃ 25, MgSO₄·7H₂O 1.2, glucose 11.1, CaCl₂·2H₂O 1.6). Vasorelaxation of mesenteric arteries was assessed after U46619-pre-constriction (100 nmol/L) in response to HNO donors (NCA: 0–300 μmol/L or AS: 0–100 μmol/L). The concentration-response curves to NCA were constructed in a cumulative fashion. Tension experiments were carried out using one or two vessels per intervention derived from at least 4 different WT animals.

**Intravital microscopy.** 12-week-old male C57BL/6NCrl mice (Charles River) were used for intravital microscopy of the microcirculation in the cremaster muscle in vivo as described previously. Experiments were in accordance with the German animal protection law and approved by the Ministerium für Energiewende, Landwirtschaft, Umwelt und ländliche Räume of Schleswig-Holstein. Mice were anaesthetised with fentanyl (0.05 mg/kg), midazolame (5 mg/kg), and dexmedetomidine (0.5 mg/kg) by intraperitoneal injection. After insertion of a catheter into the right jugular vein, anesthetic drugs were continuously infused. A tube was inserted via tracheotomy and the animals were ventilated with a stroke volume of 0.225 mL at 160 strokes per minute using a respirator (MiniVent, Harvard Apparatus). The right cremaster muscle was exposed and spread over a coverslip to allow intravital microscopy. It was continuously superfused with a 35°C warmed saline solution (containing in mmol/L: NaCl 118.4, KCl 3.8, CaCl₂ 2.5, MgSO₄ 1.2, NaHCO₃ 20, KH₂PO₄ 1.2) with a pH of 7.4 achieved by gassing with 5% CO₂ in N₂. In each mouse 7 to 18 arterioles were studied using an optical microscope (Eclipse E600, Nikon) and a 20-fold objective. The microscope was equipped with a digital camera (Zeiss Axioscam 105) connected to a PC to allow image storage (Zen2 lite, Zeiss) and later offline analysis. Arteriolar inner diameters were measured by a code written in the laboratory using the commercially available software (LabVIEW, National Instruments). Arteriolar diameters were assessed before and during application of NCA (10 μmol/L; group 1) or AS (3 to 50 μmol/L; group 2). The respective solvent (0.1% DMSO for NCA or 0.1 mmol/L NaOH for AS) was also evaluated. To assess vascular reactivity, the effect of acetylcholine (ACh, 10⁻⁶ mol/L; group 1) or AS (3 to 50 μmol/L; group 2) was also assessed. All substances were added to the superfusion solution using a roller pump and the final concentration on the preparation is indicated. After application of each substance, vessels were allowed to recover for 3 to 5 min and to return to their resting diameter. At the end of the experiment, mice were sacrificed by intravenous injection of pentobarbital (24 mg) and both cremaster muscles of each animal were harvested for western blot analysis. While the right cremaster muscle was exposed to the respective substances during the experiment, the left cremaster muscle was prepared immediately after pentobarbital injection and served as untreated control for western immunoblot analysis.

**Statistical analysis.** Statistical comparisons were performed by one-way ANOVA (*in vitro* kinase assays with HEK-293 lysates) or two-way ANOVA (*in vitro* kinase assays with bovine PKGIα; FRET analysis) followed by Bonferroni's multiple comparisons test; Student's t-test comparing samples with the same cGMP concentration (cGMP-binding assays). Arteriolar diameter changes are normalised to the respective maximal possible response:

\[
\% \text{ of maximal response} = \left( \frac{D_{\text{D_subst}} - D_{\text{D_con}}} {D_{\text{D_max}} - D_{\text{D_con}}} \right) \times 100
\]

where \(D_{\text{D_subst}}\) is the diameter in the presence of the substance, \(D_{\text{D_con}}\) the control diameter before application and \(D_{\text{D_max}}\) the respective maximal diameter observed for each vessel during the experiment. Data within groups were compared using paired t-test and corrected according to Bonferroni for multiple comparisons. Quantitative data are given as mean ± S.E.M and a value of \(P < 0.05\) was considered significant.

**Data availability.** The authors declare that all data supporting the findings of this study are available within the paper and its supplementary file.

**Results** To investigate whether PKGIα is a direct HNO target and is modified by cysteine oxidation, recombinant PKGIα was exposed to two chemically distinct HNO donors, namely 1-nitrosocyclohexyl acetate (NCA) or Angeli’s salt (AS). The oxidation state of cysteines in PKGIα was then analysed by NOxICAT methodology, involving differential isotopic labeling of reduced or oxidised thiol groups, and subsequent mass spectrometry. Without prior reduction with DTT, recombinant PKGIα is fully oxidised by air. A DTT concentration-response curve was performed initially with recombinant PKGIα (Supplementary Figure 1A,B) to determine the optimal DTT concentration. In the NOxICAT experiments, we observed increased oxidation of Cys42, Cys117 and Cys195 in PKGIα after exposure to either donor (Fig. 1). Despite DTT-pretreatment, Cys42 was still found partially oxidized in control samples (Fig. 1B, left panel). However, upon treatment with NCA or AS, Cys42 was detected exclusively in the oxidised form. Cys117 as well as Cys195 were predominantly found in the reduced state under basal conditions, but were completely oxidised after NCA treatment (Fig. 1B, middle and right panels); AS induced oxidation of these cysteines as well, albeit to a lesser extent than NCA (Fig. 1B, middle and right panels). The fact that Cys42 was almost completely oxidised by both donors suggests that HNO induces Cys42 intradisulfide formation between PKGIα monomers. The similar extent of Cys117 and Cys195 oxidation following AS or NCA treatment was consistent with the possibility of the formation of an intradisulfide bond between these two cysteines. Indeed, an intradisulfide bond between them within the cGMP-binding pocket of PKGIα has been reported, but was considered a constitutive modification. To investigate the potential impact of intradisulfide formation after HNO-exposure on kinase activity, wildtype (WT), Cys42Ser, Cys117/195Ser or Cys42/117/195Ser PKGIα that cannot form the intradisulfide, intradisulfide or either, respectively, were expressed in HEK-293 cells. Kinase
Figure 1. Identification of HNO-modified cysteines in PKGIα by NOxICAT. (A) Exposure of recombinant human PKGIα to HNO donors induced oxidation of Cys42, Cys117 and Cys195 in PKGIα. The table summarises the mass charge (m/z) in molecular ions (mi) of the detected peptides containing the cysteines in the reduced (blue) and oxidised (pink) state. (B) Fourier Transform Mass Spectrometry spectra are expressed as the relative abundance of the detected peptides in m/z containing Cys42 (left), Cys117 (middle) and Cys195 (right), vehicle-treated (top), after exposure to 1-nitrosocyclohexyl acetate (NCA; 25 mmol/L, 15 min) (middle) or Angeli’s salt (AS; 25 μmol/L, 15 min) (bottom). (C) Quantification of the detected oxidised (pink) versus reduced (blue) peptide fraction is shown as percent of the total detected peptides.
Figure 2. Effect of HNO-mediated PKG↓α oxidation on kinase activity. HEK-293 cells were transfected to express PKG↓α WT or various mutants Cys42Ser, Cys117/195Ser, Cys42/117/195Ser, exposed to (A) NCA (100 µmol/L, 30 min) or (B) AS (500 µmol/L, 15 min). In vitro kinase (IVK) assays were performed in cell lysates by addition of recombinant His6-tagged C1-M-C2 domain of cardiac myosin-binding protein C as a substrate in the presence of γ32P-ATP. Substrate phosphorylation was detected by autoradiography (figure shows cropped version, full representation in online supplement). Bar charts represent the results of 5 independent experiments. Data are expressed in comparison to the WT response after HNO treatment. *P < 0.05, **P < 0.01, ***P < 0.001 by comparison against each respective unstimulated control or WT after HNO-exposure. Western immunoblot analysis for PKG↓α under non-reducing (NR) or reducing (R) conditions was performed in the same samples (figure shows cropped reducing blots, full representation in online supplement). Representative immunoblots show PKG↓α migrating at 75 kDa (monomer) and 150 kDa (dimer). Data are representative of 5 independent experiments. (C) Analysis of HNO-induced oxidation of PKG↓α. HEK-293 cells
were transfected to express PKGIα WT or various mutants Cys42Ser, Cys117Ser, Cys195Ser, Cys42/117/195Ser, exposed to NCA (100 μmol/L, 30 min, upper panel) or the respective controls namely DMSO or decomposed NCA donor compound or AS (500 μmol/L, 15 min, bottom panel) or the respective controls NaOH, nitrite (NO2−) or decomposed AS. Western immunoblot analyses for PKGIα under non-reducing (NR) or reducing (R) conditions were performed. Representative immunoblots show PKGIα migrating at 75 kDa (monomer) and 150 kDa (dimer) under non-reducing and at 75 kDa under reducing conditions. Data are representative of 5 independent experiments. n.s.: non-significant; dec: decomposed.

Activity and oxidation status of PKGIα after exposure to HNO donors was assessed by analysing substrate phosphorylation with radiolabeled γ32P-ATP and autoradiography or western immunoblotting under non-reducing conditions (Fig. 2). The activity of WT PKGIα was significantly increased in response to both HNO donors as reflected by enhanced phosphorylation of recombintant C1-M-C2 domain of cardiac myosin-binding protein C, an established substrate of PKGIα (Fig. 2A,B top panels IVK, bar charts). Replacement of Cys42 or Cys117/195 by oxidation-resistant serine significantly reduced substrate phosphorylation in response to each of the HNO donors, whilst expression of the Cys42/117/195Ser mutant completely abolished substrate phosphorylation. This is consistent with the involvement of Cys42, Cys117 and Cys195 in HNO-mediated enhancement of PKGIα activity. Isolated murine mesenteric arteries were investigated in vitro. To further substantiate these findings, cGMP-binding assays were performed in vitro. Purified bovine PKGIα was reduced by dithiothreitol (DTT; Fig. 3B; black line) or oxidised by NCA (Fig. 3B; red line). (3H)-cGMP binding to PKGIα was assessed in the presence of increasing amounts of unlabeled cGMP (Fig. 3B). This revealed significantly reduced cGMP binding to the NCA-oxidised kinase when compared to that reduced by DTT. To investigate whether the NCA-induced modifications of PKGIα modulate cGMP-induced changes in kinase activity, in vitro kinase assays were performed in the presence or absence of cGMP. Purified bovine PKGIα was reduced with DTT (Fig. 3C; open bars) or oxidised with NCA (Fig. 3C; red bars). In the absence of cGMP, PKGIα displayed basal kinase activity. Addition of cGMP to DTT-PKGIα significantly enhanced kinase activity 3.7 fold versus no cGMP. This cGMP-induced potentiation of PKGIα-activity was not observed in the NCA-oxidised kinase. These results are in accordance with the data obtained by FRET analysis and suggest that intradisulfide formation by NCA reduces subsequent cGMP binding to the kinase concluding that NCA-induced modification of PKGIα activates the kinase, but delimits its further activation by subsequent cGMP addition.

The correlation between HNO-mediated intradisulfide formation in endogenous PKGIα and vasorelaxation was investigated in isolated murine mesenteric arteries in vitro and in the skeletal muscle microcirculation in vivo in WT mice. Both HNO donors induced concentration-dependent vasorelaxation in mesenteric arteries (NCA: EC50 = 2.96 ± 0.47 μmol/L; AS: EC50 = 0.44 ± 0.6 μmol/L; 1–2 vessels from 4 mice), which was paralleled
by enhanced inter- and intradisulfide formation in PKGI\(\alpha\) as evidenced by western immunoblot analysis under non-reducing conditions (Fig. 4A, right panel). Similar concentration-dependent vasorelaxation was observed after exposure of the microcirculation to HNO donors \(\textit{in vivo}\) (Fig. 5A NCA; \(n = 68\) arterioles in 6 mice; Fig. 5B AS; \(n = 73\) arterioles in 6 mice). Arterioles studied here exhibited maximal diameters between 13 and 49 \(\mu\text{m}\) and their means were not different between the intervention groups (NCA: 27.6 ± 0.8 \(\mu\text{m}\); AS: 29.2 ± 0.9 \(\mu\text{m}\)). Vessels exhibited spontaneous tone (calculated as the quotient of resting and maximal diameter) amounting to 40 ± 1%
(ranging from 10 to 85%). This spontaneous constriction level was similar in both groups (NCA: 41 ± 2%; AS: 40 ± 2 of maximal diameter; P = 0.63). The endothelium-dependent dilator acetylcholine (10 µmol/L) induced in both groups a significant dilation (NCA: 82 ± 3%; AS: 91 ± 2 of maximal response) assuring intact endothelial function and dilator capacity. Exposure to 10 µmol/L NCA dilated the arterioles from 11.0 ± 0.9 to 24.6 ± 0.8 µm. This dilation was further enhanced after exposure to 50 µmol/L NCA (26.3 ± 0.8 µm; P < 0.05 vs. 10 µmol/L NCA). The solvent alone (0.1% DMSO) resulted in a small constriction (from 11.4 ± 0.8 to 10.0 ± 0.6 µm; P < 0.05) (Fig. 5A). Exposure to 3 µmol/L AS dilated the arterioles from 11.6 ± 0.8 to 19 ± 0.8 µm and the highest concentration used (50 µmol/L) dilated the vessels to 23.8 ± 0.8 µm. The solvent alone (0.1 mmol/L NaOH) induced a small dilatation (from 11.6 ± 0.8 to 19.6 ± 0.8 µm) (Fig. 5B).

In vivo vasorelaxation by both HNO donors was accompanied by oxidation of PKGIα into both its inter- and intradisulfide form in this tissue as shown in the western immunoblots under non-reducing conditions (Fig. 5A, B, right panels).

Discussion
Oxidation of Cys42 in PKGIα has been reported as a mechanism that regulates protein kinase activity potentially by modulating substrate targeting, independently of the second messenger cGMP. This discovery has highlighted a role for oxidants in the regulation of important physiological functions such as blood pressure.

PKGIα contains 11 cysteines, some of which have been described previously as redox sensors. Whilst interprotein disulfide formation via Cys42 has been extensively investigated and characterised, information regarding the impact of Cys117 and Cys195 oxidation on PKGIα functions is scarce. Intraprotein disulfide formation in endogenous PKGIα with HNO-mediated vasorelaxation in vitro. The effect of NCA (A) or AS (B) on vasorelaxation was assessed in mesenteric arteries from wildtype (WT; black) mice. U46619 (100 nmol/L) was used to preconstrict vessels and then increasing concentrations of NCA (0.1, 0.3, 1, 3, 10, 30, 100, 300 µmol/L) or AS (0.1, 0.3, 1, 3, 10, 30, 100 µmol/L) were administered. Experiments were performed in 1-2 vessels derived from at least 4 animals for each HNO donor. Western immunoblot analysis for PKGIα was performed under reducing (R) or non-reducing (NR) conditions in vessel homogenates exposed to 30 µmol/L NCA or 3 µmol/L AS.
bond formation was shown previously to occur after exposure of PKGIα to metal-ion induced oxidative stress in vitro. This intradisulfide, which forms in the high affinity cGMP-binding site and therefore may be considered to be in a more rational position for regulating kinase activity than the Cys42 in the leucine zipper domain, was also observed in a crystal structure of the regulatory domain of PKGIα31. This would be consistent with a constitutive disulfide. Indeed, in these studies from the Dostmann laboratory, it was suggested that this intradisulfide most likely mediated the oxidative activation of the kinase. However, it is clear from the observations reported here that cellular PKGIα is predominantly reduced under basal conditions, with the disulfide only accumulating when oxidant levels are elevated. It is likely that the intradisulfide can be observed crystallographically in recombinant protein, because the kinase is removed from cellular thiol-reducing buffers, such as thioredoxin or glutathione, that maintain solvent accessible cysteinyl thiols in their reduced state. Indeed, interprotein disulfides in PKGIα or protein kinase A RI were originally reported as constitutive structural bonds48, 49, but subsequently were shown to be absent basally without pro-oxidant interventions23, 50. Recently, S-guanylation of Cys195 by nitro-cGMP was reported as a novel modification of PKGIα, leading to persistent kinase activity51. As S-guanylation occurs at reduced thiols, this provides independent corroboration of the observations made here that the intradisulfide is largely not present in the absence of pro-oxidant conditions.

By using a differential isotope-based thiol-trapping mass spectrometry-based approach, we identified oxidative modification of Cys42, Cys117 and Cys195 in PKGIα in response to HNO. This approach has the advantage of providing quantitative information about the redox state of individual cysteines. These data showed Cys42 was susceptible to oxidation also under control conditions, whilst Cys117 and Cys195 only oxidised upon addition of HNO. Although the matching levels of oxidation of Cys117 and Cys195 caused by NCA or AS treatment is consistent with structural studies showing that PKGIα forms an intradisulfide at this location34, NOxICAT analysis cannot provide a formal demonstration of this modification. However, inter- or intradisulfides can be assessed by

Figure 5. Correlation of intradisulfide formation in endogenous PKGIα with HNO-mediated vasorelaxation in vivo. Both HNO donors induced concentration-dependent dilations in arterioles in vivo. (A) The effect of NCA was studied in 68 vessels from 6 mice. (B) The effect of AS was studied in 73 vessels from 6 mice. Data are given as mean ± SEM. *** indicates P < 0.001 for paired comparisons (t-test). Western immunoblot analysis for PKGIα was performed under reducing (R) and non-reducing (NR) conditions in isolated cremaster muscles exposed to 50 µmol/L NCA or AS. Black arrows indicate the positions of monomeric (75 kDa), inter- (150 kDa) and intradisulfide (below 150 kDa) PKGIα.
migration changes using non-reducing western immunoblotting, as the oxidations induce higher or lower molecular weight gel shifts, respectively that can be normalised by reducing agents\textsuperscript{44, 52}. Indeed, non-reducing western immunoblot analysis of cell homogenates supported the NOxICAT data, with a basal level of Cys42 interdisulfide detected under control conditions, whereas the intradisulfide only formed upon oxidant-exposure. It is notable that the intradisulfide induced by HNO had low stoichiometry, but this is because this intervention will concomitantly elevate cGMP, which as shown here competes with the formation of the intradisulfide. It is evident that the interplay of mechanisms that control PKGI\textsubscript{α} activity is complex especially as cGMP binding or Cys42 oxidation also reciprocally negatively regulates each other. Another layer of complexity is added by the observation that cGMP binding negatively influences oxidation of Cys42\textsuperscript{28, 46, 53}. A significant finding of this study is that the interdisulfide formed at Cys42 attenuated the formation of the intradisulfide, as Cys42Ser PKGI\textsubscript{α} showed markedly potentiated intradisulfide formation in response to HNO. This complex array of interrelated, interacting and modulating mechanisms may serve as a feedback that delimits over-activation or over-recruitment of the kinase to regulatory stimuli.

In addition to the molecular detection and characterisation, it was important to investigate potential functional effects of nascent intradisulfide on PKGI\textsubscript{α} activity. This was achieved using \textit{in vitro} kinase assays performed on lysates from HEK-293 cells that were transfected with WT PKGI\textsubscript{α} or various 'redox-dead' mutants of PKGI\textsubscript{α} in the presence or absence of HNO donors. An obvious conclusion from this experiment was that both HNO donors activated the WT kinase, but the activation was impaired in the various cysteine mutants that are resistant to inter- or intradisulfide formation. The overall important conclusion to reiterate is that the primary novel finding of this work is, namely that PKGI\textsubscript{α} forms an intradisulfide and this activates the kinase. This conclusion is rationally based on the structure and is consistent with S-guanylation also activating the kinase\textsuperscript{31, 51}. Perhaps either of these oxidative activation mechanisms that target Cys195 disrupts the interaction of the autoinhibitory domain with the catalytic domain to enable catalytic competence. This is in essence similar to what cGMP achieves when it binds to its high affinity-binding site that also contains this redox active cysteine. Interestingly, the HNO donors used in our study showed apparent differences concerning their impact on kinase activity. NCA-induced kinase activity was almost completely abolished in the intradisulfide-deficient mutant, whilst activity after AS treatment was greatly reduced in the Cys42Ser mutant. These results are in accordance with the NOxICAT results showing that NCA and AS modify the same spectrum of cysteines in PKGI\textsubscript{α}, albeit to a different extent. Both donors oxidise the highly susceptible Cys42, however, NCA was superior to AS with regard to oxidation of Cys117 and Cys195. One explanation for the discrepant oxidation efficiency of the HNO donors may relate to their different kinetics of HNO release. NCA has a half-life of 2 hours\textsuperscript{32}, whereas it is only 2 to 5 minutes for AS\textsuperscript{54}. Releasing initially large amounts of HNO by AS may facilitate self-consumption, potentially by reducing the amount of HNO available for target oxidation. Alternative explanations could relate to the release of byproducts released by the two different HNO donors. When AS decomposes into HNO an equivalent of nitrite and a small amount of NO\textsuperscript{35} is also generated, whereas NCA releases HNO and a thiol-reactive moiety of its scaffold\textsuperscript{33}, each of them exerts biological effects that have to be taken into account\textsuperscript{55}. Distinct behavior with regard to target oxidation and functional effects induced between different HNO donors has in fact been described previously\textsuperscript{34}. 

---

**Figure 6.** Illustration of the domain structure of the PKGI\textsubscript{α} homodimer and the localisation of cysteines that are modified by exposure to HNO donors to form inter- and intradisulfide bonds. DD: docking and dimerisation domain; AI: autoinhibitory domain; cGMP-A: high affinity cGMP-binding domain; cGMP-B: low affinity cGMP-binding domain; SW: switch helix; ATP: ATP-binding domain; sGC: soluble guanylate cyclase.
However, as both donors induce the same post-translational modification and induce the same biological action, it is logical to conclude that this is likely due to HNO release, especially as this is the only common characteristic of these compounds.

FRET experiments performed in VSMCs isolated from transgenic mice constitutively expressing a FRET-biosensor comprising the cGMP-binding sites of PKGIα, allowed the impact of HNO-induced intradisulfide on PKGIα activity to be defined in a cellular context and compared to that induced by cGMP. Indeed, exposure to HNO increased the FRET response in the absence of cGMP, suggesting activation of the kinase. This activation was comparable to that achieved by cGMP-dependent stimulation and substantiates that the kinase is activated by intradisulfide formation and is consistent with the modification relieving the autoinhibition of the catalytic domain. Exposure to increasing concentrations of cGMP did not potentiate kinase activity induced by HNO, consistent with the intradisulfide limiting cGMP binding. Future experiments involving recombinant mutant PKGIα would be valuable in establishing whether the difference in cGMP-binding to PKGIα is definitely mediated by oxidant-induced intradisulfide formation.

The HNO-mediated intradisulfide formation in PKGIα that is reported here most likely contributes mechanistically to the previously described impact of HNO on blood pressure reduction. This is supported by the fact that HNO induces significant amounts of intradisulfide in endogenous PKGIα. This activates the kinase, and so rationally significantly underlies the correlation between vasorelaxation and oxidation of endogenous PKGIα observed in two different vascular beds in vivo.

Taken together, our study describes a novel oxidative mechanism for the activation of PKGIα by HNO. Our observations are consistent with the intradisulfide inducing the direct activation of the kinase, essentially mimicking cGMP binding, with the intradisulfide mediating substrate targeting as it was recently shown29. This explains why disrupting either of these disulfides by mutagenesis impairs oxidant-induced kinase activity and substrate phosphorylation.

References

1. Eberhardt, M. et al. H₂S and NO cooperatively regulate vascular tone by activating a neuroendocrine HNO-TRPA1-CGRP signaling pathway. *Nature communications* 5, 4381, doi:10.1038/ncomms5381 (2014).
2. Cortese-Krott, M. M. et al. Key bioactive reaction products of the NO/HH₂S interaction are S/N-hybrid species, polysulfides, and nitroxyl. *Proceedings of the National Academy of Sciences of the United States of America* 112, E4651–E4660, doi:10.1073/pnas.1509927112 (2015).
3. Sabbah, H. N. et al. Nitroxyl (HNO): A novel approach for the acute treatment of heart failure. *Circ Heart Fail* 6, 1250–1258, doi:10.1161/CIRCHEARTFAILURE.111.000632 (2013).
4. Arcaro, A., Lembo, G. & Tocchetti, C. G. Nitroxyl (HNO) for treatment of acute heart failure. *Curr Heart Fail Rep* 11, 227–235, doi:10.1007/s11897-014-0210-z (2014).
5. Kemp-Harper, B. K., Horowitz, J. D. & Ritchie, R. H. Therapeutic Potential of Nitroxyl (HNO) Donors in the Management of Acute Decompensated Heart Failure. *Drugs* 76, 1337–1348, doi:10.1007/s40265-016-0631-y (2016).
6. Andrews, K. L. et al. Nitroxyl: a vasodilator of human vessels that is not susceptible to tolerance. *Clin Sci (Lond)* 129, 179–187, doi:10.1042/CS20140759 (2015).
7. Andrews, K. L. et al. A role for nitroxyl (HNO) as an endothelium-derived relaxing and hyperpolarizing factor in resistance arteries. *Br J Pharmacol* 157, 540–550, doi:10.1111/1476-5381.2009.01510.x (2009).
8. Longhi-Balbinot, D. T. et al. The nitroxyl donor, Angeli’s salt, reduces chronic constriction injury-induced neuropathic pain. *Pain* 158, 1947–1956, doi:10.1097/01.alx.0000597627.17034.52 (2007).
9. Shoman, M. E. & Aly, O. M. Nitroxyl (HNO): A possible strategy for fighting cancer. *Curr Top Med Chem* 16, 2464–2470 (2016).
10. Gao, W. D. et al. Oxidation of cysteines activates cGMP-dependent protein kinase. *Oxidation of cysteines activates cGMP-dependent protein kinase. Nature communications* 7: 9938, doi:10.1038/s41598-017-09275-1 (2016).
11. Zhu, G. et al. Soluble guanylate cyclase is required for systemic vasodilation but not positive inotropy induced by nitroxyl in the mouse. *Hypertension* 65, 385–392, doi:10.1161/HYPERTENSIONAHA.114.04285 (2015).
12. Miranda, K. M. et al. Further evidence for distinct reactive intermediates from nitroxyl and peroxynitrite: effects of buffer composition on the chemistry of Angeli’s salt and synthetic peroxyxinitrite. *Archives of biochemistry and biophysics* 401, 134–144, doi:10.1006/abbi.2000.1931 (2002).
13. Orescanin, Z. S., Milovanovic, S. R., Spasic, S. D., Jones, D. R. & Spasic, M. B. Different responses of mesenteric artery from normotensive and spontaneously hypertensive rats to nitric oxide and its redox congeners. *Cardiovasc Res* 79, 315–322 (2007).
14. Donzelli, S. et al. Pharmacological characterization of 1-nitrosocylohexyl acetate, a long-acting nitroxyl donor that shows vasorelaxant and antaggregatory effects. *J Pharmacol Exp Ther* 334, 339–347, doi:10.1124/jpet.111.199836 (2013).
15. Bullen, M. L. et al. Vasorelaxant and antaggregatory actions of the nitroxyl donor isopropylamine NONOate are maintained in hypercholesterolemia. *Am J Physiol Heart Circ Physiol* 301, H1405–1414, doi:10.1152/ajpheart.00489.2011 (2011).
16. Miller, T. W. et al. The effects of nitroxyl (HNO) on soluble guanylate cyclase activity: interactions at ferrous heme and cysteine thiols. *The Journal of biological chemistry* 284, 21788–21796, doi:10.1074/jbc.M109.014282 (2009).
17. Favaloro, J. L. & Kemp-Harper, B. K. The nitroxyl anion (HNO) is a potent dilator of rat coronary vasculature. *Cardiovasc Res* 73, 587–596, doi:10.1016/j.cardiores.2006.11.018 (2007).
18. Paolucci, N. et al. Nitroxyl anion exerts redox-sensitive positive cardiac inotropy in vivo by calcitonin gene-related peptide signaling. *Proceedings of the National Academy of Sciences of the United States of America* 98, 10463–10468, doi:10.1073/pnas.181191198 (2001).
19. Hofmann, F., Bernhard, D., Lukowski, R. & Weinmeister, P. cGMP regulated protein kinases (cGK). *Handb Exp Pharmacol* 137–162, doi:10.1007/978-3-540-63966-9_8 (2009).
20. Geiselhöringer, A., Gaiss, M., Hofmann, F. & Schlossmann, J. Distribution of IRAg and cGKI isoforms in murine tissues. *FEBS Lett* 557, 19–22, doi:10.1016/j.febslet.2004.08.030 (2004).
21. Feil, R., Lohmann, S. M., de Jonge, H., Walter, U. & Hofmann, F. Cyclic GMP-dependent protein kinases and the cardiovascular system: insights from genetically modified mice. *Circ Res* 93, 907–916, doi:10.1161/01.RES.0000103990.68771.CC (2003).
22. Scholten, A., Aye, T. T. & Heck, A. J. A multi-angular mass spectrometric view at cyclic nucleotide dependent protein kinases: in vivo characterization and structure/function relationships. *Mass Spectrom Rev* 27, 331–353, doi:10.1002/mas.20166 (2008).
23. Burgoyne, J. R. et al. Cysteine redox sensor in PKGIα enables oxidant-induced activation. *Science* 317, 1393–1397, doi:10.1126/science.1144318 (2007).
24. Landgraf, W., Regulla, S., Meyer, H. E. & Hofmann, F. Oxidation of cysteines activates cGMP-dependent protein kinase. *The Journal of biological chemistry* 266, 16305–16311 (1991).
Kalyanaraman, H., Zhuang, S., Pilz, R. B. & Casteel, D. E. The Activity of cGMP-dependent Protein Kinase 1alpha is not Directly Regulated by oxidation-induced disulfide formation at cysteine 43. *The Journal of biological chemistry*. doi:10.1074/jbc.C117.787358 (2017).

Grassie, M. E., Moffat, L. D., Walsh, M. P. & MacDonald, J. A. The myosin phosphatase targeting protein (MYPPT) family: a regulated mechanism for achieving substrate specificity of the catalytic subunit of protein phosphatase type 1delta. *Archives of biochemistry and biophysics* **510**, 147–159, doi:10.1016/j.abb.2011.01.018 (2011).

Przybyazhna, O. & Eaton, P. Redox regulation of cGMP-dependent protein kinase 1alpha in the cardiovascular system. *Frontiers in pharmacology* **6**, 139, doi:10.3389/fphar.2015.00139 (2015).

Przybyazhna, O. et al. Phosphodiesterase 5 inhibition limits doxorubicin-induced heart failure by attenuating protein kinase G alpha oxidation. *The journal of biological chemistry* **291**, 17427–17436, doi:10.1074/jbc.M116.724070 (2016).

Scotch, J. et al. Disulfide-activated protein kinase G 1alpha regulates cardiac diastolic relaxation and fine-tunes the Frank-Starling response. *Nature communications* **7**, 13187, doi:10.1038/ncomms13187 (2016).

Przybyazhna, O., Rudyk, O. & Eaton, P. Single atom substitution in mouse protein kinase G eliminates oxidant sensing to cause hypertension. *Nat Med* **18**, 286–290, doi:10.1038/nmm.2603 (2012).

Osborne, B. W. et al. Crystal structure of cGMP-dependent protein kinase reveals novel site of interchain communication. *Structure* **19**, 1317–1327, doi:10.1016/j.str.2011.06.012 (2011).

Shoman, M. E. et al. Acyloxy nitroso compounds as nitroxyl (HNO) donors: kinetics, reactions with thiols, and vasodilation properties. *J Med Chem* **54**, 1059–1070, doi:10.1021/jm101432z (2011).

Sha, X., Isbell, T. S., Patel, R. P., Day, C. S. & King, S. B. Hydrolysis of acyloxy nitroso compounds yields nitroxyl (HNO). *The Journal of the American Chemical Society* **128**, 9687–9692, doi:10.1021/ja062363a (2006).

Rudyk, O. et al. Protein kinase D selectively targets cardiac troponin I and regulates myofilament Ca2+ sensitivity in ventricular myocytes. *Circ Res* **100**, 864–873, doi:10.1161/RES.0000260809.15393.f (2007).

Lindemann, C. & Leichert, L. I. Quantitative redox proteomics: the NOxICAT method. *Methods Mol Biol* **532**, 153–163, doi:10.1007/978-1-61779-459-3_6 (2012).

Thunemann, M. et al. Transgenic mice for cGMP imaging. *Circ Res* **113**, 365–371, doi:10.1161/CIRCRESAHA.113.301063 (2013).

Russwurm, M. et al. Design of fluorescence resonance energy transfer (FRET)-based cGMP indicators: a systematic approach. *Biochem J* **407**, 69–77, doi:10.1042/BJ20070348 (2007).

Schindelin, J. et al. Fiji: an open-source platform for biological-image analysis. *Nat Methods* **9**, 676–682, doi:10.1038/nmeth.1922 (2012).

Thunemann, M., Fomin, N., Krawutschke, C., Russwurm, M. & Feil, R. Visualization of cGMP with cG biosensors. *Methods Mol Biol* **1020**, 89–110, doi:10.1007/978-1-62703-459-3_6 (2013).

Stathopoulou, K. et al. S-glutathiolation impairs phosphoregulation and function of cardiac myosin-binding protein C in human heart failure. *FASEB J* **30**, 1849–1864, doi:10.1096/fj.201500048 (2016).

Thoven, R. et al. Molecular screen identifies cardiac myosin-binding protein C as a protein kinase G-lalpha substrate. *Circ Heart Fail* **8**, 1115–1122, doi:10.1161/CIRCHEARTFAILURE.115.002908 (2015).

de Wit, C., von Bismarck, P. & Pohl, U. Mediator role of prostaglandins in acetylcholine-induced vasodilation and control of resting vascular diameter in the hamster cremaster microcirculation in vivo. *J Vasc Res* **30**, 272–278 (1993).

Eaton, P. Protein thiol oxidation in health and disease: techniques for measuring disulfides and related modifications in complex protein mixtures. *Free Radic Biol Med* **48**, 1189–1199, doi:10.1016/j.freeradbiomed.2010.03.037 (2010).

Rudyk, O. et al. Protein kinase G oxidation is a major cause of injury during sepsis. *Proceedings of the National Academy of Sciences of the United States of America* **110**, 9909–9913, doi:10.1073/pnas.1301026110 (2013).

Burgoyne, J. R., Przybyazhna, O., Rudyk, O. & Eaton, P. cGMP-dependent activation of protein kinase G precludes disulphide activation: implications for blood pressure control. *Hypertension* **60**, 1301–1308, doi:10.1161/HYPERTENSIONAHA.112.198754 (2012).

Takio, K. et al. Guanosine cyclic 3′,5′-phosphate dependent protein kinase, a chimeric protein homologous with two separate protein families. *Biochemistry* **23**, 4207–4218 (1984).

Sarma, G. N. et al. Structure of D-ACKAP2:PKA RI complex: insights into AKAP specificity and selectivity. *Structure* **18**, 155–166, doi:10.1016/j.str.2009.12.012 (2010).

McKenzie, C. E. & Gill, G. N. Structural analysis of cGMP-dependent protein kinase G protein limited proteolysis. *The Journal of biological chemistry* **255**, 7067–7070 (1980).

Brennan, J. P. et al. Oxidant-induced activation of type I protein kinase A is mediated by RI subunit interprotein disulfide bond formation. *The Journal of biological chemistry* **281**, 21827–21836, doi:10.1074/jbc.M603952200 (2006).

Akashi, S. et al. Persistent Activation of cGMP-Dependent protein kinase G by a nitrated cyclic nucleotide via site specific protein S-glycation. *Biochemistry* **55**, 751–761, doi:10.1021/acs.biochem.5b00774 (2016).

Rudyk, O. & Eaton, P. Biochemical methods for monitoring protein thiol redox states in biological systems. *Redox Biol* **2**, 803–813, doi:10.1016/j.redox.2014.06.005 (2014).

Muller, P. M. et al. H2O2 lowers the cytosolic Ca2+ concentration via activation of cGMP-dependent protein kinase 1alpha. *Free Radic Biol Med* **53**, 1574–1583, doi:10.1016/j.freeradbiomed.2012.08.011 (2012).

Hughes, R. M. & Cammack, R. Synthesis, chemistry, and applications of nitroxyl ion releasers sodium tioxidonitrate or Angelil's salt and Piloty's acid. *Methods Enzymol* **301**, 279–287 (1999).

Dutton, A. S., Fukushima, M. J. & Houk, K. N. Mechanisms of HNO and NO production from Angelil's salt: density functional and CBS-QB3 theory predictions. *Journal of the American Chemical Society* **126**, 3795–3800, doi:10.1021/ja0391614 (2004).

Lundberg, J. O., Weitberg, E. & Gladwin, M. T. The nitrate-nitrite-nitric oxide pathway in physiology and therapeutics. *Nature reviews. Drug discovery* **7**, 156–167, doi:10.1038/nrd2466 (2008).

Acknowledgements

We thank Prof. Dr. Thomas Braulke for advice on cGMP-binding assays and Dagmara Nelson for support in the UKE isotope facility. This work was supported by the German Center for Cardiovascular Research and the German Ministry of Research and Education (SoD, MG, KS, KS, SiD, VP, HS, SS, VON, Cdw, FC), the Deutsche Forschungsgemeinschaft (SPP 1710 grant LE 2905/1-1 to LIL; CU 53/2-1 to FC; FOR 2060 projects FE 438/5-1, FE 438/6-1 to RF), the Werner-Otto-Stiftung (22/86 to FC; 4/83 to SoD), the British Heart Foundation (PE) and the European Research Council (ERC Advanced award to PE).
Author Contributions
S.o.D. contributed to NOxICAT, cGMP-binding assays, cell transfections, pharmacological treatments and western immunoblotting; M.G. contributed to cell transfections, pharmacological treatments, cGMP-binding assays, mesenteric artery isolation and western immunoblot analysis; K.j.S. performed vasodilation experiments in cremaster arterioles; M.W. performed FRET experiments in VSMCs; K.o.S. contributed to P.K.G. activity assays, cell transfections, pharmacological treatments and western immunoblot analysis; S.i.D. contributed to mesenteric artery isolation and western immunoblot analysis; O.P. performed vasorelaxation experiments in isolated mesenteric arteries; V.P. helped with initial optimisation of experiments; J.S. generated the P.K.G. mutant constructs; C.D. helped with cGMP-binding assays; HS provided antibody advise; E.B. provided purified bovine PKGIα; A.K. helped with vasorelaxation experiments in mesenteric arteries; SS discussed data; S.B.K. provided NCA; VON provided advise with NOxICAT experiments and data analysis; R.F. designed FRET experiments in VSMCs and provided his transgenic mice; P.E. designed the study, discussed data and wrote the manuscript; F.C. contributed to NOxICAT experiments, P.K.G. activity and binding assay, vasodilation experiments in cremaster arterioles, designed the study, obtained funding and wrote the manuscript. All authors provided corrections and proofread the manuscript.

Additional Information
Supplementary information accompanies this paper at doi:10.1038/s41598-017-09275-1

Competing Interests: The authors declare that they have no competing interests.

Publisher's note: Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2017