

# Ablation in Atrial Fibrillation with Ventricular Pacing Results in Similar Catheter Stability and Arrhythmia Recurrence Compared to Ablation in Sinus Rhythm with Atrial Pacing

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## Abstract

**Background:** Improved catheter stability is associated with decreased arrhythmia recurrence after atrial fibrillation (AF) ablation. Recently, atrial voltage mapping in AF was demonstrated to correlate better with scar as compared to mapping in sinus rhythm (SR). However, it is unknown whether ablation of persistent AF in sinus rhythm with atrial pacing or in atrial fibrillation with ventricular pacing results in differences in catheter stability or arrhythmia recurrence. **Methods:** We analyzed 53 consecutive patients undergoing first-time persistent AF ablation with pulmonary vein and posterior wall isolation: 27 were cardioverted, mapped, and ablated in sinus rhythm with atrial pacing, and 26 were mapped and ablated in AF with ventricular pacing. Ablation data was extracted from the mapping system and analyzed using custom MATLAB software to determine high-frequency (60Hz) catheter excursion as a novel metric for catheter spatial stability. **Results:** There was no difference in catheter stability as assessed by maximal catheter excursion, mean catheter excursion, or contact force variability between the atrial-paced and ventricular-paced patients. Ventricular-paced patients did have significantly greater mean contact forces compared to atrial-paced patients. One year arrhythmia-free survival was similar between the atrial paced and ventricular paced patients (78% vs 67%,  $p = 0.31$ ). **Conclusion:** For patients with persistent AF, ablation in AF with ventricular pacing results in similar catheter stability and arrhythmia recurrence as compared to cardioversion and ablation in sinus rhythm with atrial pacing. Given the improved fidelity of mapping in AF, mapping and ablating during AF with ventricular pacing may be preferred.

## Background

Radiofrequency (RF) ablation is a widely accepted and effective therapy in the management of atrial fibrillation (AF)<sup>1-3</sup>. However, ablation of persistent atrial fibrillation remains a therapeutic challenge, with recurrence rates of 30-60% at one year, and a frequent need for repeat ablation.<sup>4-7</sup> Improved catheter stability results in more consistent catheter-tissue contact, both allowing for more effective transmural lesion formation and preventing excessive force that could result in cardiac perforation. Furthermore, improved catheter stability has been shown to be associated with decreased arrhythmia recurrence following AF ablation.<sup>8</sup> Multiple strategies and techniques exist for improving catheter stability, including high frequency jet ventilation<sup>9-11</sup>, steerable catheter sheaths<sup>11-13</sup>, and rapid atrial pacing<sup>14</sup>. However, catheter stability may be affected by the underlying atrial rhythm. In patients with persistent AF, achieving stable sinus rhythm and reliable atrial pacing may be challenging prior to ablation, and pacing can only be performed in the ventricle. It is unknown

whether ablation in AF with ventricular pacing versus in sinus rhythm (SR) with atrial pacing has any effects on catheter stability, lesion quality, or clinical outcomes. In the present study, we sought to compare the ablation characteristics and clinical outcomes between patients with persistent AF who were mapped and ablated in sinus rhythm with atrial pacing compared to atrial fibrillation with ventricular pacing.

## Methods

### Patient population

We identified 53 consecutive patients at a single experienced electrophysiology center who underwent first time RF ablation for persistent AF. Of these, 27 were cardioverted then mapped and ablated in sinus rhythm with atrial pacing, and 26 were mapped and ablated in AF with ventricular pacing. Pacing was performed at a cycle length of 500-600ms, regardless of pacing site. All procedures were performed under general anesthesia with high frequency jet ventilation. Electroanatomical mapping was performed using either the circular Lasso<sup>®</sup> or five-spine PentaRay<sup>®</sup> mapping catheter and the CARTO3<sup>®</sup> mapping system, version 4 (Biosense Webster Inc., USA). Radiofrequency ablation was performed using the ThermoCool SmartTouch<sup>®</sup> force-sensing catheter (Biosense Webster Inc., USA) using point-by-point ablation at a power of 50W. All patients underwent pulmonary vein isolation (PVI) via wide antral circumferential ablation of the left and right pulmonary veins, as well as posterior wall isolation (PWI) via superior and inferior posterior wall lines connecting the PVI lesion sets. Additional ablation of the left and right carinas and/or residual electro-active areas within the posterior wall were performed at the discretion of the operator. VisiTag lesion stability settings were set to 2mm and 5s. Electrical isolation, including entrance and exit block, was confirmed using differential pacing and adenosine administration.

### Ablation analysis and catheter spatial stability

Our group had recently described a method for precise analysis of catheter-tip spatial stability, utilizing custom developed MATLAB script applied on X-Y-Z axis tip position data acquired at 60Hz<sup>8</sup>. Briefly, ablation data was extracted from CARTO<sup>®</sup> and analyzed using custom MATLAB script (Mathworks, USA). For each ablation lesion, the mean 3-dimensional (3-D) catheter position was calculated, and catheter excursion was defined as the distance between the position of the catheter tip, sampled at 60Hz, to the mean 3-D catheter position for the lesion. The mean and maximal catheter excursion were then calculated for each lesion (see Figure 1). Contact force variability was defined as the standard deviation of the contact force for the lesion.

### Ablation and clinical endpoints

The primary ablation endpoint was catheter stability, defined by mean and maximal catheter excursion. The secondary clinical endpoint was one year freedom from AF recurrence, following a 3-month blanking period. Recurrence was defined as an atrial arrhythmia lasting longer than 30 seconds on ambulatory monitoring, or an atrial arrhythmia documented on a standard 12 lead electrocardiogram. Recurrences within 3 months were included if they necessitated a repeat ablation.

### Statistics

All statistical analysis was performed using SPSS Statistics 26 (IBM Corp., USA) and graphs were constructed using Prism 8 (GraphPad Software Inc., USA). Continuous variables are expressed as mean  $\pm$  standard deviation, and categorical variables are expressed as percentages. Normality of data samples was assessed using Shapiro-Wilk test. Two sample hypothesis testing was performed using either Student's *t*-test if samples had normal distributions or Mann-Whitney *U* test if samples did not have normal distributions. Hypothesis testing for categorical variables was performed using Fisher's exact test. For Kaplan-Meier survival analysis, significance testing between groups was performed using log-rank test.

All patients provided written informed consent for their procedures, and all data collection and analysis was approved by the NYU Langone Medical Center Institutional Review Board.

## Results



Catheter ablation of persistent atrial fibrillation remains a therapeutic challenge, with high recurrence rates and frequent need for repeat ablations. The arrhythmogenic substrate in persistent atrial fibrillation is complex and incompletely elucidated, and although the pulmonary veins and posterior wall are frequently implicated, there is no consensus regarding optimal ablation targets or strategy<sup>15–19</sup>. Whereas the majority of ectopic foci in paroxysmal AF reside within the pulmonary vein-left atrial interface<sup>20,21</sup>, the mechanisms behind persistent AF are more complex and often the result of long-standing atrial remodeling<sup>22,23</sup>. Underlying atrial myopathy and fibrosis, particularly within the posterior wall of the left atrium, can serve as additional triggers and propagators of persistent AF. Voltage mapping can be performed to identify areas of scar and fibrosis, although previous data has shown that voltage amplitudes are affected by the underlying atrial rhythm<sup>24–26</sup>. A recent study by Qureshi et al. found that mapping during atrial fibrillation, compared to sinus rhythm, may actually be more sensitive and specific in identifying low voltage regions that correlate with atrial fibrosis on cardiac MRI<sup>27</sup>. More accurate intra-procedural voltage mapping would provide invaluable information regarding identifying arrhythmogenic substrate and refining ablation strategy.

Overall one year arrhythmia-free survival was 73%, which is in line with previous data for persistent AF ablation<sup>4,6,7</sup> and there was no significant difference in recurrence between the atrial-paced and ventricular-paced patients. The ventricular-paced patients had significantly greater LA diameters, time since AF diagnosis, as well as trend toward greater LA volume indices and lower LV ejection fraction. These measures have been shown to be associated with increased arrhythmia recurrence<sup>28–31</sup> thus sample size may have confounded recurrence outcomes.

## Limitations

This was a retrospective study of a consecutive cohort with a follow up time of 1 year. The small sample size was insufficient for logistic regression analysis to assess for predictors of arrhythmia recurrence. Patients ablated in AF with ventricular pacing may represent a subset with more complex substrate, evident by resistance to DCCV.

## Conclusions

Ablation of persistent atrial fibrillation in atrial fibrillation with ventricular pacing, compared to sinus rhythm with atrial pacing, results in similar catheter stability and lesion quality as assessed by impedance decline and ablation parameters. Given the recent evidence suggesting voltage mapping during atrial fibrillation may provide more accurate assessment of atrial fibrosis, it may be preferable, at least in patients presenting in AF, to ablate during atrial fibrillation with ventricular pacing, and defer cardioversion until after procedure completion.

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## Tables and Figures

Table 1. Baseline Clinical Characteristics

	Overall (n = 53)	A-paced (n = 27)	V-paced (n = 26)	p
Sex (% male)	74% (39/53)	74% (20/27)	73% (19/26)	1
Age (yrs)	63 ± 12	64 ± 10	62 ± 14	0.92
BMI (kg/m <sup>2</sup> )	30.7 ± 6.5	29.3 ± 4.6	32.1 ± 7.8	0.12
LVEF (%)	55 ± 12	57 ± 12	53 ± 13	0.11
LA Diameter (cm)	4.6 ± 0.7	4.4 ± 0.8	4.9 ± 0.6	0.02
LA Volume Index (mL/m <sup>2</sup> )	37 ± 13	33 ± 8	41 ± 15	0.14
Years Since AF Diagnosis	3.4 ± 4.5	2.5 ± 4.2	4.4 ± 4.7	0.02
CHA <sub>2</sub> DS <sub>2</sub> -VASc	2.1 ± 1.7	2.0 ± 1.6	2.2 ± 1.7	0.78
HTN	62% (33/53)	59% (16/27)	65% (17/26)	0.78
DM	15% (8/53)	19% (5/27)	12% (3/26)	0.7
CHF	21% (11/53)	22% (6/27)	19% (5/26)	1
CVA	6% (3/53)	4% (1/27)	8% (2/26)	0.61

	Overall (n = 53)	A-paced (n = 27)	V-paced (n = 26)	p
Vascular Disease	4% (2/53)	7% (2/27)	0% (0/26)	0.49
OSA	36% (19/53)	37% (10/27)	35% (9/26)	1
Post-Ablation Medications				
Anti-arrhythmic	87% (46/53)	81% (22/27)	92% (24/26)	0.42
BB or CCB	85% (45/53)	78% (21/27)	92% (24/26)	0.25
Post-Ablation Monitoring				
ILR, PPM, or ICD	11% (6/53)	11% (3/27)	12% (3/26)	1
Number of Monitors	2.3 ± 1.1	2.4 ± 1.0	2.1 ± 1.2	0.41

BMI = body mass index, LVEF = left ventricular ejection fraction, LA = left atrium, AF = atrial fibrillation, HTN = hypertension, DM = diabetes mellitus, CHF = congestive heart failure, CVA = cerebrovascular accident, OSA = obstructive sleep apnea, BB = beta-blocker, CCB = calcium channel blocker, ILR = implantable loop recorder, PPM = permanent pacemaker, ICD = implantable cardioverter-defibrillator

Table 2. Ablation Characteristics

	Overall (n = 53)	A-paced (n = 27)	V-paced (n = 26)	p
Count	146 ± 44	136 ± 37	156 ± 49	0.10
Lesion Duration (s)	7.4 ± 0.8	7.4 ± 0.7	7.4 ± 0.8	0.53
Average Force (g)	13.7 ± 2.2	13.0 ± 1.7	14.5 ± 2.4	0.02
Contact Force Variability (g)	4.1 ± 0.9	4.2 ± 1.0	4.1 ± 0.9	0.58
Max Power (W)	51.1 ± 1.0	51.2 ± .4	50.9 ± 1.4	0.22
Base Impedance (ohms)	120 ± 14	120 ± 12	120 ± 16	0.90
Impedance Decline (ohms)	7.4 ± 1.6	7.7 ± 1.5	7.1 ± 1.7	0.15
Percent Impedance Decline	6.0 ± 1.2%	6.3 ± 1.2%	5.8 ± 1.2%	0.12
FTI (gs)	101 ± 19	95 ± 15	107 ± 20	0.02
Mean Excursion (mm)	0.78 ± 0.12	0.79 ± 0.15	0.77 ± 0.10	0.62
Max Excursion (mm)	2.15 ± 0.40	2.14 ± 0.45	2.15 ± 0.35	0.94

Contact force variability was defined as the standard deviation of the contact force. FTI = force time integral

Table 3. Procedure Characteristics

	Overall (n = 53)	A-paced (n = 27)	V-paced (n = 26)	p
RF Time (min)	42 ± 13	41 ± 13	43 ± 13	0.64
Total Procedure Time (min)	160 ± 36	151 ± 36	169 ± 34	0.08
Number of DCCVs	1.0 ± 0.7	0.8 ± 0.6	1.2 ± 0.7	0.09?;?
2 DCCVs (%)	19% (10/53)	11% (3/27)	27% (7/26)	0.18

RF = radiofrequency, DCCV = direct current cardioversion

Figure 1. Figure 1. Example ablation lesion set with excursion measures. A) For each ablation lesion, catheter position was sampled at 60Hz. The orange dotted trail represents all sampled positions during a single RF lesion. Mean and maximal catheter excursion were calculated for each ablation lesion, and the radius of each sphere represents the mean catheter excursion for that lesion.

B) All patients underwent pulmonary vein isolation and posterior wall isolation. Full lesion set is projected with the left atrium removed.

Figure 2. Measures of catheter stability in atrial-paced and ventricular-paced patients. There was no significant difference in mean excursion (A), maximal excursion (B), or standard deviation of contact force (C) between the two groups. Ventricular paced patients had significantly greater average contact force, compared to atrial paced patients ( $p = 0.02$ ) (D).

Figure 3. One year arrhythmia free survival. There was no significant difference in one year arrhythmia free survival between the atrial-paced and ventricular-paced patients (78% vs 67%,  $p = 0.31$ ).

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