Quantification of fiber orientation in the canine atrial pacemaker complex using optical coherence tomography

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Abstract The atrial pacemaker complex is responsible for the initiation and early propagation of cardiac impulses. Optical coherence tomography (OCT), a nondestructive imaging modality with spatial resolutions of ~1 to 15 μm, can be used to identify unique fiber orientation patterns in this region of the heart. Functionally characterized canine sinoatrial nodes (SAN) (n = 7) were imaged using OCT up to ~1 mm below the endocardial tissue surface. OCT images were directly compared to their corresponding histological sections. Fiber orientation patterns unique to the cri
ta terminals (CT), SAN, and surrounding atrial myocardium were identified with dominant average fiber angles of 89 ± 12 deg, 110 ± 16 deg, and 95 ± 35 deg, respectively. Both the CT and surrounding atrial myocardium displayed predominantly unidirectionally based fiber orientation patterns within each specimen, whereas the SAN displayed an increased amount of fiber disarray manifested quantitatively as a significantly greater standard deviation in fiber angle distribution within specimens [33 ± 7 deg versus 23 ± 5 deg, atrium (p = 0.02); 18 ± 3 deg, CT (p = 0.0003)]. We also identified unique, local patterns of fiber orientation specific to the functionally characterized block zone. We demonstrate the ability of OCT in detecting components of the atrial pacemaker complex which are intimately involved in both normal and abnormal cardiac conduction. © 2012 Society of Photo-Optical Instrumentation Engineers (SPIE).

Keywords: conduction; myocardium; structure; imaging; arrhythmia.

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1 Introduction

The sinoatrial node (SAN) is a complex, heterogenous structure located in the upper, posterior region of the right atrium (RA) responsible for initiating cardiac impulses during normal conduction. Structural investigations of the SAN have revealed it to be a diffuse, cigar-shaped structure extending between the superior and inferior vena cavae,1 and electrically isolated from the remaining bulk of atrial tissue.2-4 The location of the leading pacemaker within the larger SAN is highly variable and can be affected by both vagal tone and changes in extracellular ion concentrations.5,6 Early experiments investigating the mechanism of impulse formation and propagation through the SAN have pointed to the presence of discrete exit pathways,7,8 which have more recently been functionally and anatomically identified in both the canine and human SAN.9,10

Atrial arrhythmias, such as flutter and fibrillation, have been shown to involve both the SAN and its exit pathways.11 These arrhythmias can be diagnosed during routine electrophysiology studies and potentially eradicated by catheter ablation procedures. Successful procedural outcomes oftentimes depend on the precise in situ identification of the atrial pacemaker complex, as well as normal/abnormal atrial myocardium. Current clinical anatomical visualization techniques, including magnetic resonance imaging (MRI), computed tomography (CT), and ultrasound, deliver relatively low spatial resolutions and cannot be performed in real time during electrophysiology procedures.

Optical coherence tomography (OCT), on the other hand, is an imaging technique, allowing for the visualization of micron-scale structures within nontransparent biological tissues, with resolutions ranging from submicrometer through 15 μm.12,13 Despite OCT’s high spatial resolution, however, it is currently limited by its depth penetration, estimated to be 2 to 3 mm in biological tissues13 and recently shown to be 1 to 2 mm in human cardiac tissue.14 Furthermore, OCT can be used in situ for real-time, high-resolution visualization of cardiac structures with catheter-based imaging systems.15,16

The use of OCT in visualizing components of the human heart vital to both normal and abnormal conduction, including high-resolution identification of the SAN, was evaluated in our recent study.14 Additional previous studies have shown the utility of OCT in imaging components of the conduction system in both rabbit17,18 and human hearts.14 We have also explored the relationship between structure, imaged using OCT, and function, visualized using optical mapping, in the rabbit SAN.19 The rabbit SAN, however, is essentially a two-dimensional structure,20 as opposed to the three-dimensional canine7,9 and human10,21 SANs. In this study, we aimed to apply ex vivo OCT imaging to canine SANs, which had been functionally characterized via optical mapping as published in our previous study.1 It is important to note that OCT requires highly reflective tissue surfaces, hence it cannot be used to quantify local fiber orientation in these areas important in the generation and maintenance of both normal and abnormal cardiac conduction.

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2 Materials and Methods

2.1 Experimental Preparation

The experimental protocol was approved by the Institutional Animal Care and Use Committee of Washington University. Canine right atrial preparations (n = 7) were initially isolated and functionally characterized via optical mapping with a voltage-sensitive dye. Functional data describing the activation of the SAN and the presence of discrete exit pathways was published in our previous study.9 Immediately after the optical mapping experiment, the tissue preparations were perfused and fixed with 3.7% formaldehyde overnight and then transferred to 20% sucrose for 2 days. Specimens were dissected to include the right atrial appendage (RAA), crista terminalis (CT), and 2 to 3 cm of tissue beyond the CT, where the SAN is located.5, 22 The tissues were photographed and clearly marked with pins to correlate OCT images with corresponding structural and functional datasets. It should be noted that the seven preparations imaged via OCT in this study were initially optically mapped and the electrophysiological results published earlier.9

2.2 OCT Imaging

The basics of our OCT system have been described previously.14,19,23 In short, we imaged each specimen (n = 7) in 4.5 × 4.5 mm overlapping segments from the endocardial surface, with final axial and lateral resolutions of 10 μm (in air).

2.3 Histology and Immunohistochemistry

After imaging, tissue specimens (n = 5) were embedded and frozen in Tissue-Tek® OCT compound (Electron Microscopy Sciences, Hatfield, PA) and stored at −80°C. The tissues were then cryosectioned at 16 μm (n = 4, parallel to endocardium; n = 1, perpendicular to endocardium) and mounted on Superfrost Plus glass slides (Fisher Scientific). Sister sections were then stained using either Masson trichrome (IMEB, San Marcos, CA) or antibodies for the gap junctional protein connexin 43 (Sigma-Aldrich, St. Louis, MO), the cardiac myocyte marker-α-actinin (Sigma-Aldrich, St. Louis, MO), and the fibroblast marker vimentin (Progen Biotechnik, Heidelberg, Germany), as described previously.24 Imaging of both histology and immunohistochemistry was performed on a Nikon C1/S8i confocal microscope with an attached AxioCam MRc5 brightfield camera (Carl Zeiss).

2.4 OCT Image Processing and Analysis

OCT images were processed using MATLAB (Mathworks), as described previously.14 Individual OCT images were acquired six times, and then averaged to reduce background noise. Median filtering was also performed with a 3 × 3 smoothing kernel to further diminish speckle noise. Custom-designed analysis programs were developed in MATLAB for virtual tissue slicing in order to visualize fiber orientation in relation to the tissue surface,19 as well as to quantify the depth of OCT imaging.14 Quantification of 2D fiber orientation was performed using an automated gradient-based algorithm developed and validated in cardiac tissue.25 In short, this method initially calculated the local gradient of each pixel the image of interest. Then, within a small local window of the largest image, the dominant local direction of the gradient was determined. Cardiac fiber angles were assigned perpendicular to this gradient. Areas of interest from which fiber orientation was assessed included the CT, atrial myocardium, SAN, block zone, and exit pathway(s). These areas were identified via a combination of histology, immunohistochemistry, and functional datasets. The SAN, however, was only identifiable in six out of seven hearts imaged via OCT.

Quantitative data are expressed as mean ± standard deviation. A Student’s t-test was used to determine statistical significance. A p-value of <0.05 was considered statistically significant.

3 Results

3.1 Identification of the Atrial Pacemaker Complex

OCT imaging of the canine atrial pacemaker complex achieved an average depth penetration of 1.28 ± 0.23 mm from the endocardial surface across all specimens. The approximate thickness of tissue incorporating and surrounding the atrial pacemaker complex was 2 to 3 mm. Although the SAN is defined as a primarily subepicardial structure,5 inferior portions of the node are located closer to the endocardial surface bringing them within range for OCT.19 Consequently, although OCT could not consistently image throughout the entire thickness of tissue in the area of the SAN, the depth penetration was such that key components of the atrial pacemaker complex could be visualized. OCT failed, however, at identifying nodal tissue in one heart out of seven total imaged.

The initial functional and structural identification of the atrial pacemaker complex was achieved through the merging of several data sets as shown in Fig. 1. The primary field of view for all imaging modalities encompassed tissue ranging from the crista terminalis (CT) to the interatrial septum (IAS) [Fig. 1(a)], which incorporated both nodal tissue and normal atrial myocardium into each data set. Optical mapping revealed the location of the primary pacemaker [Fig. 1(b)], the block zone, and two (or more) exit pathways from the SAN.9 In addition, both histology [Fig. 1(c)] and immunofluorescent labeling for connexin 43 [Fig. 1(d)] pinpointed the location of the SAN, as distinguished from surrounding normal atrial myocardium.

Resolution-wise, OCT is comparable to conventional histological analyses,12-14 which allows for the direct comparison of both techniques in identifying structures within the myocardium. Figure 2 illustrates this concept from two overlying fields of view. The images shown in Fig. 2(a) and 2(b) are located approximately 0.44 mm below the endocardial surface, whereas those images shown in Fig. 2(c) and 2(d) are located at a depth of approximately 0.69 mm. Major identifying structures, such as the sinus node artery and a pinhole (as indicated by the red arrowheads), are apparent using both conventional and virtual histology. In addition, abrupt changes in fiber orientation and organization can be identified and confirmed with OCT, as indicated by the arrows in Fig. 2.

3.2 Quantification of Fiber Orientation

Fiber orientation, as related to impulse propagation in the atrial pacemaker complex, was assessed from three fields of view in each heart. Figure 3 shows typical fiber orientation patterns from areas identified as the crista terminalis, as indicated by the green boxes in Fig. 3(b) and 3(c), and the interatrial septum, as indicated by the blue boxes in Fig. 3(d) and 3(e), at two depths (0.17 and 0.49 mm) below the endocardial tissue surface. As
is evident in the OCT images, these areas of tissue exhibit strong unidirectional fiber orientation with dominant fiber angles of $70 \pm 20$ deg and $137 \pm 16$ deg as measured in the crista terminalis and interatrial septum, respectively.

OCT fields of view encompassing nodal tissue, on the other hand, exhibit very different orientation patterns as shown in Fig. 4 and Video 1. Initially, immediately under the endocardial tissue surface [Fig. 4(b), depth = 0.17 mm], there is a tendency for a strong unidirectional fiber orientation with a dominant angle of $130 \pm 14$ deg, similar to the interatrial septum. As the field of view progresses from the endocardium towards the epicardium and through nodal tissue, the cardiac fibers display an increased amount of disarray as shown in Fig. 4(e) through 4(g). This fiber disarray is representative of SA nodal tissue, which propagates impulses at slower conduction velocities than normal atrial myocardium.

Specific fiber orientation analyses, as shown in Fig. 5 in a single preparation, illustrate the changing orientation and fiber angle distributions in the CT, endocardial atrial layer above the SAN, and the SAN itself. Both the CT and endocardial atrial tissue layers exhibit strong unidirectional fiber orientations with dominant angles centered around 69 deg and 130 deg as shown in Fig. 5(a) through 5(f). Nodal tissue, as verified via standard histology and immunohistochemistry, exhibits a higher degree of fiber disarray as reflected in a wider range of fiber angles ($112 \pm 34$ deg), as shown in Fig. 5(g) through 5(i). Overall average fiber orientation among all preparations, as summarized in Fig. 6, was consistent across samples within the CT (89 ± 12 deg), however, the endocardial atrial layer located above the SAN exhibited a wide range of dominant fiber angles (36 to 128 deg) between specimens. Within each specimen there was still a strong tendency for the endocardial atrial layer above the SAN to exhibit unidirectional fiber orientation as in Fig. 5(d) through 5(f). The overall average angle of
the SAN itself was relatively consistent among the specimens at 110 ± 16 deg, however each individual specimen did exhibit a high degree of fiber disarray within the node, such as that illustrated in Fig. 5(g) through 5(i). The amount of fiber disarray was evaluated based on the standard deviation of angles within each structure (CT, endocardial atrial layer, and SAN) as shown in Fig. 7 among all preparations. Overall, the SAN demonstrated significantly higher standard deviations (33 ± 7 deg) than both the endocardial atrial layer (23 ± 5 deg, $p = 0.02$) and the CT (18 ± 3 deg, $p = 0.0003$) indicative its higher degree of fiber disarray.

### 3.3 Additional Identifiable Features of the Atrial Pacemaker Complex

In addition to evaluating fiber orientation in the CT, endocardial atrium, and SAN, fibers were also identified with unique orientation patterns in the block zone $^{26}$ as shown in Fig. 8 and Video 2. The bright white area, as indicated by the white arrow in Fig. 8(c), most likely contributes to the functional block zone, as shown in Fig. 8(a). The pattern of fibers indicated by the red arrow in Fig. 8(e) appear to “dead-end” in the block zone. Consequently, an impulse originating from the...
SAN would not be able to traverse these fibers and would instead have to bypass the block zone, as indicated by the dotted white line in Fig. 8(e), in order to excite the remaining bulk of atrial tissue.

4 Discussion

It is well known that cardiac function is intimately linked to underlying tissue structure. In this study, we had the distinct advantage of performing structurally based imaging on electrophysiologically characterized preparations. We focused specifically on the structure-function relationship in the canine SAN with particular emphasis on fiber orientations patterns as they relate to SA nodal conduction. Our previous functional study demonstrated the presence of the block zone, nodal tissue, and discrete exit pathways in the canine heart. In addition, these structures were also identified using more traditional histological and immunohistochemical techniques. Now for the first time, we have demonstrated the existence of these structures using a nondestructive imaging modality.

We have identified unique fiber orientation patterns specific to the CT (Fig. 3), SAN (Fig. 4), atrial myocardium (Figs. 3 and 4), and block zone (Fig. 8). In addition, we quantified fiber orientation distributions, using an automated gradient-based algorithm, in these areas (Fig. 5). We found that the CT and atrial myocardium tended to display unidirectionally based fiber orientations, whereas the SAN exhibited an increased amount of fiber disarray manifested as a significantly greater standard deviation (Fig. 7). Although we were unsuccessful in concretely identifying the exit pathways using OCT, we believe their presence may be masked by surrounding fibers with similar orientation patterns. We were, most importantly, able to distinguish SA nodal tissue from the surrounding atrial myocardium (Figs. 4, 5, and 7).

In addition to being a key component of normal cardiac conduction, the atrial pacemaker complex, consisting of the SAN, block zone, exit pathway(s), and surrounding atrial myocardium, has also been shown to be involved in arrhythmias, such as atrial flutter and fibrillation. These arrhythmias are oftentimes associated with SA nodal dysfunction and, more importantly, are accompanied by both functional and structural

Fig. 6 Average fiber angle. Average fiber angles across all hearts in three locations: the atrium (endocardial to the SAN), SAN, and CT.

Fig. 7 Standard deviation of the average fiber angle. Standard deviation of the average fiber angle as an indicator for fiber disarray across all hearts in three locations: the atrium (endocardial to the SAN), SAN, and CT.

Fig. 8 Identification of the block zone with OCT. (a) Photograph of an SA nodal preparation as viewed from the endocardium with the OCT field of view for the SA nodal block zone identified by the white box. OCT images of the fiber orientation are shown at (b) 0.07 mm, (c) 0.12 mm, (d) 0.18 mm, (e) 0.24 mm, (f) 0.32 mm, and (g) 0.42 mm below the endocardium. The white arrow in (c) indicates the probable anatomic correlate of the functional block zone. The dotted white lines indicate the pathways for impulse propagation around the block zone. The red arrow indicates fibers that “dead-end” in the functional block zone. Abbreviations: SVC, superior vena cava; IVC, inferior vena cava; RAA, right atrial appendage; IAS, interatrial septum. (Video 2, MOV, 5.9 MB) [URL: http://dx.doi.org/10.1117/1.JBO.17.7.071309.2]
changes in the SAN. Structural changes, such as alterations in interstitial fibrosis, within the SAN could potentially be monitored with high spatial resolution over time with OCT, as an indicator of disease progression. In addition, an increased amount of fibrosis or abrupt changes in fiber orientation patterns may result in a higher propensity for reentrant arrhythmias.

Recently, it has been suggested that there is a unique structure running parallel to the human sinoatrial node termed the paranodal area. This region was shown to have unique molecular and histological characteristics in the human heart, with properties intermediate between the SAN and normal right atrial myocardium. In our study, we were unable to identify this area in the canine heart using both traditional histology and OCT, despite the functional similarities we have found between the human and canine SA nodes. The identification of the paranodal area in the human heart via OCT still remains to be explored.

OCT is a nondestructive imaging technique with superior spatial resolution as compared to other anatomical visualization and reconstruction techniques. Most importantly, OCT can be miniaturized into a catheter-based imaging system to be threaded through the vasculature and into the cardiac chambers. In this way, physicians can directly relate, in real time and in situ, cardiac function to its underlying structure during electrophysiology studies and ablation procedures. In addition, the continued development of automated visualization and analysis tools for OCT will only serve to power its movement into mainstream cardiology.

5 Conclusion

We have demonstrated the ability of OCT in detecting components of the atrial pacemaker complex, which are intimately involved in both normal and abnormal cardiac conduction. We identified and quantified unique fiber orientation patterns specific to the CT, SAN, atrial myocardium, and block zone using an automated gradient-based algorithm. Based on this study, we believe OCT is continuing to emerge as a key imaging modality with the potential to identify, in real time and in situ, structural substrates responsible for both normal and abnormal cardiac conduction.

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