Procedural and anatomical predictors of renal denervation efficacy using two radiofrequency renal denervation catheters in a porcine model

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INTRODUCTION

Hypertension is the most prevalent cardiovascular disease worldwide and is associated with poor cardiovascular outcome [1,2]. Despite the availability of multiple effective antihypertensive drugs, control rates remain unacceptably low [3]. Catheter-based renal denervation (RDN) is under investigation to disrupt renal sympathetic nerve activity and produce a reduction of blood pressure in patients with uncontrolled hypertension [4–7]. The recently published Symply HTN-OFF MED trial has proven the biological proof of principle for the blood pressure-lowering efficacy of RDN in the absence of antihypertensive medication [8]. The treatment effects, however, were variable and, in a relevant proportion of patients undergoing the procedure, blood pressure was not reduced sufficiently [9]. The neutral outcome of the SYMPLECTICITY HTN-3 study underscores the considerable heterogeneity in individual responses and the potential limitations of a poorly conceived ablation protocol. In the SYMPLECTICITY HTN-3 trial a mono-electrode catheter was utilized [10], whereas in the Symply HTN-OFF-MED trial, a revised procedural approach using a multielectrode catheter was executed [8]. Previous studies in the porcine model have shown that radiofrequency energy delivered to the branches and the main renal artery resulted in less variability and greater...
effect on norepinephrine (NEPI) reduction and axonal degeneration when compared with conventional ablation in the main artery alone [11,12]. A detailed understanding of the effect of procedural parameters and different ablative strategies and their outcomes is mandatory for further procedural and catheter refinements. The present porcine study aimed at investigating different procedural parameters and their value in predicting treatment efficacy following RDN with two different radiofrequency devices.

METHODS AND MATERIAL

Study design
A cohort of 13 female Yorkshire cross swine was approved for use by the Institutional Animal Care and Use Committee (IACUC) of the test facility, which is accredited by the Association for Assessment and Accreditation of Laboratory Animal Care (AAALAC). All procedures for this study were performed at Synchrony Labs Inc (Durham, North Carolina, USA) in compliance with USDA Regulations and the Animal Welfare Act (9 CFR Parts 1–3). The Guide for the Care and Use of Laboratory Animals was followed. Study animals were acclimated for 10 days prior to intervention. The animals were assembled in three groups. Group SPY (n = 5, SP1–SP5) and Group IBB (n = 5, IB1–IB5) underwent RDN procedures. Pigs were treated with radiofrequency energy in the proximal, mid, and/or distal regions of both main renal arteries as well as branch arteries measuring at least 3.0 mm in diameter and at least 10 mm in length. The number of ablations was determined based on the length and diameters of the arterial segments. Sham group (n = 3, Sham1–Sham3) served as sham controls for the purposes of obtaining NEPI in animals, which did not undergo radiofrequency catheter placement or radiofrequency energy delivery. Table 1 displays the treatment matrix and procedural details.

Device overview
Two different RDN devices were used. The IberisBloom (Terumo, Tokyo, Japan) RDN system consists of a multielectrode catheter, with four electrodes arranged helically and its multichannel generator [13]. The Symplicity Spyral (Medtronic, Santa Rosa, California, USA) RDN system consists of a multielectrode catheter with four electrodes mounted approximately 5 mm apart at 90° of separation from each other in a helical pattern, thus providing automated four-quadrant ablation treatments. Figure 1 visualizes the two systems.

Catheter-based renal sympathetic denervation
All animals were treated with acetylsalicylic acid (325 mg per os) and clopidogrel (75 mg per os) 24 h prior to the procedure and administered daily until the day of euthanasia. Animals were anesthetized with tiletamine (4 mg/kg intramuscularly) followed by isoflurane anaesthesia by

![FIGURE 1](a) Symplicity Spyral (SPY) RDN catheter. (b) IberisBloom (IBB) RDN catheter. [Pictures: (a) www.medtronicrdn.com; (b) http://www.terumomedical.com.]

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**TABLE 1. Treatment matrix and procedural details**

| Group     | Animal ID | Left Branch | Main | Total | Right Branch | Main | Total | Total/animal | Mean ± SD | Left | Right | Mean ± SD |
|-----------|-----------|-------------|------|-------|--------------|------|-------|--------------|-----------|------|-------|-----------|
| Group SPY| SP1       | 14          | 5    | 19    | 11           | 9    | 20    | 39           | 33.2 ± 3.3 | 2.8  | 1.2  | 1.4 ± 0.7 |
|           | SP2       | 7           | 6    | 13    | 8            | 10   | 18    | 31           | 1.2 ± 0.8  |      |       |           |
|           | SP3       | 8           | 4    | 12    | 10           | 9    | 19    | 31           | 2.0 ± 1.1  |      |       |           |
|           | SP4       | 10          | 5    | 15    | 8            | 9    | 17    | 32           | 2.0 ± 0.9  |      |       |           |
|           | SP5       | 8           | 9    | 17    | 6            | 10   | 16    | 33           | 0.9 ± 0.6  |      |       |           |
| Group IBB | IB1       | 7           | 6    | 13    | 7            | 10   | 17    | 30           | 26.2 ± 3.3 | 1.2  | 0.7  | 0.9 ± 0.5 |
|           | IB2       | 8           | 6    | 14    | 5            | 8    | 13    | 27           | 1.3 ± 0.6  |      |       |           |
|           | IB3       | 5           | 4    | 9     | 6            | 6    | 12    | 21           | 1.3 ± 1.0  |      |       |           |
|           | IB4       | 2           | 10   | 12    | 4            | 11   | 15    | 27           | 0.2 ± 0.4  |      |       |           |
|           | IB5       | 8           | 5    | 13    | 4            | 9    | 13    | 26           | 1.6 ± 0.4  |      |       |           |
| Sham group| Sham1     | No ablations|      |       |              |      |       |              |           |      |       |           |
|           | Sham2     | No ablations|      |       |              |      |       |              |           |      |       |           |
|           | Sham3     | No ablations|      |       |              |      |       |              |           |      |       |           |

Top: five animals were treated with the Symplicity Spyral system (group SPY). All animals of this group received 33.2 ± 3.3 ablations in main and branch arteries. The ratio of number of ablations (branch/main) was 1.4 ± 0.7. Middle: five animals were treated with the IberisBloom system (group IBB). All animals of this group received 26.2 ± 3.3 ablations in main and branch arteries. The ratio of number of ablations (branch/main) was 0.9 ± 0.5. Bottom: three animals received sham procedure (Sham Group) with zero ablations. IBB, IberisBloom; SPY, Symplicity Spyral.
mask to facilitate endotracheal intubation. Animals were maintained on isoflurane anaesthesia for the duration of the procedure and butorphanol (0.3 mg/kg i.m.) was administered for analgesia. Animals were prepared for aseptic surgery and vascular access was obtained percutaneously via the femoral artery. A 6 French introducer sheath was placed, and intravenous heparin was administered to achieve an activated clotting time of at least 250 s. The right and left renal arteries of the pig were engaged using a 6 French guide catheter (Medtronic PLC, Dublin, Ireland) and quantitative vascular analysis measurements were performed. Thereafter, a 6 French renal denervation catheter was delivered over an 0.014” guidewire (Abbott Vascular, Santa Clara, California, USA) introduced and positioned in the right and left renal artery via fluoroscopic guidance. Bilateral RDN was initiated within each renal artery starting in a distal position then repositioning more proximally for sequential treatments, overlap of treatments was avoided. All procedures were performed by the same interventionist. Meticulous adherence to the protocol with respect to four quadrant ablation and spacing between ablation sites was followed. All animals were survived for 7 days.

**Angiography**

Angiography was performed before and after RDN. Baseline angiography was done after placement of the guide catheter into the ostium of the intended artery. Arteriographic images of the vessel were obtained and target locations for denervation treatment were measured using quantitative vascular analysis (QVA). Terminal angiography of the treated renal arteries was performed using the same projection as during the initial procedure and QVA measurements were performed.

**Gross pathology and tissue collections**

Prior to sacrifice, renal angiography was performed to assess renal artery changes (e.g. stenosis, aneurysm) then a ventral midline laparotomy was performed to access both kidneys. Samples of renal cortical tissue were obtained and frozen in 1 g aliquots for NEPI measurements. The animals were then euthanized, and a gross pathology was performed in order to evaluate macroscopic changes to the kidneys and surrounding musculature. Kidney tissue norepinephrine concentration

All tissue specimens harvested for NEPI analysis in this study were obtained from the renal cortex. A total of six sections per kidney of renal cortical tissue were harvested. Three samples from cranial, mid, and caudal were obtained from the dorsal aspect of the kidney and three samples from cranial, mid, and caudal from the ventral aspect of the kidney. Each tissue sample was approximately 2–3 mm thick and weighed 0.5–1.0 g. Samples were flash frozen before being sorted at −80 °C. For evaluation a HPLC-MS ASSAY was used.

**Statistical analyses**

Data are presented as mean ± SD. Quantitative vessel analysis data were compared by unpaired *t*-test between Group SPY and Group IBB. NEPI renal tissue concentration of each group was compared by Tukey’s multiple comparison test. To assess main effects of arterial segments and devices and interactive effects, histomorphometric data were analysed by two-way analysis of variance (ANOVA). Pearson’s correlation coefficients were calculated to investigate the correlation among NEPI reduction and histopathological parameters. To investigate effects of number of ablations normalized by arterial length on variation of renal tissue NEPI reduction, means and SDs of NEPI reduction were calculated in kidneys whose number of ablations per millimeter length renal artery was more and less than median, respectively, and compared by unpaired *t*-test. All statistical analyses were performed with GraphPad Prism 7 (GraphPad Software, San Diego, California, USA). A *P* value greater than 0.05 was considered to indicate statistical significance.

**RESULTS**

**Angiography**

Late lumen loss and diameter stenosis are depicted in Table 2 and exhibited no significant difference between either of the treatment groups (baseline mean lumen of main and branch arteries: SPY: 4.63 ± 1.2 mm, IBB: 4.48 ± 1.1 mm, *P* = 0.60; late lumen loss of main and branch arteries: SPY: 0.29 ± 0.4 mm, IBB: 0.30 ± 0.4 mm, *P* = 0.90; diameter stenosis of main and branch arteries: SPY: 6.6 ± 10.9%, IBB: 7.7 ± 9.1%, *P* = 0.67).

**Gross pathology and histopathology**

There were no gross pathology changes noted in any of the animals treated with RDN. Lesion depth and lesion area
were comparable in both treatment arms (Table 3). Lesion depth was the greatest in main arteries compared with branch artery in both treatment groups (SPY: 6.26 ± 1.62 mm in main and 3.49 ± 1.06 mm in branch; IBB: 5.93 ± 1.77 mm in main and 3.26 ± 1.15 mm in branch; intergroup main vs. branch, \( P < 0.001 \); intergroup SPY vs. IBB, \( P = 0.54 \)). Lesion area between main or branch ablations was not different (SPY: 43.5 ± 29.5 mm\(^2\) in main and 45.0 ± 38.0 mm\(^2\) in branch; IBB: 52.3 ± 34.8 mm\(^2\) in main and 44.0 ± 42.6 mm\(^2\) in branch; intergroup main vs. branch, \( P = 0.77 \); intergroup SPY vs. IBB, \( P = 0.73 \)). The luminal surface of the renal artery was almost completely endothelialized by day 7, regardless of the device used or the number of ablations. Effects on adjacent structures, including focal psoas muscle necrosis, injury to the ureter, and focal necrosis of lymph node were observed with both devices. Ablation of anterior and posterior branch arteries showed more necrosis of kidney tissue. Localized necrosis of ureter was prevalent in both segments. Overall collateral changes did not pose a safety concern because of their low severity and limited extension. The renal veins were not affected (Fig. 2). Increasing the number of treatments along the length of the artery positively correlated with an increase in the number of nerves directly affected (Pearson’s correlation coefficient = 0.67, \( P = 0.001 \)).

**Kidney tissue norepinephrine concentration**

Treatment with RDN (SPY and IBB) resulted in a significant reduction in kidney tissue NEPI concentration compared with the Sham group, whereas there was no significant difference between the treatment groups (SPY vs. Sham, \(-95 ± 3\%\), \( P < 0.001 \); IBB vs. Sham, \(-88 ± 11\%\), \( P < 0.001 \); SPY vs. IBB, \( P = 0.33 \); Fig. 3). Interestingly, a more intensive ablation strategy led to a higher NEPI reduction with less variation. In pigs treated with less or more than the median of 0.2 ablations per 1 mm artery length, NEPI reduction was reduced by 88.5 ± 10.9 and 94.9 ± 3.3%, respectively. When assessing the effectiveness of the procedure in the main compared with the branch arteries separately, less ablations were needed per 1 mm arterial length in the branches. The median number in the main renal arteries was 0.26 ablations per 1 mm (NEPI reduction in <0.26 ablations per 1 mm: 88.4 ± 10.8% vs. >0.26 ablations per 1 mm: 95.0 ± 3.3%, \( P = 0.08 \)) compared with a median number of 0.16 ablations per 1 mm in branch renal arteries (NEPI reduction in <0.16 ablations per 1 mm: 89.8 ± 10.4% vs. >0.16 ablations per 1 mm: 93.6 ± 6.0%, \( P < 0.001 \); IBB vs. Sham, \( P < 0.001 \); SPY vs. IBB, \( P = 0.08 \); Fig. 4), indicating that ablations in branch arteries were associated with higher efficacy and less variability compared with main arteries.

**DISCUSSION**

The present study explored procedural and anatomical predictors of RDN efficacy using two different radiofrequency catheters in a porcine model. Both systems were highly effective in reducing NEPI kidney tissue content with no significant difference between them. Further, lesion depth and lesion areas were comparable in both groups. Lesion depth was highest in main arteries compared with branch arteries, which is in line with previous reports [12]. There was no clear dose–response relationship with number of ablations and NEPI reduction established. However, histological evaluation demonstrated that ablations in the branches were associated with higher efficacy based on nerve necrosis and distal nerve atrophy and less variability compared with main arteries.

Catheter-based RDN is currently under investigation for treatment of hypertension [4,14,15]. Early studies in patients with resistant hypertension have shown mixed results [10,16], whereas the recently published Spyral HTN-OFF MED study elegantly provided the biological proof of principle for the blood pressure-lowering efficacy of

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**TABLE 2. Angiographic changes following catheter-based renal denervation**

|                | Group SPY |                | Group IBB |                | \( P \) values of two-way ANOVA |
|----------------|-----------|----------------|-----------|----------------|------------------------------|
|                | \( N \)    | Baseline mean lumen diameter (mm) | Late lumen loss (mm) | Diameter stenosis (%) | Main vs. branch arteries | Group SPY vs. Group IBB | Intergroup effect |
|----------------|-----------|-----------------------------|----------------|-------------------|----------------|----------------|----------------|
| Group SPY      |           |                            |                |                   |                         |                          |                |
| All            | 32        | 4.63 ± 1.21                | 0.29 ± 0.42    | 6.6 ± 10.9        | >0.001           | 0.54            | 0.92           |
| Main           | 10        | 6.26 ± 0.43                | 0.37 ± 0.42    | 6.2 ± 7.2         | >0.001           | 0.73            | 0.67           |
| Branch         | 22        | 3.90 ± 0.51                | 0.25 ± 0.42    | 6.7 ± 12.4        | >0.001           | 0.71            | 0.96           |
| Group IBB      |           |                            |                |                   |                         |                          |                |
| All            | 28        | 4.48 ± 1.08                | 0.30 ± 0.39    | 7.7 ± 9.1         | >0.001           | 0.33            | 0.68           |
| Main           | 10        | 5.74 ± 0.27                | 0.18 ± 0.46    | 3.3 ± 8.5         | >0.001           | 0.33            | 0.68           |
| Branch         | 18        | 3.77 ± 0.58                | 0.37 ± 0.34    | 10.2 ± 8.7        | >0.001           | 0.33            | 0.68           |

Comparable lesion depth in lesion area in both arms. Lesion depth is significantly greater in main than in a branch. Data are presented as mean ± SD. IBB, IberisBloom; SPY, Symplicity Spyral.
RDN in the absence of antihypertensive medication [8]. Although several confounding factors have been identified and addressed in ongoing studies, such as the exclusion of isolated systolic hypertensive patients and revised procedural protocols with ablations in the main and branch arteries, the blood pressure-lowering effectiveness was variable [16–19]. Identification of patients not responding to the RDN therapy remains a major objective of future studies in the field.

One reason for nonresponse to the procedure may, in part, relate to an ineffective targeting of renal afferent and efferent sympathetic nerves, which are located in the adventitia of renal arteries. Disruption of these structures reduces the sympathetic nerve innervation of the kidney and through alteration of afferent nerve signalling whole body sympathetic activity [20]. The distribution pattern and density of sympathetic nerves along the renal arteries have been identified to impact the success of RDN [21]. Human histological studies as well as preclinical animal studies indicate that the maximal number of nerves are located around the proximal and middle segments of the renal artery, whereas lower number of nerves are located around distal structures. However, the mean distance of the sympathetic afferent and efferent nerves to the lumen of the renal arteries is shorter in the distal arterial segments [21]. Even if the mean lesion depth, which is influenced by surrounding structures [12], is less pronounced in branch arteries, the effectiveness of radiofrequency RDN is enhanced in this area likely because of the proximity of the nerves to the treated artery. One of the major unresolved issue of the procedure is how to monitor treatment success intraprocedurally. As no feedback is provided to the interventionist during the procedure, targeting the wrong arterial segments and performing too few ablations per artery could result in a high variability of blood pressure reduction or even in a complete negative outcome. On the other hand, ablations that exceed the required threshold needed to reduce local sympathetic activity are unnecessarily exposing patients to prolonged procedures and potential long-term vascular safety risk. Intraprocedural guidance on the number of ablations that need to be performed is currently lacking, though urgently desired. Herein, numerically more ablations per length were needed in the main when compared with branches to increase

**FIGURE 2** Focal changes following radiofrequency renal denervation shown within dotted lines. Focal changes were documented in both treatment groups. Ablation of main arteries (A) led to focal affection of lymph node (LN) and of psoas muscle (M). Ablation of anterior and posterior branch arteries (AB, PB) shows more irritations of kidney tissue (KI). Inflammation of ureter (U) was prevalent in both segments. The renal veins (V) have not been affected.

**FIGURE 3** Relative change in kidney norepinephrine concentration. Data are presented as mean ± SD.
In conclusion, the two devices evaluated in this study demonstrated similar histopathological effects (lesion depth, lesion area, nerve necrosis, nerve atrophy) and similar reduction of renal cortical NEPI levels. Ablations performed in the branches improved efficacy and reduced variability in treatment effects. This confirms the importance of anatomic and procedural parameters when performing RDN and deserves further investigation in clinical studies.

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Conflicts of interest

A.S. is an employee of Terumo Corporation. K.K. reports grants from Medtronic Japan Co. Ltd.; honoraria from Terumo Corp. and Otsuka Pharmaceutical Co. Ltd. during the conduct of the study. M.B. reports Speaker’s fees and scientific advice honoraria to Medtronic, Abbot Vascular, Servier, Novartis, Boehringer-Ingelheim. F.M. is supported by Deutsche Hochdruckliga and Deutsche Gesellschaft für Kardiologie and has received speaker honoraria and consultancy fees from Medtronic and Recor.

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