Monitoring Vascular Permeability and Remodeling After Endothelial Injury in a Murine Model Using a Magnetic Resonance Albumin-Binding Contrast Agent

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Background—Despite the beneficial effects of vascular interventions, these procedures may damage the endothelium leading to increased vascular permeability and remodeling. Re-endothelialization of the vessel wall, with functionally and structurally intact cells, is controlled by endothelial nitric oxide synthase (NOS3) and is crucial for attenuating adverse effects after injury. We investigated the applicability of the albumin-binding MR contrast agent, gadofosveset, to noninvasively monitor focal changes in vascular permeability and remodeling, after injury, in NOS3-knockout (NOS3−/−) and wild-type (WT) mice in vivo.

Methods and Results—WT and NOS3−/− mice were imaged at 7, 15, and 30 days after aortic denudation or sham-surgery. T1 mapping (R1=1/T1, s−1) and delayed-enhanced MRI were used as measurements of vascular permeability (R1) and remodeling (vessel wall enhancement, mm2) after gadofosveset injection, respectively. Denudation resulted in higher vascular permeability and vessel wall enhancement 7 days after injury in both strains compared with sham-operated animals. However, impaired re-endothelialization and increased neovascularization in NOS3−/− mice resulted in significantly higher R1 at 15 and 30 days post injury compared with WT mice that showed re-endothelialization and lack of neovascularization (R1 [s−1]=15 days: NOS3−/−4.02 [interquartile range, IQR, 3.77–4.41] versus WT 2.39 [IQR, 2.35–2.92]; 30 days: NOS3−/−4.23 [IQR, 3.94–4.68] versus WT 2.64 [IQR, 2.33–2.80]). Similarly, vessel wall enhancement was higher in NOS3−/− but recovered in WT mice (area [mm2]=15 days: NOS3−/−5.20 [IQR, 4.68–6.80] versus WT 2.13 [IQR, 0.97–3.31]; 30 days: NOS3−/−7.35 [IQR, 5.66–8.61] versus WT 1.60 [IQR, 1.40–3.18]). Ex vivo histological studies corroborated the MRI findings.

Conclusions—We demonstrate that increased vascular permeability and remodeling, after injury, can be assessed noninvasively using an albumin-binding MR contrast agent and may be used as surrogate markers for evaluating the healing response of the vessel wall after injury. (Circ Cardiovasc Imaging. 2015;8:e002417. DOI: 10.1161/CIRCIMAGING.114.002417.)

Key Words: gadofosveset trisodium ■ magnetic resonance imaging ■ permeability ■ vascular remodeling

Despite the beneficial effects of percutaneous transluminal coronary angioplasty and stent implantation, these procedures may damage the vessel wall, particularly the endothelial layer, leading to increased vascular permeability and remodeling. Therefore, noninvasive assessment of these processes may provide a more comprehensive understanding of the focal responses of the vessel wall to injury.

See Clinical Perspective

In healthy vessels, functional endothelium produces nitric oxide through endothelial nitric oxide synthase (eNOS or NOS3). Nitric oxide controls the vasoactivity of the vessel wall, inhibits platelet and leukocyte adhesion and smooth muscle cell proliferation and migration, promotes re-endothelialization, and regulates endothelial cell and tight junction morphology and vascular permeability.1,2 Dysfunctional endothelium because of vascular injury or cardiovascular risk factors5–7 is characterized by reduced NOS3-derived nitric oxide resulting in the attenuation of endothelial-dependent vasodilation5 and accelerated vascular remodeling.9–12

Genetically modified NOS3−/− mice (1) vasodilate only in response to endothelial-independent stressors,13 (2) have...
impaired re-endothelialization and (3) have increased inflammatory response after injury collectively leading to increased vascular remodeling. However, despite some of the similarities between the vascular response observed in NOS3−/− mice and patients that undergo percutaneous interventions, endothelial denudation followed by vascular remodeling, as seen in this model, may represent the worst case outcome compared with human interventions. Interestingly, in a rabbit model, viral and nonviral transfection of NOS3 and placement of NOS3 gene–eluting stents significantly reduced vascular remodeling and improved re-endothelialization.

Uptake into the vessel wall of albumin-binding dyes (Evans blue, I-albumin, rhodamine) has been traditionally used as a surrogate marker of vascular permeability in ex vivo experiments. MRI is a noninvasive modality that allows the study of both endothelial function and vascular remodeling with high spatial resolution. An albumin-binding MRI contrast agent, gadofosveset, has been recently used as the in vivo counterpart of albumin-binding dyes and allowed the assessment of focal changes in vascular permeability and vascular remodeling in atherosclerosis, and chronic myocardial infarction. Gadofosveset is a clinically approved gadolinium-based, blood pool contrast agent. After injection in mice, ≈70% of circulating gadofosveset binds reversibly to serum albumin and this increases its r1 relaxivity by 5- to 10-fold (r1=25 mmol L−1 s−1) compared with the free fraction (r1=5 mmol L−1 s−1). Uptake of gadofosveset into the vessel wall of pathological tissues has been shown to occur through damaged endothelium and leaky neovessels.

Endothelial damage and vascular remodeling may occur after vascular injury. Re-endothelialization of the vessel wall with functionally and structurally intact endothelial cells is crucial for minimizing vascular remodeling and maintaining the long-term patency of the treated vessel. We hypothesized that augmented vascular permeability because of impaired re-endothelialization and increased neovascularization, observed after injury in the NOS3−/− mice, is associated with enlarged vascular remodeling that could be monitored noninvasively using gadofosveset-enhanced MRI in vivo.

**Methods**

**Animals/Endothelial Denudation**

Wild-type (WT) C57BL/6 mice and B6.129P2-Nos3tm1Unc/J knock-out mice (8–12 weeks old) were purchased from the Jackson Laboratory (Bar Harbor, ME). Surgery was performed under isoflurane anesthesia using a dissection microscope (Leica, Wetzlar, Germany). Before surgery, the abdominal fur was removed with lotion hair and the surgical field was disinfected. After a midline incision of the abdomen, the aorta, renal, and iliac arteries were exposed and isolated. Subsequently, 2 ligatures (surgical silk, size 4-0, Aragó, Zaragoza, Spain) were placed around the aorta, below the renal arteries, and above the iliac bifurcation, respectively (Figure 1A; left). A 30G syringe was introduced into the abdominal aorta via an aortic puncture followed by a bolus infusion of 1-mL saline buffer. This procedure was repeated 3× to remove the endothelial cells (Figure 1A; right). Finally, the needle was removed, the aortic puncture was repaired, and the ligatures were removed to restore blood flow. The muscle and skin incisions were sutured (5-0 Vicryl, Ethicon Inc, Somerville, NJ), and the animals were allowed to recover. Each surgery lasted 30 minutes. Sham-operated animals underwent the same surgical procedure described above without injuring the endothelial layer. All procedures used in these studies were performed in accordance with the guidelines of the UK Home Office.

**In Vivo MRI Protocol at 3T**

In vivo vessel wall imaging was performed using a Philips Achieva MR scanner (Philips Healthcare, Best, The Netherlands) equipped
with a clinical gradient system (30 mT m⁻¹, 200 mT m⁻¹ ms⁻¹) and a single-loop surface coil (diameter=47 mm). Mice were imaged before, 7, 15, and 30 days after the surgery (Figure I in the Data Supplement). Four sham-operated mice and 6 injured mice per strain were imaged per time point. Animals were imaged in supine position before and 30 minutes after intravenous administration of 0.03 mmol/kg gadofosveset (Lantheus Medical Imaging, North Billerica, MA). Anesthesia was induced with 5% and maintained with 1% to 2% isoflurane during the MRI experiments. After 3-dimensional (3D) gradient echo scout scans, contrast-enhanced angiography images were acquired for visualization of the vasculature with a field of view (FOV)=35×35×16 mm, matrix=32×32×233, in-plane resolution=0.15×0.15 mm, slice thickness=0.5 mm, repetition time/echo time (TR/TE)=28/6 ms, and flip angle=40°. The maximum intensity projection images were used to plan the subsequent delayed enhancement (DE) and T₁ mapping scans. A 2D-look–Locker sequence planned perpendicular to the aorta was used to determine the optimal inversion time (TI) for blood signal nulling. Acquisition parameters were FOV=30×30 mm, matrix=76×75, in-plane resolution=0.4×0.4 mm, slice thickness=2 mm, TR/TE=18/8.3 ms, TR between subsequent IR pulses=1000 ms, and flip angle=10°. An inversion-recovery 3D fast gradient echo sequence was acquired 30 minutes post injection and was used for DE-MRI and visualization of contrast uptake. Imaging parameters were FOV=35×35×12 mm, matrix=348×348, in-plane resolution=0.1×0.1×1 mm (normal vessel wall thickness=0.2 mm), slice thickness=24 mm, TR/TE=27/8.2 ms, TR between subsequent infrared pulses=1000 ms, and flip angle=30°. T₁ mapping was performed using a sequence that uses 2 nonselective inversion pulses each followed by 8-segmented readouts. The 2 imaging trains result in a set of 16 images per slice with increasing inversion times ranging from 20 to 2000 ms. For T₁ mapping, the acquisition parameters were FOV=36×22×10 mm, matrix=180×102, in-plane resolution=0.2×0.21×0.5 mm, slice thickness=20 mm, TR/TE=9/4.6 ms, and flip angle=10°. R₁ values were computed on a pixel-by-pixel basis using in house Matlab software.²⁴

MRI Data Analysis
Vascular remodeling was calculated by manually segmenting the visually contrast-enhanced area of the vessel wall as seen on the DE-MRI images using OsiriX (OsiriX Foundation, Geneva, Switzerland). To ensure that the segmented area encompassed the vessel wall, the DE-MRI images were coregistered and fused with the magnetic resonance angiographic images. DE-MRI areas were measured on consecutive slices along the aorta and were added for each animal to express the total extent of vessel wall remodeling. R₁ values were measured on consecutive slices along the aorta and were averaged for each animal.

Histology

Tissue Harvesting
For all histological procedures, mice were anesthetized with isoflurane and perfused through the left ventricle with physiological saline (for transmission electron microscopic [TEM] studies and Evans blue dye [EBD] staining) or with physiological saline followed by 10% buffered formalin (for hematoxylin and eosin [H&E] staining and immunohistochemistry).

EBD Staining
Vascular permeability was assessed by visualizing and quantifying the leakage of EBD into the vessel wall. Mice were anesthetized, and EBD (0.1 mL of 1% dye in saline) was injected in the left ventricle. After 10 minutes, the injured abdominal aorta was isolated, dried, and weighed. EBD was extracted by incubation in formamide for 24 hours at 60°C and the absorbance was measured at 620 nm. Standard curves for pure EBD were used to calculate the total amount of dye in the tissue. Four animals were analyzed per group at each time point.

H&E staining and immunohistochemistry
Abdominal aortas were fixed in 10% buffered formalin for 48 hours at 4°C, embedded in paraffin, and sectioned transversely (5-μm thick). H&E staining was used to investigate vessel wall morphology. Immunohistochemistry using antibodies to platelet endothelial cell adhesion (PECAM-1; 1:50, BD Pharmingen, Oxford, UK) and serum albumin (1:5000, Abcam, Cambridge, UK) was used to study endothelial cells and serum albumin present in the vessel wall, respectively. Five slices (5 μm) were stained per animal. Digital images were used to analyze the histological sections using ImageJ (NIH).

Vascular remodeling was used to describe 2 changes that occur in the vessel wall after injury: neo-intima formation that involves the inward deposition of cellular components and extracellular matrix proteins at the luminal surface, and the accumulation of extracellular matrix/collagen and neovessels at the periphery of the adventitial contour of the vessel wall. Vascular remodeling was calculated using the H&E images [adventitia area–the luminal area (mm²)]. Immunopositive areas were analyzed on digital images by computerized planimetry. The immunopositive area was segmented on the images and expressed as normalized albumin area (%; albumin immunopositive area/area of vascular remodeling×100). Individual measurements performed within each animal were averaged and 4 animals were analyzed per group at each time point.

Transmission Electron Microscopy
Preselected segments of the aorta that showed enhancement on MRI (n=1 per time point per group) were pinned down and fixed in 2% glutaraldehyde in 0.1 mol/L sodium phosphate buffer (pH 7.4) for 2 hours, washed with sodium phosphate buffer for 2 hours, and post fixed in 1% OsO₄ for 2 hours. Each aorta was divided into 3 levels (upper, middle, and low) starting from the left renal branch to the iliac bifurcation and processed for histological analysis. Subsequently, 3 sections were examined from the aorta of each animal. Samples were dehydrated through a graded series of ethanol and embedded in epoxy resin. Semithin sections (0.2 μm) were stained with toluidine blue for light microscopy examinations and were used to guide sampling for TEM studies. Thin sections (0.09 μm) were collected on 150 mesh copper grids and double stained with uranyl acetate and lead citrate for electron microscopy examinations (H7650, Hitachi, Tokyo, Japan).

Statistical Analysis
GraphPad Prism 5.00 was used for the statistical analysis. Two-group comparisons of continuous variables were performed with a Mann–Whitney nonparametric exact test after the variables were ranked. The data are presented as the median±interquartile range, and P values <0.05 were used to define statistical significance.

Results

Endothelial Denudation Protocol
Histological validation and characterization of the surgical protocol was shown in our previous work, where we show that this procedure removes virtually all endothelial cells over the
area being subjected to the saline infusion procedure. Representative images of the surgical protocol are illustrated in Figure 1A. EBD staining, a histological marker of endothelial permeability, was performed in a subgroup of WT animals, showing endothelial denudation 1 hour after surgery. En face photographs of the extracted aortas showed increased uptake of EBD demonstrating higher vascular permeability in injured animals compared with the sham-operated mice (Figure 1B). H&E staining and PECAM-1 immunohistochemistry further demonstrate that the endothelium was successfully removed after vascular injury compared with sham-operated animals (Figure 1C and 1D).

**MRI Findings**

Vascular permeability was measured using relaxation rate (R1) maps that allow in situ quantification of the gadolinium concentration in the vessel wall (Figure 2). This analysis revealed significantly higher R1 in denuded animals, in both strains, 7 days after injury compared with sham-operated mice. Importantly, NOS3−/− mice showed significantly higher R1, 15 and 30 days after injury compared with WT mice that show restoration of the R1 values close to baseline (Figure 2; graph). These results suggest healing of the vessel wall and normalization of vascular permeability in WT whereas in NOS3−/− mice vascular permeability remained increased after injury.

Vascular remodeling was quantified using DE-MR images after administration of gadofosveset. Similar to the R1 values, DE-MRI areas were higher in the abdominal aorta of both WT and NOS3−/− mice 7 days after injury compared with sham-operated animals (Figure 3, second row). Importantly, in NOS3−/− mice, the area of vessel wall enhancement remained higher 15 and 30 days after injury compared with that observed in WT mice at the same time points (Figure 3, third and fourth rows). The area of vessel wall enhancement in WT mice 15 and 30 days post injury was similar to that observed in sham-operated animals (Figure 3, first column). The quantitative changes in the DE-MRI area are also illustrated (Figure 3; graph). Finally, correlation analysis showed a strong linear correlation between DE-MRI and R1 (r=0.702; P<0.001; Figure IIA in the Data Supplement). Collectively, these data suggest that the differential response in vascular permeability and remodeling, after injury, between NOS3−/− and WT can be assessed noninvasively by R1 mapping and DE-MRI after gadofosveset administration.

**Histological Findings**

Ex vivo histological analysis corroborated the in vivo MRI data. EBD (Figure 4A) and albumin immunohistochemistry (Figure 4B) showed that at 7 days after injury both NOS3−/− and WT had focal uptake of the dye and albumin immunopositive areas indicating uptake of the EBD and albumin into the vessel wall because of increased vascular permeability. However, NOS3−/− showed persistent and significantly higher uptake of EBD and albumin in the injured segment of the abdominal aorta at 15 and 30 days post injury compared with WT mice. Vessels from sham-operated mice, of both strains, showed low uptake of EBD and albumin immunopositive areas located primarily in the adventitial layer and to a lesser extent in the endothelium.

PECAM-1 immunohistochemistry showed impaired re-endothelialization in NOS3−/− compared with WT mice after injury (Figure 5A). Furthermore, PECAM-1 immunohistochemistry revealed neovessel formation within the remodeled area (Figure 5B), breaks in-between adjacent endothelial cells (Figure 5C), and regions of denudation (Figure 5D) in NOS3−/− mice but not in WT mice, which showed complete re-endothelialization 30 days after surgery (Figure 5E). H&E staining showed increased vascular remodeling in NOS3−/− compared with WT mice after injury (Figure 5F). Quantitative differences in vessel wall area between the 2 mice strains were significant at 7, 15, and 30 days post injury (Figure 5F; graph).

Qualitative ultrastructural changes after vessel wall injury were studied with TEM (Figure III in the Data Supplement). At 7 days post injury, the endothelial layer was absent in both strains. In WT mice, the vessel wall was re-endothelialized with endothelial cells that appeared elongated and had cytoplasmic vacuolization and microvilli at 15 and 30 days. Conversely, NOS3−/− mice showed re-endothelialization with cuboidal endothelial cells and partial fragmentation of elastin fibers.

Correlation analysis between the MRI measurements and histology showed a strong linear correlation between the vessel wall R1 and EBD uptake (r=0.782; P<0.001; Figure IIB in the data Supplement) and between the R1 and normalized albumin area (r=0.83; P<0.001; Figure IIC in the Data Supplement). There was also a statistically significant correlation between EBD and normalized albumin area (r=0.80; P<0.001; data not shown). The correlation between the DE-MRI and histology was not statistically significant when all time points after injury were included (r=0.41; P=0.02; Figure IID in the Data Supplement). However, the correlation was significant.
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\( r = 0.78; \) \( P < 0.001; \) Figure IID in the Data Supplement) after the acute phase (7 days), when measurements derived from sham-operated animals and at 15 and 30 days post injury were included.

### Discussion

In this study, we investigated the applicability of the albumin-binding MR contrast agent, gadofosveset, to noninvasively monitor focal changes in vascular permeability and remodeling, after injury, in NOS3-knockout (NOS3\(^{−/−}\)) and WT mice in vivo. We demonstrate that (1) increased vascular permeability, measured in situ using T1 mapping, showed higher vessel wall R1 rates in NOS3\(^{−/−}\) mice at 15 and 30 days postinjury compared with WT animals, and correlated with measures of vascular permeability, including EBD and albumin immunohistochemistry. In addition, histological staining, including PECAM-1, showed impaired re-endothelialization and neovessel formation in NOS3\(^{−/−}\) compared with WT animals. (2) DE-MRI using an albumin-binding contrast agent, gadofosveset, enables in vivo quantification of vascular remodeling after injury. NOS3\(^{−/−}\) mice showed significantly larger vascular remodeling, measured as increased contrast-enhanced area by MRI, compared with WT animals 15 and 30 days after injury. Ex vivo H&E validated the presence of extensive vascular remodeling in NOS3\(^{−/−}\) mice compared with WT animals. We propose that uptake of gadofosveset into the vessel wall measured with T1 mapping and DE-MRI could provide surrogate markers to assess the healing response of the vessel wall in response to vascular injury.

Several experimental murine models of induced vascular injury have been described.\(^{34–37}\) Arterial ligation and mechanically induced denudation are among the 2 most common procedures. Although the model of arterial ligation provided important insights in the biological processes involved in vascular remodeling, it did not allow evaluation of the role of the endothelium that was 1 of the primary goals of our study. Mechanically induced vascular injury using guide wires causes endothelial denudation but it could also tear the elastic lamina. To overcome this complication, non–guide wire methods have been developed\(^ {12,36}\) that showed similar histological vessel wall changes to those observed after percutaneous interventions, both in animal models\(^ {37,38}\) and humans.\(^ {39}\) Particularly, using a non–guide wire surgical procedure, we characterized the differences in the vascular response between NOS3\(^{−/−}\) and WT mice by histology.\(^ {12}\)

In our study, we found that the highest vascular permeability occurred 7 days after vascular injury, as measured by in vivo and ex vivo methods, and was similar between the 2 strains. However, in vivo measurements of vascular remodeling, as measured by DE-MRI at 7 days overestimated the vessel wall area compared with that measured by H&E staining.
This might be because of the lack of the endothelial barrier, which allows for unobstructed leakage of molecules in the blood (e.g., albumin, gadofosveset) into the vessel wall, leading to high permeability that is not necessarily accompanied by the presence of vascular remodeling. This data suggest that in an acute phase of vascular injury, changes in permeability occur first and precede formation of vascular remodeling. Whether increased permeability will be followed by
remodeling may partly depend on the subsequent re-endothelialization of the vessel wall. However, after the acute phase of endothelial injury (7 days), DE-MRI was in good agreement with vascular remodeling quantified by histology. For this reason, the differences in the long-term vascular response between NOS3−/− and WT mice could be deciphered. NOS3−/− mice showed persistently higher R1 values and vessel wall enhancement on DE-MRI at 15 and 30 days after injury, which correlated with histological measures of both vascular permeability and remodeling, respectively. Conversely, R1 and DE-MRI values decreased close to baseline levels in WT animals. It is unclear if gadofosveset interacts nonspecifically with matrix proteins or other components of the vessel wall, but in our previous publication, we have shown that gadofosveset accumulates within the vessel wall without showing preferential localization in specific plaque components using x-ray film gadolinium spectra. In this study, we propose that longitudinal measurements of both vascular permeability and remodeling in a single MRI examination provide a more comprehensive understanding of the focal responses of the vessel wall to injury.

Our histological analysis using PECAM-1 immunohistochemistry showed impaired re-endothelialization and neovessel formation only in NOS3−/− mice 15 and 30 days after injury. Ultrastructural description using TEM showed some evidences of morphological changes of the endothelium in both NOS3−/− and WT animals, including large cytoplasmatic vacuoles, altered endothelial cells shape, microvilli, and partial fragmentation of elastin fibers. Similar morphological changes have been previously reported in specimens from both animal models40 and humans41 with different cardiovascular risk factors. Nevertheless, a comprehensive morphometric and quantitative analysis of the TEM images would be required to elucidate the mechanism responsible for the differences in permeability observed between the 2 strains. Impaired re-endothelialization and increased neovascularization in NOS3−/− mice resulted in increased vascular permeability as measured by EBD and vascular remodeling measured by H&E. These results are in good agreement with previous histological studies that showed the importance of NOS3-derived nitric oxide in a large number of vascular diseases.15,42

In our study, in vivo measurements of R1 are complimentary to that of EBD. Although the 2 techniques rely on the same biological process, the units of each one are not identical and thus cannot be directly compared. The patterns of the changes, however, between animals and time points are similar between these 2 techniques (see Figures 2 and 4A). The measurements of DE-MRI area and H&E expressed in mm² might be more suitable for direct comparison (Figures 3 and 5F). However, in our study, the DE-MRI areas measured on consecutive slices along the aorta and were added for each animal to express the extent of vessel wall remodeling occurring in each time point. Conversely, histology was not performed continuously along the aorta and were added for each animal to express the extent of vessel wall remodeling occurring in each time point. The views expressed are those of the authors and not necessarily those of the NHS, the NIHR, or the Department of Health. (4) the National Institute for Health Research (NIHR) comprehensive Biomedical Research Centre award to Guy’s and St Thomas’ National Health Service (NHS) Foundation Trust in partnership with King’s College London and King’s College Hospital NHS Foundation Trust. The views expressed are those of the authors and not necessarily those of the NIH, the NIHR, or the Department of Health. (4) Ministerio de Economía y Competitividad SAF 2011 to 28375 and (5) European Cooperation in Science and Technology (COST) action TD1007 Bimodal PET-MRI molecular imaging technologies and applications for in vivo monitoring of disease and biological processes.

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Disclosures

None.

References


CLINICAL PERSPECTIVE

Despite the beneficial effects of percutaneous coronary intervention and stent implantation, these procedures may damage the vessel wall, particularly the endothelial layer, leading to increased vascular permeability and remodeling. Re-endothelialization of the vessel wall with functionally and structurally intact endothelial cells is crucial for minimizing vascular permeability and remodeling and maintaining the long-term patency of the treated vessel. Therefore, longitudinal measurements of both vascular permeability and remodeling in a single MRI examination provide a more comprehensive understanding of the focal responses of the vessel wall to injury. We propose that uptake of gadofosveset, a clinically approved albumin-binding contrast agent, into the vessel wall using $T_1$ mapping and DE-MRI protocols provides measurements of vascular permeability and remodeling. These biological processes could be used as surrogate markers to assess the healing response of the vessel wall in response to vascular injury.