The continuous challenge of AF ablation: From foci to rotational activity

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Abstract Pulmonary vein isolation (PVI) is central to ablation approaches for atrial fibrillation (AF), yet many patients still have arrhythmia recurrence after one or more procedures despite the latest technology for PVI. Ablation of rotational or focal sources for AF, which lie outside the pulmonary veins in many patients, is a practical approach that has been shown to improve success by many groups. Localized sources lie in atrial regions shown mechanistically to sustain AF in optical mapping and clinical studies of human AF, as well as computational and animal studies. Because they arise in localized atrial regions, AF sources may explain central paradoxes in clinical practice – such as how limited ablation in patient specific sites can terminate persistent AF yet extensive anatomical ablation at stereotypical locations, which should extinguish disordered waves, does not improve success in clinical trials. Ongoing studies may help to resolve many controversies in the field of rotational sources for AF. Studies now verify rotational activation by multiple mapping approaches in the same patients, at sites where ablation terminates persistent AF. However, these studies also show that certain mapping methods are less effective for detecting AF sources than others. It is also recognized that the success of AF source ablation is technique dependent. This review article provides a mechanistic and clinical rationale to ablate localized sources (rotational and focal), and describes successful techniques for their ablation as well as pitfalls to avoid. We hope that this review will serve as a platform for future improvements in the patient-tailored ablation for complex arrhythmias.

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Introduction

Ablation is increasingly used to treat atrial fibrillation (AF), yet the 1-2 year success rate of pulmonary vein isolation (PVI) remains 40-50% for persistent AF and 50-65% for paroxysmal AF. Studies of the often extensive ablation of lines, empiric electrogram targets and particularly the posterior left atrium often have not improved success versus PVI alone in multicenter trials.

Targeting AF sources has gained much attention in recent years. In the source model, AF is sustained by rotational (rotor) or focal sources in localized patient-specific regions of either atria. This model is now supported by wide evidence in many patients ranging from optical mapping of
human AF,7 multicenter (non-randomized) clinical trials of AF rotor ablation,8-13 optical mapping of animal AF,14 and mathematical analyses.15 Moreover, this model can reconcile the paradox that limited ablation can terminate persistent AF in some patients,8,16,17 while extensive (untargeted) ablation of left atrial regions can be ineffective in others.1,2,18

The source model may contradict the multwavelet hypothesis, in which AF is caused by disordered waves alone,19,20 although ongoing studies on the interaction between organized sources and disordered waves may reconcile this. A central tenet of the 'pure' disordered model is that extensive ablation limits the critical mass for wave propagation and will increase success, yet this is contradicted by multicenter trials1,2,6 with suboptimal results even in some surgical studies51 compared to original reports.

This review provides an overview of the science for localized drivers of human AF, and technical factors explaining why they may be revealed by some but not all mapping approaches. In particular, we focus on potential explanations for why some clinical ablation studies have been disappointing despite promising results at many independent centers. We hope to provide a mechanistic/clinical foundation to help reconcile debates in this field.

Initiation of human atrial fibrillation

Human arrhythmias have dynamic and static mechanisms.22 In AF, triggers such as ectopic beats,23 bursts of atrial tachycardia24 or varying autonomic balance25,26 may on occasion initiate AF as a dynamic process, despite a relatively static atrial architecture and fibrosis.27

Dynamics in the physiology of atrial repolarization and conduction can explain how triggers initiate AF. In Figure 1A, an ectopic beat produces dramatic oscillations of human left atrial action potential duration (APD),24,28,29 because the graph relating APD to diastolic interval (DI, time between beats; Figure 1B)30 is steep, so that an early beat (short DI) drastically shortens APD, lengthening subsequent DI, producing APD alternans and wavebreak. Figure 1C shows that triggers, such as from the PVs, may also abruptly slow human atrial conduction to facilitate reentry and AF (Figure 1D).31 These dynamics likely interact with atrial anatomy15 and/or fibrosis,31 explaining why sites of spiral wave reentry may initiate at spatially conserved sites for diverse triggers.

Mechanisms that sustain human atrial fibrillation after it has been triggered

Once AF is initiated by triggers from pulmonary veins23 or other sites, two central hypotheses may explain how disorganized wavefronts in AF sustain. In the multi-wavelet hypothesis,19 disorganized activity generates new wavelets in a stochastic fashion, where no specific atrial region plays a crucial role in the AF maintenance. This hypothesis therefore precludes "special" regions of the atria, such that structural elements e.g. fibrosis distribution are not critical.
to the maintenance of AF. A second, alternative, mechanism is that "special" regions of the atria do indeed exist and act as functional sources, manifest electrically in the form of spiral waves or focal activation patterns. These organized sources generate wavefronts that break down ("fibrillatory conduction") to explain the disorder of AF.

Some of the most important evidence for localized drivers of human AF comes from optical mapping. Figure 2 shows results from the gold standard of optical mapping of human AF. Rotational drivers are seen (Figure 2A) where ablation terminated AF to sinus rhythm (Figure 2B) and, in another specimen, 2 concurrent AF sources are seen in left atrium (Figure 2C). Sources hence lay in either atrium in these diseased atria, stabilized by micro-reentry around fibrotic and slowly conducting regions, which are less reported in animal models. One important methodological issue arising from this work, that we will revisit later, is that action potentials in AF may be quite regular (Figure 2D), yet traditional electrograms show additional deflections in AF ("noise", "far-field activation") which will reduce accuracy for mapping.

Reconciling differences between mapping studies of AF rotational drivers

Despite data in favor of rotational or focal drivers for human AF, there is an active debate on this issue. First, historical AF mapping shows only disorganized waves with no driving regions, albeit in patients rarely referred for ablation (with permanent AF at non-arrhythmia surgery). Second, organized drivers by frequency analysis may be unstable by activation or phase maps. This may reflect mapping of the epicardium in these studies, where drivers are less stable on optical maps. Third, AF-driver ablation outcomes are disappointing at some centers (Table 1 summarizes PVI and driver ablation studies). Mapping differences will alter reported AF mechanisms, and Rudy et al. show that combined activation with phase may be optimal in AF. The method used by our group, Focal Impulse and Rotor Modulation (FIRM), uses such a combined activation and phase approach. However, it is unresolved to what extent conflicting studies reflect mapping methods, patient selection, species specificities or other differences.

Since long-term ablation success varies dramatically between centers even for PVI, reconciling differences in AF mapping requires other study designs. On the basis that sites where ablation terminates persistent AF may have mechanistic relevance, we are systematically comparing maps created by different techniques for analyzing the same raw electrographic data in cases of clear termination of persistent AF by ablation in the ongoing COMPARE-AF study.

Figure 3 shows a patient in whom ablation outside the PVs before PVI terminated persistent AF to sinus rhythm. In panels C-E, detailed analysis of raw AF electrograms prior to ablation revealed sustained rotational activation by 3 methods. Of these methods, traditional activation maps of AF electrograms may be confused by spurious deflections, perhaps reflecting far field (e.g. Figure 2, compared to optical maps), and often showed only disorder at AF termination sites where phase and combined phase/activation maps showed rotations.

We reported an early series of such patients, which was the inspiration for the international COMPARE-AF study (NCT02997254) from which mapping data and code are being made available online to accelerate developments in AF mapping.

Reconciling heterogeneity in the success of source-guided ablation

Initial reports of mapping-guided ablation of AF drivers were positive, using endocardial mapping by FIRM in
<table>
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<tr>
<th>Year</th>
<th>Authors</th>
<th>N</th>
<th>Persistent AF (N)</th>
<th>Source mapping method</th>
<th>Automatic map read software?</th>
<th>Endpoint</th>
<th>Estimated cases/operator</th>
<th>1-Proc AF Free All (%)</th>
<th>1-Proc AF/AT Free Persist AF (%)</th>
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<tr>
<td>2012</td>
<td>Narayan et al.</td>
<td>107</td>
<td>76</td>
<td>FIRM</td>
<td>No</td>
<td>Termination, AF slowing</td>
<td>50</td>
<td>82</td>
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<td>2014</td>
<td>Haissaguerre</td>
<td>103</td>
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<td>2016</td>
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<td>42</td>
<td>42</td>
<td>FIRM</td>
<td>No</td>
<td>Remap, AT Not ablated</td>
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<td>47</td>
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<td>68</td>
<td>68</td>
<td>Novel Pigtaarray</td>
<td>No</td>
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<td>30</td>
<td>75</td>
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<td>170</td>
<td>106</td>
<td>FIRM</td>
<td>Yes (&gt;80% of cases)</td>
<td>Elimination on remap</td>
<td>55</td>
<td>74</td>
<td>62</td>
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<td>Total</td>
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<td>1066</td>
<td>806</td>
<td></td>
<td></td>
<td>Average</td>
<td>74.6%</td>
<td>64.9%</td>
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* With no PVI performed.
Figure 3  Rotational drivers for AF by multiple mapping methods. In a 67 year old woman, ablation (A) at the left atrial roof prior to PVI (B) terminates persistent AF to sinus rhythm. (C) Traditional activation mapping, (D) phase mapping using published methods \(^{38}\) and (E) Focal Impulse and Rotor Mapping (FIRM) each confirmed sustained rotations at the site where ablation terminated AF. Modified from references \(^{66}\) and \(^{48}\).

2011-12, \(^{8,49}\) body-surface mapping, \(^{10}\) novel mapping by Shih-Ann Chen et al.,\(^{50}\) Seitz et al.\(^{51}\) and other centers\(^{52,53}\) (Table 1). However, while independent centers continue to report positive results, \(^{11,12,54,55}\) outcomes are disappointing by some operators\(^{54-56}\) (Table 1).

Table 1 details the heterogeneity in outcomes from map-guided AF source ablation across centers. Notably, this does not appear to stratify by patient comorbidity or disease severity. For instance, Sommer, Hindricks et al.,\(^{54}\) and Miller et al.\(^{5,12}\) reported 70-80% maintenance of sinus rhythm by FIRM-ablation in populations that included challenging patients with prior failed procedures and long-standing persistent AF; as did Tomassoni,\(^{12}\) Rashid\(^{11}\) and others. Conversely, Buch, Shivkumar et al.,\(^{42}\) report 21% success and Steinberg et al. report 12% success\(^{55}\) in patients even with paroxysmal AF despite also performing PVI. It is difficult to reconcile these results based primarily on disease severity.

Much of the heterogeneity in results between centers may thus reflect technical differences. It is noted that reading AF rotational/focal source maps and/or guiding ablation accordingly are novel skills and not learned as part of traditional PVI. Indeed, in Table 1 lower success was seen at centers with fewer cases per operator, and in patients studied early (pre-2013) when no automatic tools were available to interpret maps to assist the operator. Although follow-up duration is typically a confounder, positive studies in Table 1 are reported at 1-2 years, 2 years\(^{12}\) and 3 years\(^{39}\) by separate groups.

Areas for technical improvement in source-guided ablation of AF

Three core components are required for effective source-guided ablation: effective broad-area mapping of both atria, precise identification of sources to target, and ensuring full elimination of target areas. Each of these components also requires urgent technical improvement, since they may explain much of the heterogeneity in outcomes between centers.

Our approach to widely map the atria is to use multi-polar basket catheters to analyze many wavefronts at the same time.\(^{49}\) Figure 4 shows optimal and suboptimal basket placement. Various basket sizes and types are available from an increasing number of vendors. We select the most appropriate basket size based on atrial size from intracardiac echocardiography or computed tomography. Since the atrium is not a spherical structure, the basket may have to be moved sequentially for optimal mapping. Figure 4A shows a single suboptimal left atrial basket position, which does not contact the walls (noted by its spherical shape). Figure 4B shows successive repositioning in this case to cover most of the atria in 2-3 epochs, showing spline deformation i.e. good contact. Figure 4C shows that suboptimal basket positioning may not be recognized.\(^{56}\) Figure 4D shows careful basket positioning at a high-volume center in a recent large series.\(^{57}\)

Theoretically, baskets can resolve 1-2 cm diameter reentry circuits in human AF as predicted by Allessie et al.\(^{58}\)
and confirmed in human optical maps (Figure 2A). Conversely, if electrodes are too closely spaced, calculated wave propagation may fall within measurement error. For instance, for atrial conduction velocity in AF patients of 40 cm/sec, reported errors in assigning onset time in AF (≈5-10 ms) translate to a distance of 2-4 mm (≈0.005 to 0.010). Closer electrode separation than this will be attempting to identify circuits within measurement noise, and will have less confidence. A related issue is that closely-spaced electrode arrays typically cover small distances simultaneously, which may also miss the 1-2 cm diameter circuits in human AF noted in optical maps.

Successful source ablation should fully cover the affected source areas, and contact force sensing catheters may help in this regard. Incomplete coverage may explain lower success rates in some source ablation studies. For instance, Figure 4E (Gianni et al. in patients from OASIS) shows sparse lesions that may not have eliminated sources in the long-term even if AF terminates acutely. Conversely, in the successful series in Figure 4F, ablation lesions densely cover the affected atrial regions.

As shown in Figure 5, incompletely eliminated source regions may cause recurrent post-ablation AF or AT. Thus, it is critical to ablate the entire affected region, although the science on how much atrium to ablate has yet to be fully established.

Finally, source-guided ablation does not appear to increase complication rates over traditional ablation alone, and is likely not pro-arrhythmic in most patients, and may be superior to historical endpoints of AF termination. Lack of AF termination may reflect many mechanisms including residual AF sources, but clinical results have been promising even in such patients if sources are eliminated.

Figure 4 Reported pearls and pitfalls in AF source ablation. (A) Suboptimal basket position in left atrium. (B) Multi-Basket Mapping, the method of choice in large atria. (C) Suboptimal basket in a disappointing report, consistent with inadvertent LV prolapse – conical shape, anterolateral to coronary sinus, ventricular electrograms; (D) good basket position covering atrium well; (E) sparse lesions over source area in disappointing series; (F) dense source area lesions in a more successful series.
Conclusions

Evidence continues to mount that human AF is maintained by rotational and focal sources, and that targeting these areas may improve outcomes over PVI alone. We have outlined the scientific rationale for source ablation, and practical strategies for ablation associated with promising outcomes in multicenter studies. Analysis of smaller less promising studies suggests that suboptimal basket placement, resulting in greater difficulty in reading maps and in targeting ablation may explain at least some of these discrepancies. Future improvements in AF source ablation may be facilitated by better computational interpretation of AF maps, comparative studies between potentially complementary mapping approaches, and improvements in basket design. Combined mechanistic and imaging studies may enable better and functional classification of AF that may enable better patient tailoring of AF ablation. Ultimately a better mechanistic and clinical understanding of AF may pave the way for novel drug discovery or regenerative therapies for AF. This is an exciting prospect.

Disclosures

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References

8. Narayan SM, Krummen DE, Shivkumar K, et al. Treatment of atrial fibrillation by the ablation of localized sources: the conventional ablation for atrial fibrillation with or without focal...
Foci and rotational activity in AF


