Impact of the arrhythmogenic potential of long lines of conduction slowing at the pulmonary vein area 🤖

Elisabeth M.J.P. Mouws, MD,* Lisette J.M.E. van der Does, MD,* Charles Kik, MD,† Eva A.H. Lanters, MD, * Christophe P. Teuwen, MD, * Paul Knops, BSc, * Ad J.J.C. Bogers, MD, PhD, † Natasja M.S. de Groot, MD, PhD*  

From the *Department of Cardiology, Erasmus Medical Center, Rotterdam, The Netherlands, and †Department of Cardiothoracic Surgery, Erasmus Medical Center, Rotterdam, The Netherlands.

BACKGROUND Areas of conduction delay (CD) or conduction block (CB) are associated with higher recurrence rates after ablation therapy for atrial fibrillation (AF).

OBJECTIVE Thus far, there are no reports on the quantification of the extensiveness of CD and CB at the pulmonary vein area (PVA) and their clinical relevance.

METHODS Intraoperative high-density epicardial mapping of the PVA (interelectrode distance 2 mm) was performed during sinus rhythm in 268 patients (mean ± SD [minimum–maximum] 67 ± 11 [21–84] years) with and without preoperative AF. For each patient, extensiveness of CD (conduction velocity 17–29 cm/s) and CB (conduction velocity <17 cm/s) was assessed and related to the presence and type of AF.

RESULTS CD and CB occurred in, respectively, 242 (90%) and 183 (68%) patients. Patients with AF showed a higher incidence of continuous conduction delay and block (CDCB) lines (AF: n = 37 [76%]; no AF: n = 132 [60%]; P = .046), a 2-fold number of lines per patient (CD: 7 [0–30] vs 4 [0–22], P < .001; CB: 3 [0–11] vs 1 [0–12], P = .003; CDCB: 2 [0–6] vs 1 [0–8], P = .004), and a higher incidence of CD or CB lines ≥6 mm and CDCB lines ≥16 mm (P = .011, P = .025, and P = .027). The extensiveness of CD, CB, and CDCB could not distinguish between the different AF types.

CONCLUSION Patients with AF more often present with continuous lines of adjacent areas of CD and CB, whereas in patients without AF, lines of CD and CB are shorter and more often separated by areas with normal intra-atrial conduction. However, a considerable overlap in the amount of conduction abnormalities at the PVA was observed between patients with a history of paroxysmal and persistent AF.

KEYWORDS Atrial fibrillation; Conduction; Epicardial mapping; Pulmonary veins; Sinus rhythm

(Heart Rhythm 2018;■:1–9) © 2018 The Authors. Published by Elsevier Inc. on behalf of Heart Rhythm Society. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

Introduction

The pulmonary vein area (PVA) has been of particular interest in the pathophysiology of atrial fibrillation (AF) ever since Haïssaguerre et al demonstrated bursts of rapid ectopic beats as triggers for spontaneous AF. Since then, treatment strategies for AF mainly focus on the isolation of the PVA by endocardial and/or epicardial ablation. Yet, recurrence rates are considerable for both patients with paroxysmal AF and those with persistent AF and are likely the result of either reconnection or transition of AF from a trigger-driven to a more substrate-driven disease.7

To date, AF recurrences after ablation procedures remain difficult to predict. Yet, fibrosis at the left atrial (LA) posterior wall, resulting in conduction delay (CD) or conduction block (CB), appears to be associated with higher recurrence rates.3,4 It has been suggested that the assessment of electrophysiology, including low voltages, fractionation, and conduction abnormalities, during sinus rhythm (SR) at the PVA may facilitate the identification of target sites for ablation or can be used to predict AF recurrences after ablation therapy.5–9

In several mapping studies, a line of CB running vertically between the right and left pulmonary veins during SR was identified.10–12 This CB line varied between patients in its continuity and could in some patients be altered by pacing, indicating that it was partly functional in nature.10–12 Furthermore, this line was more frequently observed in patients with AF or mitral valve regurgitation.11,12 On the basis of histological findings in postmortem hearts, the authors suggested that abnormal conduction was the result of a change in myocardial fiber direction.10 Aside from this

Dr de Groot is supported by grants from the Erasmus Medical Center fellowship, Dutch Heart Foundation (grant no. 2012TI0046), LSH-Impulse grant 40–43100–98–008, CVON AFFIP (grant no. 914728), and VIDI grant (grant no. 91717339). Dr Teuwen is supported by a grant from the Dutch Heart Foundation (grant no. 2016T071). Address reprint requests and correspondence: Dr Natasja M.S. de Groot, Unit Translational Electrophysiology, Department of Cardiology, Erasmus Medical Center, ’s Gravendijkwal 230, RG-632, 3015CE Rotterdam, The Netherlands. E-mail address: nmsdegroot@yahoo.com.
line of CB, other areas of conduction disorders were observed in only a minority of patients.\textsuperscript{10–12} However, the degree and extent of conduction abnormalities during SR at the PVA have never been quantified and correlated with the different types of AF as defined by the European Society of Cardiology guidelines.\textsuperscript{13} The goals of the present intraoperative high-resolution epicardial mapping study were therefore to detect and quantify conduction abnormalities at the PVA in a large cohort of patients during SR and to investigate the association with AF persistence.

**Methods**

**Study population**

The study population consisted of 268 successive adult patients undergoing elective coronary artery bypass grafting, aortic or mitral valve surgery, or a combination of valvular and bypass grafting surgery. This study was approved by the institutional medical ethical committee (MEC2010-054/MEC2014-393). Written informed consent was obtained from all patients, and clinical data were extracted from electronic patient files. A detailed description of the methods is provided in the Supplemental Material.

**Epicardial high-resolution mapping**

Epicardial high-resolution mapping of the PVA was performed during SR from the transverse sinus along the borders of the right and left pulmonary veins down toward the atrioventricular groove (Figure 1), as previously described.\textsuperscript{14} Local activation maps of the right and left pulmonary veins during SR were constructed by annotating the steepest negative slope of atrial potentials recorded at every electrode (see also Supplemental Figure 1). Heterogeneity in conduction was determined by quantifying the amount, number, and length of lines of CD, CB, and continuous conduction delay and block (CDCB) and its differences between patient groups on a 2 mm resolution scale. Lines of CD and CB were defined as time differences ($\Delta t$) of, respectively, 7–11 and $\geq 12$ ms between adjacent electrodes.\textsuperscript{15,16}

**Statistical analysis**

Normally distributed data are presented as mean ± SD (minimum–maximum). Skewed data are presented as median (minimum; interquartile range; maximum) and analyzed using Mann-Whitney $U$ tests. Categorical data are expressed as numbers and percentages and analyzed using the $\chi^2$ test.
or Fisher exact test, as appropriate. Receiver operating characteristic curves for the difference in CD and CB lengths were constructed, and cutoff values were based on sensitivity > 50% and 1 - specificity < 50%. Multivariate regression analysis was performed to identify independent predictors of CD and CB. A P value of < .05 was considered statistically significant.

Results

Study population

The characteristics of the study population (N = 268; 196 men [73%]; mean [minimum–maximum] age 67 ± 11 [21–84] years; mean [minimum–maximum] body mass index [BMI] 28 ± 5 [18–55] kg/m²) are summarized in Table 1. Patients had either ischemic heart disease (IHD) (n = 157 [59%]) or ischemic and valvular heart disease ([i]VHD) (n = 111 [41%]; only valvular disease: n = 63 [24%]). LA dilation was present in 58 patients (22%) and 49 patients (18%) had a history of AF. Most patients had normal left ventricular function (n = 203 [76%]) and used class II antiarrhythmic drugs (n = 183 [68%]).

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (y)</td>
<td>67 ± 11 (21–84)</td>
</tr>
<tr>
<td>Sex: male</td>
<td>196 (73)</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>28 ± 5 (18–55)</td>
</tr>
<tr>
<td>Underlying heart disease</td>
<td></td>
</tr>
<tr>
<td>IHD</td>
<td>157 (59)</td>
</tr>
<tr>
<td>[i]VHD</td>
<td>111 (41)</td>
</tr>
<tr>
<td>Aortic valve stenosis</td>
<td>69 (26)</td>
</tr>
<tr>
<td>Aortic valve insufficiency</td>
<td>6 (2)</td>
</tr>
<tr>
<td>Mitral valve insufficiency</td>
<td>36 (13)</td>
</tr>
<tr>
<td>Left atrial dilation &gt;45 mm</td>
<td>58 (22)</td>
</tr>
<tr>
<td>History of AF</td>
<td></td>
</tr>
<tr>
<td>Paroxysmal</td>
<td>49 (18)</td>
</tr>
<tr>
<td>Persistent</td>
<td>38 (14)</td>
</tr>
<tr>
<td>Persistent</td>
<td>11 (4)</td>
</tr>
<tr>
<td>Left ventricular function</td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>203 (76)</td>
</tr>
<tr>
<td>Mild dysfunction</td>
<td>52 (19)</td>
</tr>
<tr>
<td>Moderate dysfunction</td>
<td>11 (4)</td>
</tr>
<tr>
<td>Severe dysfunction</td>
<td>2 (1)</td>
</tr>
<tr>
<td>Antiarrhythmic drugs</td>
<td></td>
</tr>
<tr>
<td>Class I</td>
<td>2 (1)</td>
</tr>
<tr>
<td>Class II</td>
<td>183 (68)</td>
</tr>
<tr>
<td>Class III</td>
<td>12 (5)</td>
</tr>
<tr>
<td>Class IV</td>
<td>3 (1)</td>
</tr>
</tbody>
</table>

Values are presented as mean ± SD (minimum–maximum) or as n (%).

BMI = body mass index; IHD = ischemic heart disease; [i]VHD = ischemic and valvular heart disease; VHD = valvular heart disease.

Incidence of CD and CB

Most patients showed lines of CD (n = 242 [90%]) and CB (n = 183 [68%]) at the PVA during SR. The number of lines of CD (4 [0–30]) was significantly higher than that of CB lines (1 [1–12]) (P < .001), though the maximum length of CB lines was longer (median [minimum; interquartile range; maximum], CD: 6 [2; 4–10; 20] mm; CB: 8 [2; 4–12; 44] mm; P < .001) (Figure 2). A clear turning point was observed at a length of ≥8 mm from which point on the incidence of CB lines exceeded the incidence of CD lines. Most patients also had continuous lines of CDCB (n = 169 [63%]; median number 1 [0–6]; maximum length 14 [4–72] mm).

A longitudinal line of CD or CB running vertically between the left and right pulmonary veins from superior to inferior positions was observed in 14 patients (5%), though varying in its continuity and length. Typical examples of activation maps and corresponding isochrones and CD/CB maps of these patients are shown in Figure 3. The incidence of this line was similar between patients without and with AF (P = .295), as well as between patients with IHD and those with [i]VHD (P = .503). However, this line was more often observed in patients with LA dilation (n = 6 [10%]) than in patients without LA dilation (n = 8 [4%]) (P = .048). As displayed in Table 2, multivariate regression analysis revealed only the presence of AF episodes as an independent predictor of long lines of CD and CB at the PVA; clinical characteristics including IHD, [i]VHD, LA dilation, sex, BMI, older age, and left ventricular dysfunction were not.

Association between AF and heterogeneity in conduction

The upper panel of Figure 4 displays typical examples of activation maps and the corresponding CD/CB maps obtained from a patient without AF and a patient with AF.

Patients with AF more often have continuous lines of CDCB as compared with patients without AF, as demonstrated in the middle left panel (AF: n = 37 [76%]; no AF: n = 132 [60%]; P = .046).

The number of lines of CD, CB, and CDCB in patients with AF was approximately 2-fold the number observed in patients without AF (CD: 7 [0–30] vs 4 [0–22], P < .001; CB: 3 [0–11] vs 1 [0–12], P = .003; CDCB: 2 [0–6] vs 1 [0–8], P = .004).

The maximum length of continuous CDCB lines in patients with AF ranged from 8 to 72 mm, whereas in patients without AF this length ranged from 4 to 42 mm; CDCB lines ≥16 mm occurred more often in patients with AF (n = 20 [41%] vs n = 50 [25%]; P = .027).

Hence, the presence of AF episodes was strongly associated with increased heterogeneity in conduction, marked not only by a higher incidence of CB and CDCB but also by a higher number of lines of CD, CB, and CDCB and more importantly longer lines of CD, CB, and CDCB.

Thus, patients with AF more often present with continuous lines of adjacent areas of CD and CB, whereas in patients without AF, lines of CD and CB are more often separated by normal intra-atrial conduction. These findings were validated in an age- (5-year range),
BMI- (2-kg/m² range), sex-, and type of surgery–matched case-control analysis (AF N=35, No AF N=35).

In addition, patients who developed de novo AF in the early postoperative phase (70 patients without preoperative AF [32%]) showed a trend toward more CB lines (P = .055), a higher number of continuous CDCB lines (P = .030) at the PVA, and a trend toward a higher incidence of long CB lines (>6 mm) (P = .073).

Severity of conduction abnormalities vs clinical AF classification

Figure 5 provides typical examples of PVA activation combined with the corresponding CD and CB maps obtained from 2 patients with paroxysmal AF and 2 patients with persistent AF; the amount of conduction abnormalities in 1 patient with paroxysmal AF is even higher than in the patient with persistent AF.

Figure 5 shows that there is a large interindividual variation in the amount of conduction abnormalities in both paroxysmal and persistent AF groups. There is also no difference between patients with paroxysmal AF and patients with persistent AF in the number of CD, CB, and CDCB lines (P = .442, P = .535, and P = .951). Also, incidences of CD, CB, and CDCB were similar (P = .204; P = .835, and P = .708); receiver operating characteristic curve analyses could not identify a cutoff value for the length of lines distinguishing patients with persistent AF from those with paroxysmal AF. The duration of AF history was similar for patients with paroxysmal AF and those with persistent AF (P = .429).

Hence, although this is only a small group of patients, the overlap in the severity of conduction abnormalities suggests that severity of conduction abnormalities at the PVA does not seem to clearly differentiate patients with paroxysmal AF from patients with persistent AF.

Discussion

Key findings

Intraoperative high-resolution epicardial mapping of the PVA during SR for the first time quantified and characterized different types of conduction abnormalities and related it to the presence of AF episodes. Current data demonstrated that patients with AF have more and longer lines of CD, CB, and CDCB, whereas in patients without AF, short lines (<6 mm) of CD and CB separately are more diffusely present. Furthermore, the severity of conduction abnormalities at the PVA during SR does not differentiate between patients with paroxysmal AF and those with persistent AF.

Conduction abnormalities at the PVA

To our knowledge, only 3 previous studies have investigated conduction abnormalities at the posterior wall of the LA in humans during SR. In an endocardial noncontact mapping study by Markides et al., conduction at the LA was analyzed during SR in 19 patients with a history of paroxysmal AF. They observed a vertical line of CB (inter-electrode time interval 30 ms) extending from the LA roof...
across the posterior LA wall turning septally below the ostium of the right inferior pulmonary vein proceeding anteriorly toward the septal mitral annulus. This CB line was present in all patients, though varied in its continuity, particularly during pacing. On the basis of histological findings in postmortem hearts, Markides et al suggested that the line of CB was caused by an abrupt change in myocardial fiber orientation at the subendocardium.

Roberts-Thomson et al performed epicardial mapping during SR in 34 patients without AF. They observed a similar

Figure 3  Longitudinal line of CD/CB between the right and left pulmonary veins. Typical examples of activation maps with a line of CD (blue lines), CB (red lines), or CDCB running downward between the right and left pulmonary veins, though varying in its continuity. Corresponding isochrone maps (per 5 ms) and CD/CB maps are shown next to the activation maps. Arrows indicate the main wave trajectory; local activation times are provided next to the arrows. Lightning bolts indicate areas of simultaneous activation. Color classes per 10 ms. CB = conduction block; CD = conduction delay; CDCB = continuous conduction delay and block.
line of functional CD, defined as a conduction velocity between 10 and 20 cm/s, running vertically across the PVA, though only occurring in a minority of 5 patients. In contrast to the findings of Markides et al, during pacing from superior and inferior positions at the PVA, the line of CD now appeared in all patients. In a subsequent study, epicardial SR mapping after electroconversion in 16 patients without AF and 5 patients with persistent AF showed a similar vertical CD line in 2 patients without AF whereas this was observed in 4 patients with AF. However, when pacing from different sites, the CD line again appeared in all patients without AF and the line of functional CD now appeared in all patients.11

In contrast to previous studies, we observed conduction delays in 16 patients without AF and 5 patients with persistent AF showing a similar vertical CD line in 2 patients without AF whereas this was observed in 4 patients with AF. However, when pacing from different sites, the CD line again appeared in all patients without AF and the number of CD lines increased in patients with AF to a maximum of 3 vertical lines running parallel to each other across the PVA. Although the findings of Markides et al and Roberts-Thomson et al appear to contradict each other, it may be concluded that this line of CD was more evident in patients with AF and was, at least in part, functional, as it varied during different pacing conditions. Besides the vertical line of abnormal conduction, no other CD/CB lines were observed in these studies.

In contrast to these previous studies, we observed conduction abnormalities scattered across the PVA with no clear predilection site. Lines of CD occurred in almost all patients and CB in approximately 70% of the population. The fact that the aforementioned studies did not observe any other lines of CD or CB at the PVA is remarkable, especially since study populations consisted of patients with IHD, patients with AF, and patients with LA dilation due to mitral regurgitation. In all these patients, areas of fibrosis would be expected, particularly at the LA posterior wall. Our CB criteria correspond with a conduction velocity of <17 cm/s, which is in the range of the CD criteria of Roberts-Thomson et al. Therefore, although our cutoff criteria are slightly more sensitive, the higher incidence of CD/CB cannot be totally explained by differences in cutoff values. Yet, the higher resolution of the mapping system used in the present study may be the explanation for this discrepancy, as it contains the unique ability to identify lines of CD and CB with a minimum length of 2 mm. Furthermore, we did not set a minimum length or wavefront propagation criterion for lines of CD and CB, as opposed to previous studies.

In our cohort, only a minority of patients showed a longitudinal line running downward between the left and right pulmonary veins, which might be similar to the line observed in previous studies. However, this line varied in length and continuity and practically never consisted of a line of CB running continuously from the superior to the inferior of the posterior wall. The precise nature of this line so far remains unclear. If, as suggested by previous studies, a histological change in fiber direction would be the underlying cause, we would expect it to occur in the majority of patients during SR.

Conduction abnormalities and AF

In correspondence to previous studies, increased amounts of CD, CB, and CDCB at the PVA were observed in patients with AF. In patients with AF, a higher incidence of CB,
CDCB, and an almost 2-fold number of separate CD, CB, and CDCB lines per patient was observed. Also, CD, CB, and CDCB lines extended over larger areas.

These observations suggest a critical role for the spatial distribution of conduction abnormalities in AF development. A certain length of an area of abnormal conduction is required for reentry to occur; this phenomenon was first demonstrated by Ortiz et al\textsuperscript{17} in 7 canine hearts with sterile pericarditis. In this study, the critical role of the length of an area of functional block in the right atrial free wall was observed. In the case of stable atrial flutter, a functional CB line of 24 mm was observed, enabling reentry to occur.\textsuperscript{17} When the cycle length decreased, areas of slow conduction disappeared, resulting in a shorter line of functional CB with a mean length of 16 mm.\textsuperscript{17} This resulted in unstable reentrant circuits migrating across the atrial wall, giving rise to AF.\textsuperscript{17} When the atrial wall already contains continuous long lines of structural CD and CB, it is likely more vulnerable to reentry circuits to occur or for areas of functional block to connect, thereby reaching the critical length for AF initiation.

**Figure 4** Differences in electropathology between patients without and with AF. Upper panels: Typical examples of activation maps of a patient without AF and a patient with AF. Middle and lower panels: Patients with AF show more electropathology at the pulmonary vein area. Patients with AF particularly show a higher incidence of CB and CDCB, a higher number of CD, CB, and CDCB lines per patients, and also longer lengths of CD (blue lines), CB (red lines), and CDCB lines. Color classes per 10 ms. AF = atrial fibrillation; CB = conduction block; CD = conduction delay; CDCB = continuous conduction delay and block.
The future of AF therapy

Despite the fact that conduction abnormalities are more profound in patients with AF, the clinical categories of AF do not correspond with the amount of conduction disorders at the PVA during SR. In a previous study, we demonstrated a considerable intra-atrial variation in the distribution of conduction disorders across the right atrium and LA, indicating that a low amount of CB at the PVA does not necessarily implicate a low amount of CB at other atrial regions. Hence, either the arrhythmogenic substrate underlying AF is not located at the PVA in these patients or, although CD and CB measured during SR are indicators of structural conduction abnormalities, functional conduction disorders may only be revealed during triggers or AF.

To date, ablation treatment strategies for AF focus primarily on isolation of the pulmonary veins. However, recurrence rates remain unsatisfactory. Recent studies have shown the complex and heterogeneous etiology

Figure 5  Overlap in electropathology between paroxysmal AF and persistent AF. Upper left panel: Activation maps of a patient with ischemic heart disease and a patient with valvular heart disease with paroxysmal AF, both diagnosed 3 months before surgery. The corresponding CD/CB maps (blue lines, CD; red lines, CB) show a relatively small amount of CD/CB in the first patient, whereas the second patient has a large amount of CD/CB. Upper right panel: Activation maps of patients with persistent AF. Both patients underwent mitral valve surgery and were diagnosed with persistent AF, respectively, 3 and 6 months before surgery; both patients underwent electrophysiological evaluation of the atrial tissues. In this case also, one patient has a relatively small amount of electrophysiological disarray in the atrial tissues, whereas the other patient has a large amount of CD/CB. Hence, a considerable overlap in the amount of conduction disorders is observed between paroxysmal AF and persistent AF, which is also quantified in the lower panel, showing the number of lines per patient and the distribution of lengths of these lines. Color classes per 10 ms. AF = atrial fibrillation; CB = conduction block; CD = conduction delay; CDCB = continuous conduction delay and block.
of fractionated potentials, providing a possible explanation of
the low success rate of ablation therapy targeting these com-
plex fractionated potentials.19

Furthermore, recent studies have revealed that aside from
the well-known effects of increased renin-angiotensin-system
activation, cardiac endothelin-1 levels may play an in impor-
tant role in AF pathogenesis.20 Endothelin-1 expression, pro-
moting myocyte hypertrophy and interstitial fibrosis, was in-
creased during AF compared to SR.20 Particularly, endothelin-1
levels were higher at the LA and in patients with VHD, leading to
the presumption that this may play a sub-

tantial role in the vulnerability of these patients for AF de-
velopment.20 Also, increased endothelin-1 levels are regarded as
an important factor in AF persistence. As endothelin-1 produc-
tion is stretch mediated, atrial regions subjected to greater wall
stress may also produce higher levels of endothelin-1, thereby
leading to regional differences in conduction abnormalities.20

Study limitations
Whether general anesthesia influences conduction is yet to be
investigated; however, as a standard anesthetic protocol was
used for all patients, equal dispersion of possible effects can
be assumed. The number of patients with AF was relatively
small; thereby, when comparing patients with persistent and
paroxysmal AF, conclusions should be drawn cautiously. In
addition, although late gadolinium enhancement- magnetic
resonance imaging (LGE-MRI) is a feasible technique to detect
cardiac fibrosis, it was logistically and financially not possible
to perform LGE-MRI before surgery in these patients.

Conclusion
Intraoperative high-resolution epicardial mapping of the PVA
during SR for the first time quantified and characterized
different types of conduction abnormalities and demonstrated
that the presence of AF episodes is associated with continuous
lines of adjacent areas of CD and CB, whereas in patients
without AF, lines of CD and CB are shorter and more often
separated by areas with normal intra-atrial conduction. Pa-


tients with AF showed a 2-fold number of CD, CB, and
CDCB lines per patient, which also extended over longer
lengths. This study demonstrated a considerable overlap in
the amount of conduction abnormalities at the PVA between
patients with a history of paroxysmal and persistent AF. Stud-
ies quantifying the extensiveness of electrophotology
by various parameters, including conduction abnormalities,
may contribute to the future development of a more accurate
risk estimation of recurrent AF after ablation therapy and will
thereby enable more patient-tailored care in the future.

Appendix
Supplementary data
Supplementary data associated with this article can be found
in the online version at https://doi.org/10.1016/j.hrtm.2018.10.027.

References
of atrial fibrillation by ectopic beats originating in the pulmonary veins. N Engl
Allessie M, de Groot NMS. Atrial fibrillation: to map or not to map? Neth Heart
J 2014;22:259–266.
3. Ferrari R, Bertini M, Blomstrom-Lundqvist C, Dobres D, Kirchhof P, Pappone C,
Mavroudis J, Tamargo J, Tavazzi L, Vicedomini GG. An update on atrial fibrillation
4. Oukes RS, Badger TJ, Kholmovski EG, et al. Detection and quantification of left
atrial structural remodeling with delayed-enhancement magnetic resonance imag-
Altmann D, Piedra C, Hindricks G, Plokker W. Tailored atrial substrate modi-
fication based on low-voltage areas in catheter ablation of atrial fibrillation. Circ
Arrhythm Electrophysiol 2014;7:825–833.
6. Vlachos K, Efremidis M, Letsas KP, et al. Low-voltage areas detected by high-
density electroanatomical mapping predict recurrence after ablation for parox-
7. Masuda M, Fujita M, Iida O, et al. Left atrial low-voltage areas predict atrial fibril-
lation recurrence after catheter ablation in patients with paroxysmal atrial fibrilla-
8. Pachon MJC, Pachon MEI, Pachon MZ, Vargas RNA, Pachon DQV, Lopez MFJ.
Atrial line mapping on the atrial fibrillation: a new tool to assess atrial fibril-
9. Roberts-Thomson KC, Kistler PM, Sanders P, Morton JB, Haqqani HM, Stevenson I,
Vohra JK, Sparks PB, Kalman JM. Fractionated atrial electrograms during sinus rhythm:
10. Markides V, Schilling R, Ho S, Chow A, Wyn Davies D, Peters N. Characteriza-
tion of left atrial activation in the intact human heart. Circulation 2003;
107:733–739.
11. Roberts-Thomson KC, Stevenson IH, Kistler PM, Haqqani HM, Goldblatt JC,
Sanders P, Kalman JM. Anatomically determined functional conduction delay
in the posterior left atrium: relationship to structural heart disease. J Am Coll Car-
diol 2008;51:856–862.
12. Roberts-Thomson KC, Stevenson I, Kistler PM, Haqqani HM, Spence SJ,
Goldblatt JC, Sanders P, Kalman JM. The role of chronic atrial stretch and atrial
fibrillation on posterior left wall conduction. Heart Rhythm 2009;
6:1109–1117.
ment of atrial fibrillation developed in collaboration with EACTS. Eur J Heart
14. Mouws EMJP, Lanters EAH, Teuwen CP, van der Does LJME, Kik C, Knops P,
Yashk A, Bekkers JA, Bogers AJJC, de Groot NMS. Impact of ischemic and
valvular heart disease on atrial fibrillation: a high-resolution epicardial mapping
Allessie M. Electropathological substrate of longstanding persistent atrial fibril-
lation in patients with structural heart disease: epicardial breakthrough. Circulation
2010;122:1674–1683.
16. Spach MS, Dolber PC, Heidlage JF. Influence of the passive anisotropic prop-
ties on directional differences in propagation following modulation of the so-
dium conductance in human atrial muscle: a model of reentry based on
17. Ortiz J, Niwano S, Abe H, Rady Y, Johnson NJ, Wada AL. Mapping the conver-
sion of atrial flutter to atrial fibrillation and atrial fibrillation to atrial flutter: in-
Marion DMS, Brundel BJM, Bogers AJJC, Allessie MA, de Groot NMS. Spatial
distribution of conduction disorders during sinus rhythm. Int J Cardiol 2017;
19. van der Does LJME, Knops P, Teuwen CP, Serban C, Stareveld R, Lanters EAH,
Mouws EMJP, Kik C, Bogers AJJC, de Groot NMS. Unipolar atrial electrogram
morphology from an epicardial and endocardial perspective. Heart Rhythm 2018;
Wagoner DR. Association of left atrial endothelin-1 with atrial rhythm, size,
and fibrosis in patients with structural heart disease. Circ Arrhythm Electrophysiol