Point density exclusion mapping—A useful tool for mapping arrhythmias arising from the endocavitary structures

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1 INTRODUCTION

Intracardiac echocardiography (ICE) is a useful modality for imaging endocavitary structures of the ventricles and has become the standard accessory tool while ablating arrhythmias arising from these structures. However, this is not freely available in most of the electrophysiological (EP) labs globally and significantly adds to the procedural cost thus hampering its utility. Point density exclusion (PDX) mapping, a recently described simple technique, helps to visualize the endocavitary structures like the papillary muscle (PM), needs no additional tools apart from high density geometry and activation mapping, and can be done using any electroanatomical mapping (EAM) system. The following report illustrates the use of PDX mapping in precise mapping and successful ablation of a ventricular tachycardia (VT) arising from the PM (PM-VT).

2 CASE DETAILS

A 48-year-old gentleman with mitral valve prolapse and moderate mitral regurgitation had recurrent episodes of VT with one of the episodes needing electrical cardioversion. His cardiac magnetic resonance imaging showed no myocardial enhancement and normal biventricular function. He was planned for electrophysiological study and ablation for the VT. The electrocardiogram suggested a VT arising from the posteromedial PM and the procedure was done using the Ensite Precision system (Abbott, Abbott Park,55 Illinois) for EAM. The basal intracardiac intervals were normal, and the tachycardia was consistently inducible by burst pacing from the right ventricular apex but not with the extra-stimulus protocol. The left ventricular (LV) geometry was created using the Advisor HD grid mapping catheter (Abbott, Abbott Park,55 Illinois), and the activation map revealed a focal mechanism with a centrifugal spread of activation from the site of the earliest activation in the mid-level of the posteroseptal LV aspect. The local electrogram here had high frequency pre-potentials that preceded the surface QRS by 30 ms, and there was an excellent (92%) match during pace-mapping from this region (Figure 1). The Advisor HD grid was exchanged for a 4 mm irrigation-tipped Flexibility catheter (Abbott, Abbott Park,55 Illinois), and activation and pace mapping were done to reconfirm the site of origin. Subsequently, point density exclusion (PDX) technique was applied to display the geometry of the PM and the earliest activation was observed to be arising from this structure. Radiofrequency ablation was performed at this site and after a few
runs of automaticity, the VT became non-inducible. A transthoracic echocardiogram performed with the ablation catheter in situ confirmed its location on the posteromedial PM. The procedure time was 127 min, and the fluoroscopic time was 6 minutes and 27 seconds.

### 3 | PDX MAPPING

The PDX mapping has several advantages over ICE-guided imaging in EP procedures. Firstly, the acquisition of the geometrical points and activation mapping from the endocavitary structure is simultaneous and from the same catheter unlike in ICE. Secondly, there is no need of any additional hardware/software for employing this technique apart from the conventional EAM, thus improving the feasibility and reducing the procedural cost so that it can potentially be applied in any EP lab with EAM facility. Thirdly, the procedural time is less than when using ICE for imaging/image integration. Fourthly, the technique is platform neutral and can be used with any of the EAM systems.

The use of a contact force sensing catheter for ensuring a good contact with the endocavitary structure, though preferable, is not mandatory as illustrated in the current case. The near-perpendicular bending of the electrode grid and the shaft of the Advisor HD grid catheter indicates good tissue contact, and only such sites were used to derive the PDX mapping in our case.

However, there are also a few potential concerns while using PDX mapping as a replacement for ICE guided ablation. The catheter stability needs to be ensured as the PDX mapping will not be able to detect catheter movements and the resultant inaccuracy. The utility of ICE for visualizing lesion formation, and as a guide for visualizing structures like the interatrial septum, pulmonary veins, and detecting pericardial effusion obviously cannot be replaced by PDX mapping. Also, PDX mapping is a post-acquisition processing technique that is applied after obtaining the entire geometrical and the activation points and is heavily dependent on a skillful technical assistant.

### 4 | THE TECHNIQUE FOR PDX MAPPING

In the standard EAM, the inner geometrical points are automatically excluded as more outer points are added as the mapping catheter roves. Thus, endocavitary structures would be excluded from the final representation of the geometry. In PDX mapping, these internal points and the corresponding activation timing are manually included after the acquisition of a high-density geometry and thus the visualization of the endocavitary structures is made possible. For ensuring accuracy, including only the inner points with a contact force of at least 8 gms is recommended though other surrogates may be used as in the current case.

After acquisition of a high-density EAM, the surface fill is reduced to 10% so that the "negative space", which represents the inner points originally excluded from the map, is seen (Figure 2). This is due to the inner points and represents the presence of an endocavitary structure within the shell. The surface fill is akin to the interpolation used in the activation map, and reducing it increases the resolution of the geometry. Adjusting the translucency allows to see the local activation points (and not the surface activation points) with early local activation that surround the negative space. Geometry points in these sites are then copied to a new surface model that corresponds to these internal structures, and the same activation map is applied to obtain a display of activation of the endocavitary structures.¹
CONFLICTS OF INTERESTS
None of the authors have any competing interests or conflicts of interests to declare.

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