



Published in final edited form as:

J Cardiovasc Electrophysiol. 2014 May ; 25(5): 457–463. doi:10.1111/jce.12357.

Comparison of Left Atrial Area Marked Ablated in Electroanatomical Maps with Scar in MRI

Bhrihu R. Parmar, MD¹, Tyler R. Jarrett, BS¹, Nathan S. Burgon, BS¹, Eugene G. Kholmovski, PhD¹, Nazem W. Akoum, MD¹, Nan Hu, PhD², Rob S. MacLeod, PhD¹, Nassir F. Marrouche, MD¹, and Ravi Ranjan, MD PhD¹

¹CARMA Center, Division of Cardiology, University of Utah, Salt Lake City, UT

²Department of Internal Medicine, University of Utah, Salt Lake City, UT

Abstract

Background—Three-dimensional electroanatomic mapping (EAM) is routinely used to mark ablated areas during radiofrequency ablation. We hypothesized that, in atrial fibrillation (AF) ablation, EAM overestimates scar formation in the left atrium (LA) when compared to the scar seen on late-gadolinium enhancement magnetic resonance imaging (LGE-MRI).

Methods and Results—Of the 235 patients who underwent initial ablation for AF at our institution between August 2011 and December 2012, we retrospectively identified 70 patients who had preprocedural magnetic resonance angiography (MRA) merged with LA anatomy in EAM software and had a 3-month post-ablation LGE-MRI for assessment of scar. Ablated area was marked intraprocedurally using EAM software and quantified retrospectively. Scarred area was quantified in 3-month post-ablation LGE-MRI. The mean ablated area in EAM was $30.5 \pm 7.5\%$ of the LA endocardial surface and the mean scarred area in LGE-MRI was $13.9 \pm 5.9\%$ (p-value <0.001). This significant difference in the ablated area marked in the EAM and scar area in the LGE-MRI was present for each of the three independent operators. Complete pulmonary vein (PV) encirclement representing electrical isolation was observed in 87.8% of the PVs in EAM as compared to only 37.4% in LGE-MRI (p <0.001).

Conclusions—In AF ablation, EAM significantly overestimates the resultant scar as assessed with a follow-up LGE-MRI.

Keywords

atrial fibrillation; radiofrequency ablation; magnetic resonance imaging

For correspondence: Ravi Ranjan MD PhD Division of Cardiology, CARMA Center, 30 North 1900 East, Rm 4A100 University of Utah Salt Lake City, UT 84132 ravi.ranjan@hsc.utah.edu Fax: 801-581-7735 Phone: 801-213-2273.

Disclosures: Ravi Ranjan has been a consultant to Biosense Webster.

Ravi Ranjan: Concept/design, Funding secured by, Critical revision of article

R. Ranjan has been a consultant to Biosense Webster. E.G. Kholmovski participated on a research grant supported by Marrek Inc. N. Burgon is currently employed by Marrek Inc and holds stock options. N. Marrouche holds stock options in Marrek Inc. Other authors: No disclosures.

Introduction

Atrial fibrillation (AF) is increasingly managed by radiofrequency ablation, in which left atrial (LA) tissue is ablated around the pulmonary vein (PV) antra to create scar with a primary goal of electrically isolating the pulmonary veins^{1, 2}. Further ablation to the posterior wall, roof or septum varies from center-to-center and is largely operator dependent³. To guide ablation, preprocedural magnetic resonance angiography (MRA) or computerized tomography angiography is commonly acquired, segmented in an electroanatomic mapping (EAM) system, and merged with real-time LA geometry to create a 3D model of the LA. Areas of ablation are routinely marked on these merged maps to more accurately deliver contiguous lesion sets and improve procedural success⁴.

Late-gadolinium enhancement magnetic resonance imaging (LGE-MRI) is also increasingly used to visualize post-ablative scar in the LA⁵⁻⁷. Nonconductive scar takes some time to form fully after ablation and, as a result, a blanking period is required between the ablation procedure and accurate assessment of scar using LGE-MRI^{5, 8}. Several small studies have shown a correlation between the ablation-related LA scar seen on LGE-MRI and clinical outcomes^{9, 10}.

Peters et al. have shown the presence of scar at the ablation sites marked on EAM, yet studies comparing the EAM-recorded ablation tags with the eventual LA scar seen on LGE-MRI are limited¹¹. Although electrophysiologists depend heavily on the EAM to guide and track ablation, no study has evaluated the correlation between the ablated areas marked on the EAM during the procedure and the eventual scar area later seen in LGE-MRI.

The purpose of this study was to assess whether intraprocedural EAM ablated area and PV encirclement correlates with 3-month postprocedural LGE-MRI scar area and PV encirclement. We hypothesized that EAM ablated area significantly overestimates the post-procedural LGE-MRI scar in both total endocardial surface area and PV encirclement.

Methods

Patient Population

Two hundred and thirty-five patients underwent initial AF ablation at the University of Utah Hospital between August 2011 and December 2012. Of the 235 patients, 70 patients had a retrievable MRA LA models merged intraprocedurally with an EAM system and a 3-month post-ablation LGE-MRI. Study variables were collected after approval by the institutional Internal Review Board.

Ablation Procedure

All patients underwent PV antrum isolation with or without additional ablation by one of three cardiac electrophysiologists at the University of Utah Hospital. Preprocedural MRA was used to generate a LA shell for merge with the LA anatomy in the CARTO EAM system (Biosense Webster, Diamond Bar, CA, USA). Merging of the shell was accomplished by a landmark registration method using intracardiac echocardiography (ICE) and the merge quality was also verified during the procedure. A 3.5-mm irrigated tip

ablation catheter (Thermocool, Biosense Webster) and a circular 10-pole mapping catheter (Lasso, Biosense Webster) were advanced transseptally into the LA under ICE and fluoroscopic guidance. Radiofrequency energy was applied circumferentially to the PV antra for all patients with additional application to the posterior wall in a subset of patients. ICE was used to track the ablation catheter tip during the procedure to ensure good catheter tip tissue contact. After the initial ablation presence of local electrograms was checked by the lasso or ablation catheter and additional ablation was carried out if there was local electrical activity or the pulmonary vein was not isolated. Operator 1 used 50 W for LA ablations while repositioning the catheter approximately every 5 seconds. Operator 2 used 35 W with catheter repositioning approximately every 10 seconds for most of the ablation and decreased power to 30 W if close to the esophagus as identified with intracardiac echocardiography. Operator 3 used 25–30 W to ablate the posterior wall and 30–35 W for the anterior wall, moving the catheter every 10 to 12 seconds. Operator 3 also used a temperature probe in the esophagus to monitor the temperature.

During RF delivery, ablation tags with an institutional standard 2 mm radius were marked and projected onto the surface of the MRA shell approximately every 5 seconds for operator 1 and every 10 seconds for operators 2 and 3. The Lasso catheter was used to confirm the electrical isolation of PVs.

Quantification of Ablated Surface Area in EAM

Merged LA models with ablation tags from each procedure were retrieved retrospectively from the EAM software using the CARTO Merge Plus module (CARTO, Biosense Webster). Regions of the model surface covered by ablation tags were outlined by two blinded experts using the design line tool and surface areas of these regions were measured with the area measurement tool (Figure 1). Surface area distal to the PV ostia and the section representing the mitral valve were measured and excluded from the total surface area of the model to quantify LA wall area and reduce variability from PV segmentation. Gaps within ablated regions of 0.5 cm² or greater were also measured and excluded from regions with ablation tags to more accurately assess the ablated area. Ablated surface areas were then summed and expressed as a percentage of the LA wall area.

In a randomly selected subset of consecutive patients (n=10), ablation tag size was retrospectively reduced to 1 mm radius to assess for any discrepancy in EAM ablated area and the ablated area was re-calculated in EAM using the same process as for 2mm ablation tag size.

Quantification of Scarred Surface Area from LGE-MRI

All MRI studies were performed on 1.5 or 3 Tesla clinical MR scanners (Siemens Healthcare, Erlangen, Germany). Each LGE-MRI scan was acquired about 15 minutes after contrast agent injection (0.1 mmol/kg, Multihance, Bracco Diagnostic Inc., Princeton, NJ) using a 3D inversion recovery prepared, respiration navigated, electrocardiogram (ECG)-gated, gradient echo pulse sequence using an institutional standard protocol described previously¹². Typical acquisition parameters were: free-breathing using respiratory and ECG navigation, a transverse imaging volume with acquired voxel size = 1.25 × 1.25 × 2.5 mm

(reconstructed to $0.625 \times 0.625 \times 1.25$ mm). The other imaging parameters were optimized for respective field strength of scanner to improve post-ablation scar visibility and simultaneously keep scan duration acceptable for patients (< 15 minutes). For scans performed on 1.5 Tesla scanners, parameters were set as follows: repetition time = 5.4 ms, echo time = 2.3 ms, and flip angle = 20° . For scans performed on 3 Tesla scanner, parameters were: repetition time = 3.1 ms, echo time = 1.4 ms, flip angle = 14° . Depending on subject respiration pattern, typical scan time for the LGE-MRI study on 1.5 Tesla scanner was 8-12 minutes and at 3 Tesla scan time was 5-9 minutes.

Endocardial and epicardial contours were delineated in each slice and assembled in Corview software (Marrek Inc., Salt Lake City) to create a 3D model of the LA. The PVs and the mitral valve were excluded from the segmentation to isolate the LA wall, and the total area of the remaining LA endocardial surface was recorded. Scar was defined using a pixel-intensity threshold algorithm as described in previous reports^{9, 13, 14}. In brief, normal and scar tissue were distinguished based on a bimodal distribution of pixel intensities, and scar was defined at three standard deviations above the normal mean tissue pixel intensity. With determination of the scar areas, the surface area of the LA endocardial surface covered by enhancement was recorded. Scar area was expressed as a percent of total LA endocardial surface area.

Pulmonary Vein Encirclement

Two blinded observers evaluated patterns of ablation in EAM around the PV ostia by estimating the percent of ostial circumference covered by ablation tags in 10% increments. Inter-observational consensus was defined as the mean percent of circumference. A disagreement of 20 percent or greater between observers in circumferential encirclement was resolved with a third observation by a blinded electrophysiologist.

Three blinded observers evaluated scar patterns around the PV ostia in all LGE-MRI segmentations. Each observer estimated the percent of ostial circumference covered by enhancement in increments of 10% and these percentages were averaged to obtain observer consensus. If percent isolation varied by greater than or equal to 20 percent among the observers, the average of the two nearest estimates was used as observer consensus. Complete encirclement was defined as greater than 90 % of circumferential coverage of the PV antra by enhancement or ablation tags in both EAM and LGE-MRI.

Statistical Analysis

Paired Student's *t*-test was performed to compare the ablated area in EAM with LGE-MRI scar area as well as percent encirclement of the individual PVs. McNemar's test was applied to compare the number of PVs completely isolated by EAM ablation tags and by LGE-MRI scar. Left or right common PVs were excluded from this analysis. Two-sided p-values of less than 0.05 were deemed significant.

Results

Baseline characteristics of the study population are presented in Table 1. The average age was 65 years and 76% were men. Of the 70 patients, 34 had paroxysmal AF and 36 had

persistent AF. Radiofrequency energy was applied circumferentially to the PV antra for all patients (n=70) with additional application to the posterior wall (n=38, 54.3%). The mean ablated area as a percent of LA surface area in EAM and the standard deviation was $30.5 \pm 7.5\%$, and the corresponding mean scar area along with standard deviation in LGE-MRI was $13.9 \pm 5.9\%$. The difference between these two means was statistically significant (p-value <0.001 , Figure 2). Figure 3 shows two pairs of EAM and LGE-MRI, one showing a strong correlation between the ablation tags in EAM and the scar from LGE-MRI and the other showing a large discrepancy. The mean total LA surface area and standard deviation were $155.1 \pm 36.4 \text{ cm}^2$ in the EAM and $164.5 \pm 38.7 \text{ cm}^2$ in the LGE-MRI.

Ablation tag encirclement of the PVs in EAM did not consistently predict scar encirclement in LGE-MRI. While the mean percent of ostial circumference covered by ablation tags in EAM was $>90\%$ for all PVs, the corresponding mean in LGE-MRI varied between 70-73% with the exception of the left inferior vein that had a mean of 85% (Figure 4; p-value <0.001). Complete encirclement was significantly more common in EAM compared to LGE-MRI for all four PVs (Figure 5; p-value <0.001). Collectively, 87.8% of all PVs were encircled in EAM compared to 37.4% in LGE-MRI (p-value <0.001). Left superior PVs were completely encircled in 74% of the patients in EAM compared to 36% in the LGE-MRI, left inferior PVs were encircled in 86% of the patients in EAM compared to 52% in LGE-MRI, right superior PVs were encircled in 92.6% of the patients in EAM compared to 33.8% in LGE-MRI, and right inferior PVs were encircled in 98.5% of the patients in EAM compared to 27.9% in LGE-MRI (all p-values <0.001 , Figure 5).

The number of pulmonary veins isolated in any patient was also much higher in EAM compared to LGE-MRI. All four PVs were marked isolated in 47% of the patients in EAM compared to only 8.5% based on scar in LGE-MRI. Three veins were isolated in 30% in EAM and 11.4% in LGE-MRI. Two veins were isolated in 21.4% in EAM compared to 12.9% on LGE-MRI. Only one vein was isolated in 1.4% of patients in EAM compared to 37% in LGE-MRI and finally there were no patients with none of the veins marked isolated in EAM compared to 30% of patients in LGE-MRI.

In the subset of patients (n=10), the ablation area was also measured using reduced ablation point tag size of 1mm radius in EAM. The percent of LA wall area covered by 1 mm tags was $21.2 \pm 5.2\%$ and the percent of LA wall area covered by 2 mm tags in the same group of patients was $26.5 \pm 8\%$. The percent of LA area covered by scar in this group of patients based on LGE-MRI was $12.0 \pm 5.1\%$. The mean ablated area in the EAM when using 1 mm radius tags was diminished compared to 2 mm radius tags, but the disparities between the ablated area in EAM and scar area seen in LGE-MRI remained significant (1 mm: p <0.001 ; 2 mm: p <0.001).

The differences between EAM ablation area and scar area on LGE-MRI persisted among all three operators (Figure 6). Operators 1 and 2 had similar mean ablation areas in EAM of 31-32% and mean scar areas in LGE-MRI of 14%, while operator 3 had a lower mean ablation area in EAM of 25.1% and subsequently lower mean scar area on LGE-MRI of 10%. All differences between EAM and LGE-MRI were significant (p-values <0.01). The

total ablation time for all three operators was similar at 2146 seconds for operator 1, 2748 seconds for operator 2 and 2957 seconds for operator 3.

Discussion

This study sought to evaluate correlations between the ablated areas marked in EAM during the procedure with resultant scar areas as determined by follow-up LGE-MRI. One prior study has shown a relationship between the intended sites of ablation and LGE-MRI scar in 19 patients, and did not find scar at approximately 20% of the marked ablation sites upon follow-up¹¹. This study elaborates upon that result and demonstrates that the intended area of ablation as marked on EAM significantly overestimates the actual scar on LGE-MRI across multiple operators and ablation tag sizes. Similarly, ablation tags on EAM that indicate complete or nearly complete encirclement of PV ostia by ablation correlate inconsistently with encirclement by scar in LGE-MRI. Quantitatively, the percent of PV encirclement with ablation tags was also overestimated in EAM when compared to the percent PV encirclement with scar in LGE-MRI (Figure 4).

Our study also showed that this tendency of EAM to overestimate scar formation in LGE-MRI is not operator dependent but was found across three operators using different ablation techniques with varying combinations of power and catheter movement. Despite these different techniques, the difference between ablation area in EAM and scar area on LGE-MRI was significant for all the three operators.

The observed differences between EAM ablated area and LGE-MRI scar could be explained in several ways: lack of proper contact between the LA tissue and the ablation catheter, inadequate time spent at the ablation site (or the frequency at which the tags are placed on the EAM), formation of transient edema at the ablation site that recovers over time, or the size of the tags marking ablation being much bigger than the area ablated.

The use of tags with 2 mm radius is the current standard across most centers and reasonably assumes the size of lesions created using catheters with a 3.5 mm tip¹⁵. Prior work looking at ablation lesions with similar size catheter tip has shown that even 5 Watts of power can create 5 mm size ventricular lesions and significantly larger lesions at higher power¹⁶. In this study reducing the size of tags to mark the ablated area did reduce the calculated ablation area in the EAM but did not resolve the difference in area between EAM and LGE-MRI. Moreover, the size of ablation tags was reduced retrospectively during the time of the analysis, creating additional gaps in the ablation that contributed to decreased ablated area. Given that most operators prefer to create contiguous lesion sets in PV isolation, it is likely that use of 1 mm tags during the ablation procedure itself would result in these gaps being filled and an ablated area closer to that marked with 2 mm radius tags.

Adjusting ablation tag acquisition parameters based on catheter stability, wattage, temperature or impedance change might also be effective in mapping ablated areas that will correlate more consistently with resultant scar areas. But as one gets more aggressive with ablation, one has to balance the increased risk of complications with more force or increased ablation time. In addition to the catheter related features that impact ablation tag acquisition,

the frequency with which tags are placed can make a big difference in the area deemed to be ablated. However, finding optimal frequency of tag placement is challenging given the large number of variables that factor into good lesion formation. In this study we used the clinical maps created at the time of the ablation with the operator marking the spots deemed to be ablated based on well accepted current clinical practice including elimination of local electrograms. Tagging areas as ablated after eliminating local electrograms is the current clinical practice but clearly seems to overestimate the area that will result in scar. A direct feedback of tissue changes as ablation is carried out can potentially help in narrowing this discrepancy.

The use of force sensing catheters can provide some indication of tip-to-tissue contact, but the lack of any direct feedback of tissue changes will persist as a central problem for electrophysiologists^{17, 18}. Furthermore, transmuralty of lesions is dependent not only on good tissue contact but also on energy delivered, duration of the energy delivered, the catheter contact force, catheter tip area in contact with the tissue, and irrigated versus non-irrigated catheter tips. Given that acute electrical isolation of PVs confirmed with the Lasso catheter offers some evidence of encirclement of the PVs by lesions, the observed differences may stem from gaps in lesions sets with edema that contribute to acute isolation of the veins but recover over time. The possibility of acute isolation despite the presence of gaps in lesion sets has been demonstrated both computationally and experimentally¹⁹.

MRI offers a promising avenue for resolving this discrepancy. MRI has excellent soft tissue characterization ability and has been used to acutely identify ablation related tissue changes^{15, 16}. Using MRI to identify gaps in ablation lesions sets and target them is one potential way to bridge this gap between the marked area in EAM believed to be ablated and true scar as seen in LGE-MRI^{20, 21}. Even though a real-time MRI system seems to be a most promising method for direct visualization of changes to the tissue from ablation, limitations remain, and accurate determination of long-term scar from scans done acutely is still an area of active investigation.

Further work is needed to elucidate the observed discrepancies between EAM ablation and LGE-MRI scar. Prior studies have shown that smaller scar area on LGE-MRI correlates with higher AF recurrence^{9, 10}. For the first time, we show that large areas marked as ablated in the EAM do not result in long-term scar and could well be the reason for high recurrence rates of arrhythmias like AF. Further studies are needed to associate AF recurrence with the difference between EAM ablation area and LGE-MRI scar area to provide the necessary impetus to advocate for delivery of more effective lesions.

Conclusions

In AF ablation, EAM ablation mapping significantly overestimates scar formation when compared to LGE-MRI. Better visualization and ablation mapping techniques are necessary to achieve the desired LA scar.

Limitations

We acknowledge significant limitations to this study. This is a retrospective analysis of a single center, AF ablation population, although multiple operators with various ablation techniques were represented. Furthermore, variables derived from procedures may have significant inter- and intra-observer variability. To reduce inter-observer variability, we derived a consensus from two blinded independent experts for EAM and three blinded independent reviewers for LGE-MRI to document PV encirclement as well as ablated and scarred areas. Also, this study does not make an attempt to predict areas marked as ablated in EAM that will result in scar or heal over time and if this difference between areas deemed ablated versus scar affects procedural outcome. Correlating this difference between areas deemed ablated in EAM and resulting scar with procedural outcome is needed before we make any changes to our ablation approach.

Acknowledgments

The authors would like to thank Vissal Penn and Hannah Tredway for help with Corview and PV encirclement analysis.

Research reported in this publication was supported by the National Heart, Lung, And Blood Institute of the National Institutes of Health under Award Number K23HL115084 to Ravi Ranjan. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health.

This project was also supported by grants from the National Institute of General Medical Sciences (8 P41 GM103545-14) from the National Institutes of Health through the Center for Integrative Biomedical Computing (CIBC).

Bibliography

1. Haissaguerre M, Jais P, Shah DC, Takahashi A, Hocini M, Quiniou G, Garrigue S, Le Mouroux A, Le Metayer P, Clementy J. Spontaneous initiation of atrial fibrillation by ectopic beats originating in the pulmonary veins. *N Engl J Med.* 1998; 339:659–666. [PubMed: 9725923]
2. Pappone C, Rosanio S, Oreto G, Tocchi M, Gugliotta F, Vicedomini G, Salvati A, Dicandia C, Mazzone P, Santinelli V, Gulletta S, Chierchia S. Circumferential radiofrequency ablation of pulmonary vein ostia: A new anatomic approach for curing atrial fibrillation. *Circulation.* 2000; 102:2619–2628. [PubMed: 11085966]
3. Segerson NM, Daccarett M, Badger TJ, Shabaan A, Akoum N, Fish EN, Rao S, Burgon NS, Adjei-Poku Y, Kholmovski E, Vijayakumar S, DiBella EV, MacLeod RS, Marrouche NF. Magnetic resonance imaging-confirmed ablative debulking of the left atrial posterior wall and septum for treatment of persistent atrial fibrillation: Rationale and initial experience. *J Cardiovasc Electrophysiol.* 2010; 21:126–132. [PubMed: 19804549]
4. Malchano ZJ, Neuzil P, Cury RC, Holmvang G, Weichet J, Schmidt EJ, Ruskin JN, Reddy VY. Integration of cardiac ct/mr imaging with three-dimensional electroanatomical mapping to guide catheter manipulation in the left atrium: Implications for catheter ablation of atrial fibrillation. *J Cardiovasc Electrophysiol.* 2006; 17:1221–1229. [PubMed: 17074008]
5. Peters DC, Wylie JV, Hauser TH, Kissinger KV, Botnar RM, Essebag V, Josephson ME, Manning WJ. Detection of pulmonary vein and left atrial scar after catheter ablation with three-dimensional navigator-gated delayed enhancement mr imaging: Initial experience. *Radiology.* 2007; 243:690–695. [PubMed: 17517928]
6. Hunter RJ, Jones DA, Boubertakh R, Malcolm-Lawes LC, Kanagaratnam P, Juli CF, Davies DW, Peters NS, Baker V, Earley MJ, Sporton S, Davies LC, Westwood M, Petersen SE, Schilling RJ. Diagnostic accuracy of cardiac magnetic resonance imaging in the detection and characterization of

- left atrial catheter ablation lesions: A multicenter experience. *J Cardiovasc Electrophysiol.* 2013; 24:396–403. [PubMed: 23293924]
7. Badger TJ, Daccarett M, Akoum NW, Adjei-Poku YA, Burgon NS, Haslam TS, Kalvaitis S, Kuppahally S, Vergara G, McMullen L, Anderson PA, Kholmovski E, MacLeod RS, Marrouche NF. Evaluation of left atrial lesions after initial and repeat atrial fibrillation ablation: Lessons learned from delayed-enhancement mri in repeat ablation procedures. *Circ Arrhythm Electrophysiol.* 2010; 3:249–259. [PubMed: 20335558]
 8. McGann C, Kholmovski E, Blauer J, Vijayakumar S, Haslam T, Cates J, DiBella E, Burgon N, Wilson B, Alexander A, Prastawa M, Daccarett M, Vergara G, Akoum N, Parker D, MacLeod R, Marrouche N. Dark regions of no-reflow on late gadolinium enhancement magnetic resonance imaging result in scar formation after atrial fibrillation ablation. *J Am Coll Cardiol.* 2011; 58:177–185. [PubMed: 21718914]
 9. McGann CJ, Kholmovski EG, Oakes RS, Blauer JJ, Daccarett M, Segerson N, Airey KJ, Akoum N, Fish E, Badger TJ, DiBella EV, Parker D, MacLeod RS, Marrouche NF. New magnetic resonance imaging-based method for defining the extent of left atrial wall injury after the ablation of atrial fibrillation. *Journal of the American College of Cardiology.* 2008; 52:1263–1271. [PubMed: 18926331]
 10. Peters DC, Wylie JV, Hauser TH, Nezafat R, Han Y, Woo JJ, Taclas J, Kissinger KV, Goddu B, Josephson ME, Manning WJ. Recurrence of atrial fibrillation correlates with the extent of post-procedural late gadolinium enhancement: A pilot study. *JACC Cardiovasc Imaging.* 2009; 2:308–316. [PubMed: 19356576]
 11. Taclas JE, Nezafat R, Wylie JV, Josephson ME, Hsing J, Manning WJ, Peters DC. Relationship between intended sites of rf ablation and post-procedural scar in af patients, using late gadolinium enhancement cardiovascular magnetic resonance. *Heart Rhythm.* 2010; 7:489–496. [PubMed: 20122877]
 12. Vergara GR, Marrouche NF. Tailored management of atrial fibrillation using a lge-mri based model: From the clinic to the electrophysiology laboratory. *J Cardiovasc Electrophysiol.* 2010; 22:481–487. [PubMed: 21044212]
 13. Badger TJ, Oakes RS, Daccarett M, Burgon NS, Akoum N, Fish EN, Blauer JJ, Rao SN, Adjei-Poku Y, Kholmovski EG, Vijayakumar S, Di Bella EV, MacLeod RS, Marrouche NF. Temporal left atrial lesion formation after ablation of atrial fibrillation. *Heart Rhythm.* 2009; 6:161–168. [PubMed: 19187904]
 14. Ranjan R. Magnetic resonance imaging in clinical cardiac electrophysiology. *Critical reviews in biomedical engineering.* 2012; 40:409–426. [PubMed: 23339649]
 15. Dickfeld T, Kato R, Zviman M, Lai S, Meininger G, Lardo AC, Roguin A, Blumke D, Berger R, Calkins H, Halperin H. Characterization of radiofrequency ablation lesions with gadolinium-enhanced cardiovascular magnetic resonance imaging. *J Am Coll Cardiol.* 2006; 47:370–378. [PubMed: 16412863]
 16. Dickfeld T, Kato R, Zviman M, Nazarian S, Dong J, Ashikaga H, Lardo AC, Berger RD, Calkins H, Halperin H. Characterization of acute and subacute radiofrequency ablation lesions with nonenhanced magnetic resonance imaging. *Heart Rhythm.* 2007; 4:208–214. [PubMed: 17275759]
 17. Kuck KH, Reddy VY, Schmidt B, Natale A, Neuzil P, Saoudi N, Kautzner J, Herrera C, Hindricks G, Jais P, Nakagawa H, Lambert H, Shah DC. A novel radiofrequency ablation catheter using contact force sensing: Toccata study. *Heart Rhythm.* 2012; 9:18–23. [PubMed: 21872560]
 18. Reddy VY, Shah D, Kautzner J, Schmidt B, Saoudi N, Herrera C, Jais P, Hindricks G, Peichl P, Yulzari A, Lambert H, Neuzil P, Natale A, Kuck KH. The relationship between contact force and clinical outcome during radiofrequency catheter ablation of atrial fibrillation in the toccata study. *Heart Rhythm.* 2012; 9:1789–1795. [PubMed: 22820056]
 19. Ranjan R, Kato R, Zviman MM, Dickfeld TM, Roguin A, Berger RD, Tomaselli GF, Halperin HR. Gaps in the ablation line as a potential cause of recovery from electrical isolation and their visualization using mri. *Circ Arrhythm Electrophysiol.* 2011; 4:279–286. [PubMed: 21493875]
 20. Lardo AC, McVeigh ER, Jumrussirikul P, Berger RD, Calkins H, Lima J, Halperin HR. Visualization and temporal/spatial characterization of cardiac radiofrequency ablation lesions using magnetic resonance imaging. *Circulation.* 2000; 102:698–705. [PubMed: 10931812]

21. Ranjan R, Kholmovski EG, Blauer J, Vijayakumar S, Volland NA, Salama ME, Parker DL, MacLeod R, Marrouche NF. Identification and acute targeting of gaps in atrial ablation lesion sets using a real-time magnetic resonance imaging system. *Circ Arrhythm Electrophysiol.* 2012; 5:1130–1135. [PubMed: 23071143]

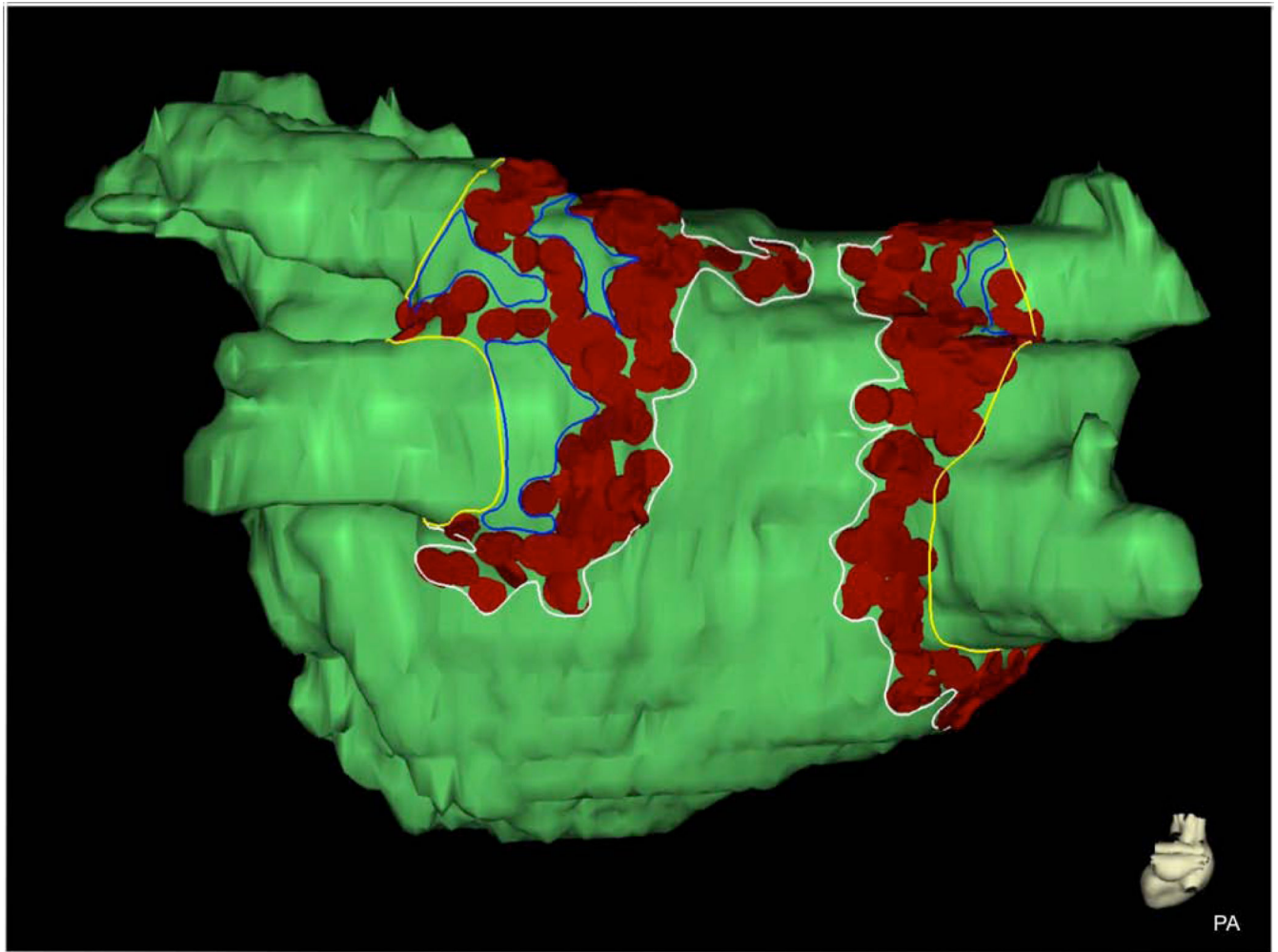


Figure 1. Measurement of ablated area in EAM. Ablation tags (red) are projected onto the surface of the LA model (green). Lines were drawn on the surface of the model to define the PV ostia (yellow), ablated regions (white), and gaps in ablated regions (blue). Gaps and PV surface area were measured and subtracted from surface area enclosed by white contours to derive ablated surface area. EAM: electroanatomic mapping; LA: left atrium; PV: pulmonary vein.

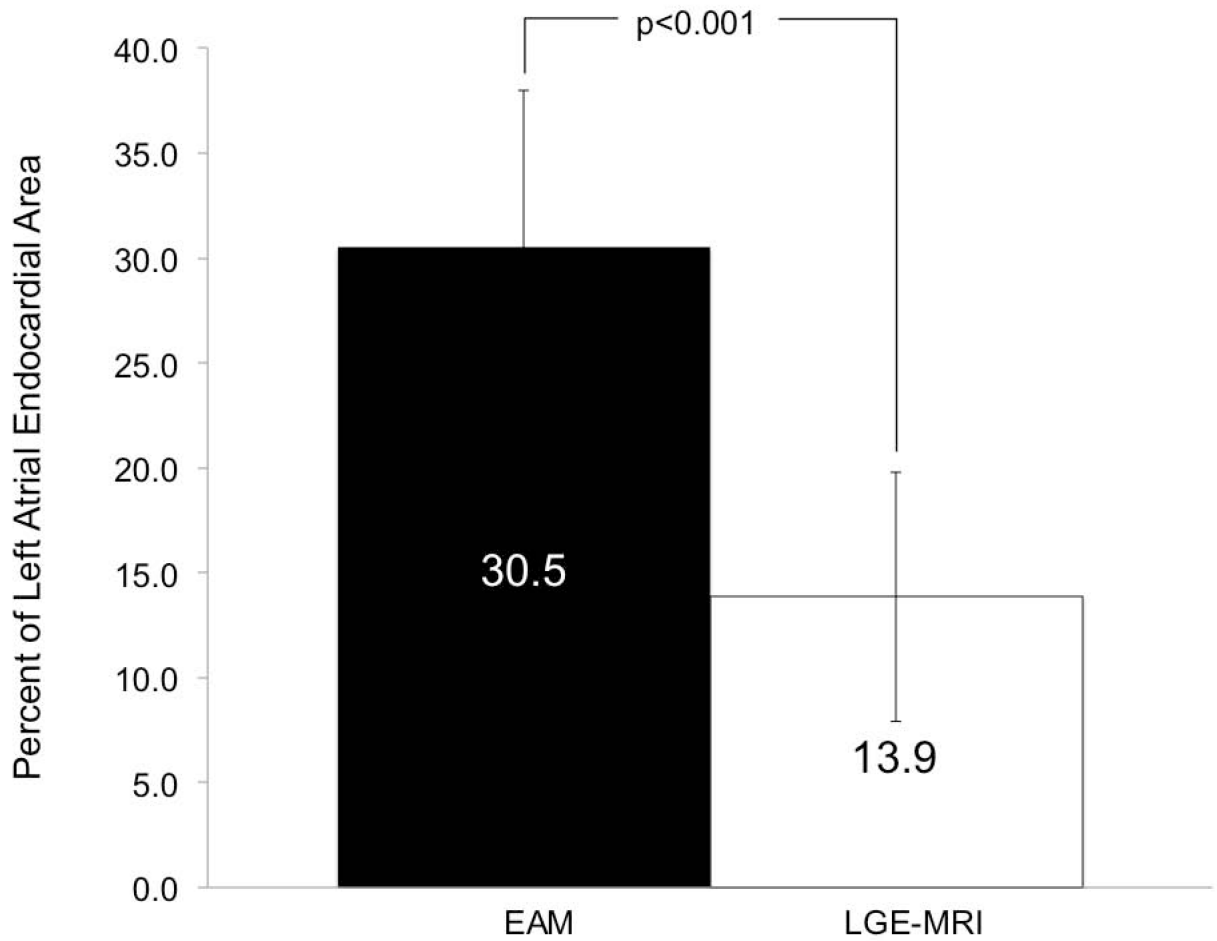


Figure 2. Comparison of the mean ablated area along with corresponding standard deviation in EAM and three-month LGE-MRI for all patient ($p < 0.001$). EAM: electroanatomic mapping; LGE-MRI: late-gadolinium enhancement magnetic resonance imaging.

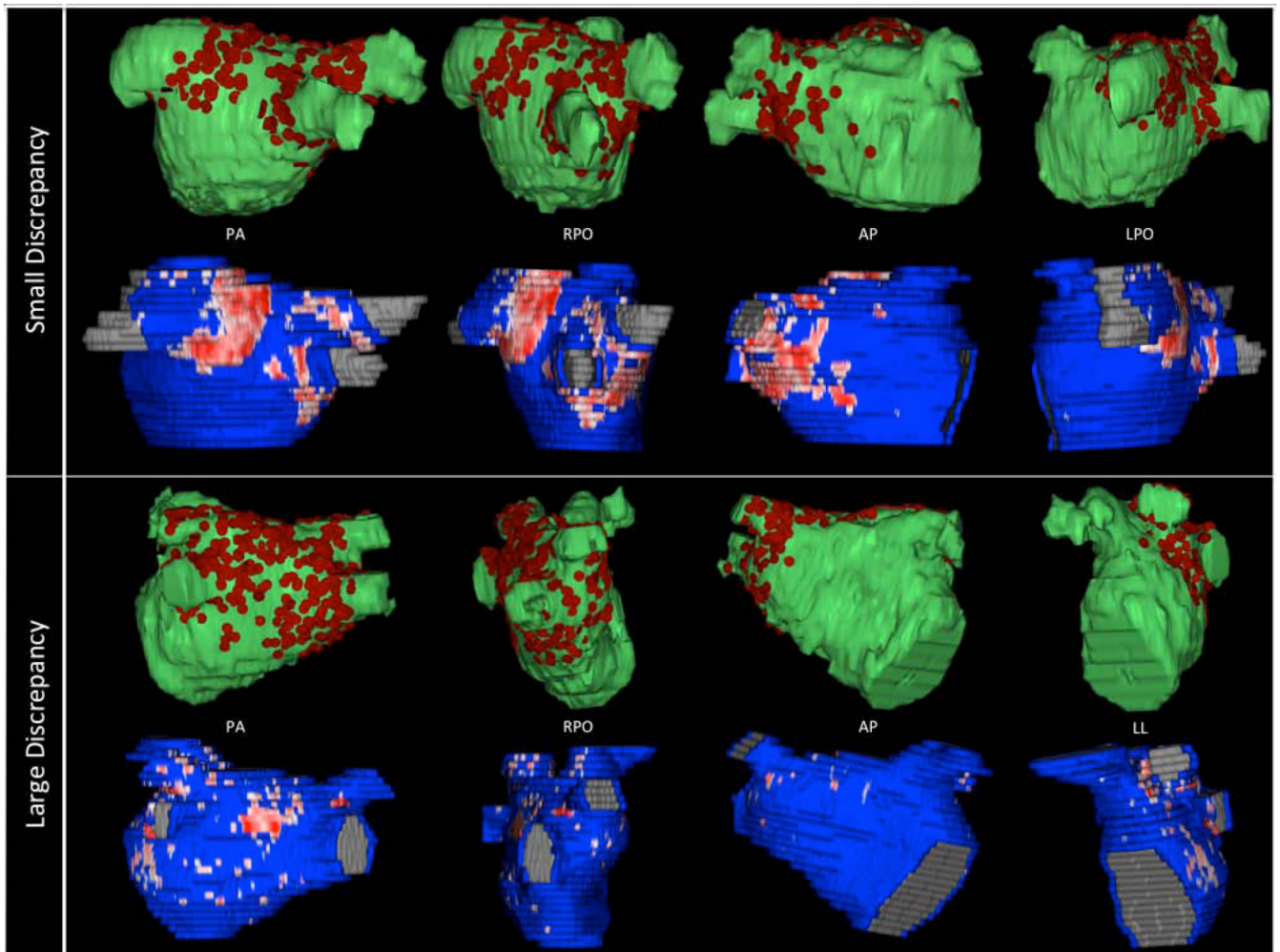


Figure 3.

Comparison of PVI in EAM and follow-up LGE-MRI in two patients. The upper rows show a close correlation between ablation tags marked in EAM (row 1) with corresponding scar in follow-up LGE-MRI (row 2). The difference between ablated and scarred areas was 5.3 %. Rows 3 and 4 demonstrate a much greater difference (30.6 %) between EAM and LGE-MRI areas. EAM: electroanatomic mapping; LGE-MRI: late-gadolinium enhancement magnetic resonance imaging; PVI: pulmonary vein isolation; AP: antero-posterior; PA: postero-anterior; RPO: right posterior oblique; LPO: Left posterior oblique, LL: left lateral.

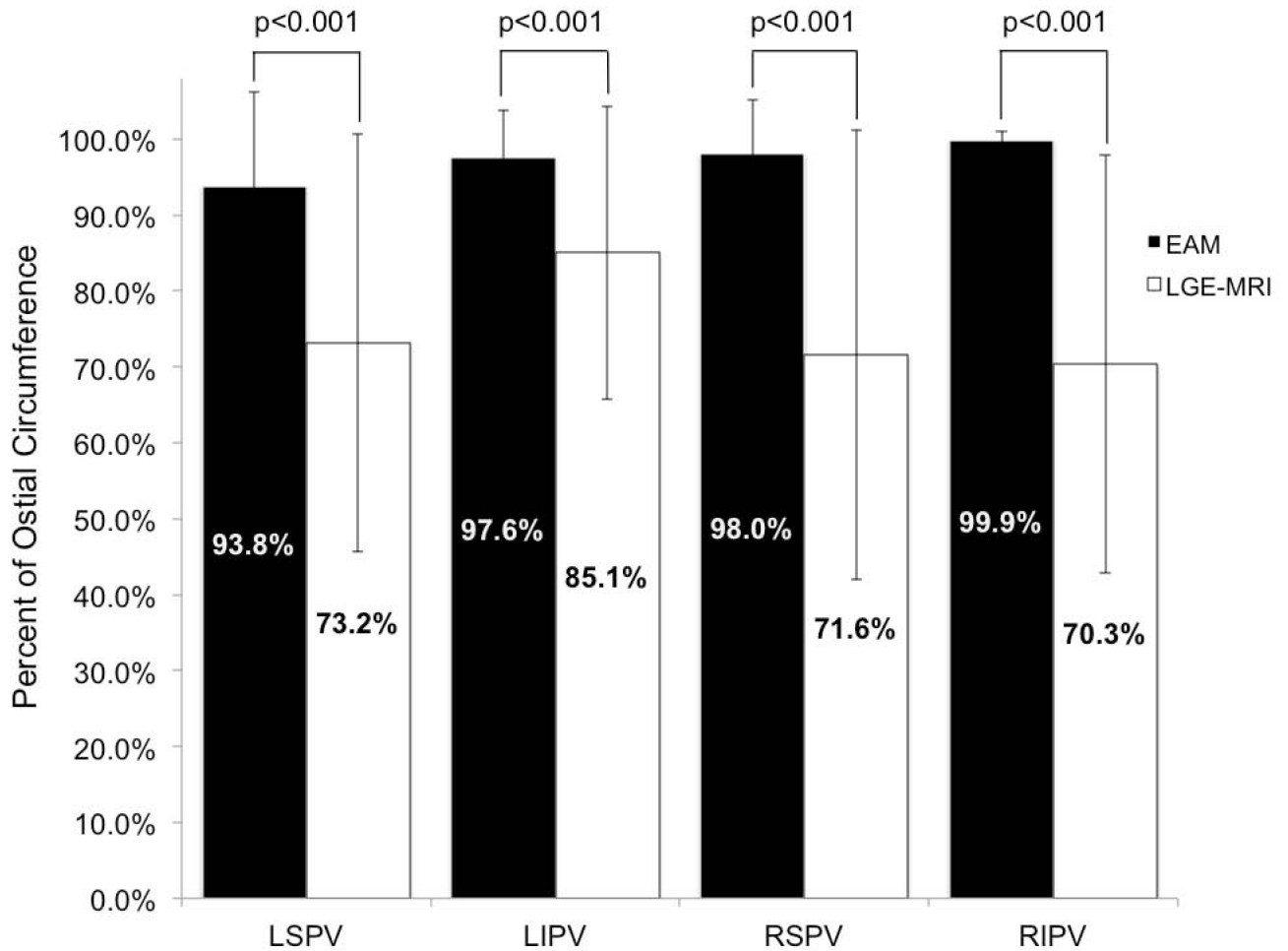


Figure 4.

Comparison of mean percent encirclement of each PV in EAM and scar from LGE-MRI (p-values <0.001 for each PV comparison). EAM: electroanatomic mapping; LGE-MRI: late-gadolinium enhancement magnetic resonance imaging; LSPV: left superior pulmonary vein; LIPV: left inferior pulmonary vein; RSPV: right superior pulmonary vein; RIPV: right inferior pulmonary vein.

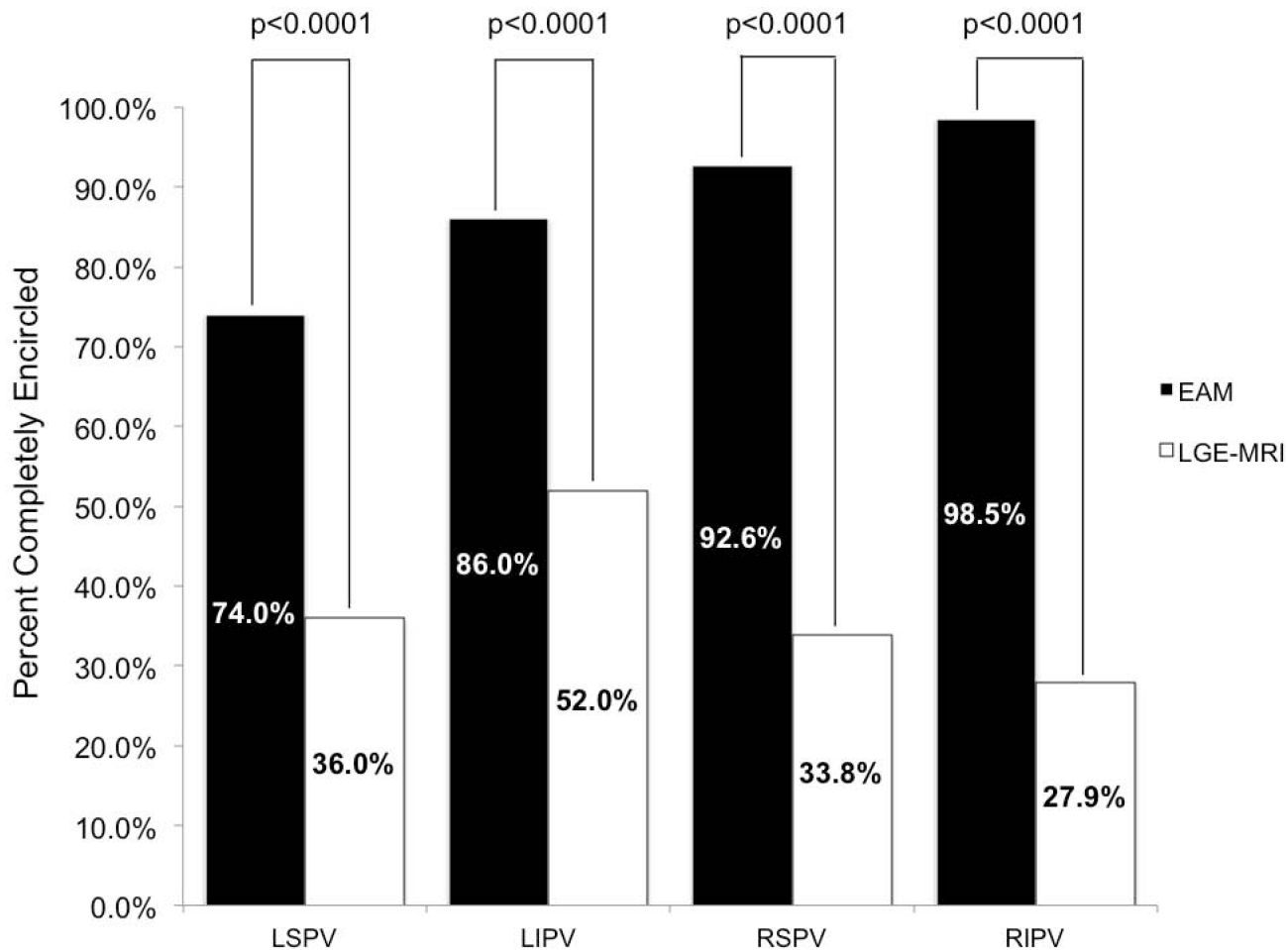


Figure 5.

Comparison of completely encircled PVs with ablation tags in EAM and scar in LGE-MRI. Complete encirclement defined as >90% encirclement of the PVs by ablation tags or scar. Differences were significant for all four vein pairs ($p < 0.001$). EAM: electroanatomic mapping; LGE-MRI: late-gadolinium enhancement magnetic resonance imaging; LSPV: left superior pulmonary vein; LIPV: left inferior pulmonary vein; RSPV: right superior pulmonary vein; RIPV: right inferior pulmonary vein.

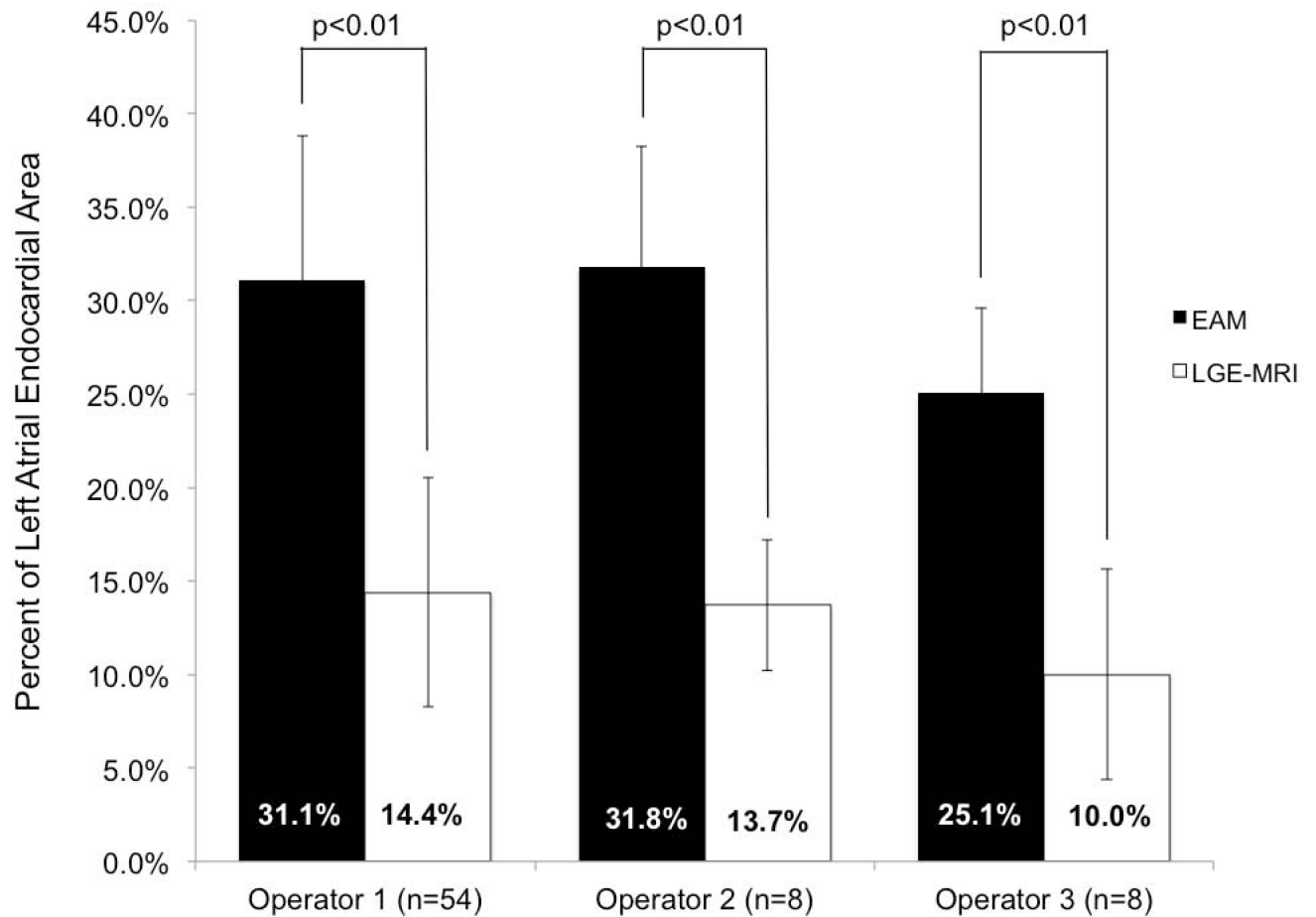


Figure 6.

Comparison of mean ablated area along with corresponding standard deviation in EAM and in LGE-MRI for each operator ($p < 0.01$). Means are presented in columns with standard deviation bars. LA: left atrium; EAM: electroanatomic mapping; LGE-MRI: late-gadolinium enhancement magnetic resonance imaging.

Table 1

Clinical characteristics of the patient population (n=70).

<u>Characteristic</u>	<u>Mean +/- Standard Deviation</u>
Age (Years)	65.0 ± 2.4
Body Mass Index	31.6 ± 7.2
	<u>Number (Percent of Population)</u>
Gender	
Male	54 (75.7%)
Female	16 (24.3%)
AF Type	
Paroxysmal	34 (48.1%)
Persistent	36 (51.9%)
Permanent	0 (0.0%)
Hypertension	41 (58.6%)
Diabetes Mellitus	8 (11.4%)
Obstructive Sleep Apnea	20 (28.6%)
Coronary Artery Disease	15 (21.4%)
Congestive Heart Failure	12 (17.1%)