The Wolff–Parkinson–White Syndrome
A Test Bed for the Assessment of Myocardial Dyssynchrony?

Sylvestre Maréchaux, MD, PhD

Among patients with a Wolff–Parkinson–White (WPW) syndrome, approximately half will experience arrhythmia during their lifetime. Radiofrequency catheter ablation of the accessory atrioventricular pathway is the treatment of choice for this condition and is associated with a high success rate. Electrophysiology study is indeed the key procedure for identifying the target of radiofrequency ablation. The present prospective study, published in this issue of Circulation: Cardiovascular Imaging by Ishizu et al, investigated the ability of a new imaging technology based on 3-dimensional speckle-tracking (ST) strain echocardiography in localizing AP in WPW syndrome.1 This tool, named isochrone activation imaging (AI), was accurate enough to assess synchronized activation in the normal heart and to detect contractile abnormalities, which approximately matched ablation site in WPW patients with left- or right-sided AP. See Article by Ishizu et al

Localization of Accessory Pathways With Echocardiography, a 40-Year-Old Story

Historically, motion-mode (M-mode) echocardiography was the first effective modality for the ultrasonic detection of early pre-ejection events, concomitant with the delta wave in WPW syndrome. Because of its high temporal resolution (1000–3000 Hz), M-mode echocardiography can display the reduced amplitude of left ventricle (LV) posterior systolic wall motion for AP, emerging from LV basal free wall. This modality can also display rapid pre-ejection posterior septal motion associated with a slow anterior motion of the septum for AP, emerging from LV basal free wall.2 Twenty years later, pulsed-wave tissue doppler imaging (TDI) and high frame-rate tissue velocity imaging enabled AP-accurate localization in 50% more patients, when compared with the M-mode method.2 High temporal resolution (at least 150 Hz) is a potential advantage of TDI; however, clinical use was limited by artifacts, caused by myocardial translational motion, nonoptimal Doppler alignment, and poor reproducibility.

Approximately 10 years ago, 2-dimensional speckle-tracking echocardiography gained popularity in quantification of amplitude and timing of myocardial deformation, with a lower influence of cardiac motion as compared with TDI.3 Optimal frame rate for speckle-tracking echocardiography is 50 to 90 Hz, a much lower rate than that of M-mode or TDI. Nevertheless, using customized software, De Boeck’s team demonstrated that speckle-tracking echocardiography mapping matched the electrophysiology localization (with a tolerance of 1 contiguous segment) of the AP in 28 patients (85%) with a WPW syndrome.4 Importantly, early deformation associated with the AP was subsequently followed by reduced local deformation during systole, especially in patients with AP involving the interventricular septum (IVS). In a cohort of 40 patients with WPW syndrome referred for radiofrequency ablation, longitudinal speckle-tracking echocardiography strain accurately identified early basal pre-ejection LV deformation in 38 patients (95%), whereas early abnormal basal motion was not observed in control patients. Moreover, contractile abnormalities were found to be at worst one adjacent segment different compared with the AP ablation site in all 38 patients. Right-sided AP was associated with IVS contractile abnormalities, as previously reported with M-mode. The decrease in IVS deformation in AP involving the interventricular septum was more obvious than the decrease in LV free wall deformation in AP involving the LV free wall. Ishizu et al are to be praised for their careful description of the value of 3-dimensional ST strain echocardiography to noninvasively localize AP in WPW syndrome.5 The advantage of 3-dimensional ST strain is the ability to assess area strain, a combination of circumferential and longitudinal strain of the entire ventricle. From a technical point of view, full volume data sets of the LV and the RV are acquired from apical views with a wide angle, to cover both ventricles, with a volume rate of at least 20 Hz (31±3 Hz, range 22–40 Hz), which is significantly lower than with other techniques. The authors used noninvasive isochrone AI, a technique previously validated against electric activation mapping in 7 patients with cardiomyopathies, 5 of them receiving cardiac resynchronization therapy (CRT) and 2 with an left bundle branch block (LBBB) during catheter ablation therapy of ventricular tachycardia.6 This method identifies the timing of deformation on each LV or RV segment and applies a time-dependent color-coded mapping either on a plastic bag model or on a polar map of the LV or RV. Segment deformation is coded by means of a graded color scale increasing in time, and color is applied when segment

The opinions expressed in this article are not necessarily those of the editors or of the American Heart Association.

From the Université Lille Nord de France/GCS-Groupement des hôpitaux de l’institut Catholique de Lille/Faculté Libre de Médecine, Cardiology Department, Université Catholique de Lille, 59000 Lille, France, and INSERM U 1088, Université de Picardie, Amiens, France.

Correspondence to Sylvestre Maréchaux, MD, PhD, GCS–Groupement des hôpitaux de l’institut Catholique de Lille, 115 rue du Grand But, 59462 Lomme Saint Philibert/Faculté Libre de Médecine, Université Catholique de Lille, Cardiology Department, 115 rue du Grand But, 59462 Lomme Cedex, INSERM U 1088, Université de Picardie, Amiens, France. E-mail sylvestre.maréchaux@gmail.com


Circ Cardiovasc Imaging is available at http://circimaging.ahajournals.org

DOI: 10.1161/CIRCIMAGING.116.005112
deformation is of at least 25% of maximal deformation. Hence, this method allows for visual identification of both early and late deformed segments. Using this approach, the sites for the first LV and RV endocardial activation (the breakthrough sites) were frequently identified at the LV and RV apical or mid-walls in normal subjects. In contrast, the breakthrough site was perianular and basal in WPW patients. In 2 patients who underwent CARTO® mapping, AI mapping completely matched the CARTO® mapping and the ablation site. Agreement between AI and AP localization was perfect in only 38% of patients, but was observed in 87% of patients when applying a 2 o’clock–range tolerance for AP localization. In addition, the AI method was more precise than the electrocardiogram to accurately detect AP localization. Method reproducibility was excellent. Unexpectedly, septal area strain was not different between controls and WPW patients with AP involving the IVS, but was significantly reduced in basal and midanterior LV walls in patients with an AP involving the LV free wall. It is noteworthy that image quality issue, similar to other ultrasonic techniques, is inherent to 3-dimensional ST strain. Despite these potential limitations, this study presents promising data regarding a new visual method to accurately diagnose short-lived early events and is thus a potential new tool for the assessment of myocardial dyssynchrony.

WPW Syndrome, Septal Preexcitation, Dyssynchrony, and Cardiac Resynchronization Therapy

Novel data released by Ishizu et al does not change the electrophysiology laboratory’s daily practice. Indeed, invasive electrophysiology is the gold standard to accurately identify and treat the AP site. Echocardiographic techniques identify...
the contractile abnormalities associated with AP, but not the exact site for ablation. However, the results of the present study may be considered alongside recent reports demonstrating that early septal activation associated with septal AP may induce reduced IVS motion, as encountered in patients with native or RV pacing-induced LBBB. The development of LBBB-mediated LV dysfunction responsive to CRT has been well documented. Reduced IVS motion and LV dyssynchrony associated with septal AP may result in LV dysfunction, which resolves after radiofrequency ablation, even in the absence of permanent tachycardia. In both septal AP (Figure A) and LBBB (Figure D), activation proceeds across the septum at the breakthrough point before reaching the LV endocardium. It then slowly propagates to the endocardium of the posterolateral wall, with a basal breakthrough site in the event of an AP and more often a mid to apical septal one in the event of an LBBB. This activation delay correlates with a specific LV contractile pattern characterized by an early rapid deformation of the IVS at the breakthrough site, associated with early stretching followed by delayed peak systolic deformation of the posterolateral wall (Figure). Two-dimensional ST longitudinal strain identified a similar contractile pattern in the event of IVS AP (Figure B and C) and LBBB (Figure D and F). Interestingly, the abnormal septal motion associated with this contractile pattern is more frequently found in patients with a prolonged QRS duration in both WPW and LBBB patients. In patients with an LBBB and heart failure receiving CRT, this classical LBBB longitudinal strain pattern has been linked to the occurrence of LV reverse remodeling, improvement in LV function, and clinical outcome, even in patients with relatively narrow QRS (120–150 ms) in whom the benefit of CRT remains uncertain. Given the excellent accuracy of AI to detect and image the contractile consequences of this activation delay in the WPW syndrome, AI may be a suitable and clinically useful tool to identify the contractile substrate of CRT response and, therefore, patients who will derive benefit from CRT. However, this hypothesis deserves further specific research.

Acknowledgments
I am indebted to Raphaëlle-Ashley Guerbaai, RN, MSc, for editing assistance.

Disclosures
None.

References

Keywords: Editorial(s) • cardiac resynchronization therapy • echocardiography • electrophysiology • speckle-tracking echocardiography • Wolff-Parkinson-White syndrome
The Wolff–Parkinson–White Syndrome: A Test Bed for the Assessment of Myocardial Dyssynchrony?
Sylvestre Maréchaux

Circ Cardiovasc Imaging. 2016;9:
doi: 10.1161/CIRCIMAGING.116.005112
Circulation: Cardiovascular Imaging is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2016 American Heart Association, Inc. All rights reserved.
Print ISSN: 1941-9651. Online ISSN: 1942-0080

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://circimaging.ahajournals.org/content/9/6/e005112

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published in Circulation: Cardiovascular Imaging can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial Office. Once the online version of the published article for which permission is being requested is located, click Request Permissions in the middle column of the Web page under Services. Further information about this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to Circulation: Cardiovascular Imaging is online at:
http://circimaging.ahajournals.org//subscriptions/