A Method for Catheter Navigation within Realistic Three-Dimensional Cardiac Anatomy

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Abstract

We recently combined noncontact mapping and intracardiac echocardiography (ICE) in a single cathetersystem that permitted detailed, single-beat, 3-D electricalanatomical imaging of the heart. In the present study we developed a nonfluoroscopic electrode-catheter localization method and integrated it with noncontact electrical-anatomical imaging. The difference in nonfluoroscopic electrode locations compared to ICE was 6.3±3.4 mm, while error in reproducibility of the nonfluoroscopic method was 0.7±1.0 mm. Therefore, accurate and reproducible nonfluoroscopic localization and navigation of standard electrode-catheters within true and detailed 3-D images of cardiac anatomy is feasible. This approach may facilitate diagnosis of arrhythmias and advance their therapy.

1. Introduction

Radiofrequency catheter ablation is widely used for treating several types of cardiac arrhythmias. In order to guide the ablation catheter to regions of interest inside the heart and to deliver the therapy successfully, visualizing the underlying anatomy and accurate three-dimensional positioning of electrode-catheters at specific locations on the endocardium are necessary.

The disadvantages of routine fluoroscopy during catheterization include radiation effects, prolonged procedure duration, limited three-dimensional localization of the catheter, and poor reproducibility. New systems achieve accurate and reproducible three-dimensional positioning of electrode-catheters by employing specialized position sensing mechanisms in the catheter [1-4]. However, present systems do not provide capabilities for imaging the cardiac anatomy and assumptions must often be made in regards to electrode locations with respect to underlying anatomical structures.

We recently developed a novel cardiac catheter imaging system and tested it in the canine intact beating heart [5]. The system was the first to efficiently integrate noncontact electrical and anatomical imaging by combining a custom multielectrode probe and a standard intracardiac echocardiography (ICE) catheter. This noncontact imaging technique accurately reconstructed single-beat endocardial surface electrograms from measured probe electrograms, provided detailed images of cardiac anatomy based on ICE, and digitally fused both modalities to produce realistic, three-dimensional electrical-anatomical images of the beating heart. The objective of the present study was to develop a nonfluoroscopic electrode-catheter positioning method that could be integrated with noncontact electrical-anatomical catheter imaging.

2. Methods

2.1. Experimental model

A catheter-system, comprised of a 9-F sheath carrying a coaxial noncontact 64-electrode lumen-probe (outer diameter = 7 mm) on the outside and a central ICE catheter (9 F, 9 MHz) on the inside, was placed inside the heart cavity. Standard, contact electrode-catheters were fixed at various endocardial sites in the LV of 3 dogs and in the RA of 2 other dogs (total = 10 different sites). To reconstruct the endocardial anatomy, the ICE catheter was pulled back inside the sheath in 2-mm steps and two-dimensional images were acquired while gating to ECG and respiration. Contact electrode locations and endocardial anatomy were identified by ICE as illustrated in Figure 1. The coaxial noncontact probe was then

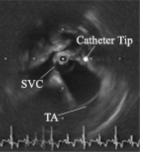


Figure 1. Anatomical imaging in the RA and electrodecatheter localization by ICE.

advanced into the heart cavity by sliding it over the sheath to fixed markers that aligned the probe with the anatomical images. Electrical current pulses (a homing signal) were emitted from the contact tip-electrodes and the generated potentials were simultaneously sensed by all probe electrodes.

2.2. **Mathematical model**

The probe is comprised of electrodes with coordinates (x_i, y_i, z_i) (i=1,2,...,n), where n is the number of probe electrodes. Probe electrodes inside the cavity represent field points and their layout is known in advance. The source point where current is injected (x, y, z) is the contact electrode at the endocardium. The aim of the localization algorithm is to determine the position of the source point (x, y, z).

Let the measured probe potentials be represented by vector $V_m = [V_{m1}, V_{m2}, ..., V_{mn}]^T$, where each V_{mi} (i=1,2,...,n) corresponds to potential on each probe electrode. According to the coordinates of the source point and the field points, V_i can be computed using

$$V_i = \frac{I_s}{4\pi\sigma r_i}$$
 (i=1,2,...,n) (1)

where I_s is the injected location current, σ is the conductivity of the blood, and r_i is the distance between the source point (x, y, z) and the *i*th field point $(r_i = \sqrt{(x-x_i)^2 + (y-y_i)^2 + (z-z_i)^2}).$

Let the calculated potentials be represented by $V_c = [V_{c1}, V_{c2}, ..., V_{cn}]^T$, where each V_{ci} (i=1,2,...,n) is a function of source point position p = (x, y, z). That is, V_c can be expressed as

$$V_c = F(\overline{p}) \tag{2}$$

The process of localizing the contact electrodes is an inverse problem. Here, we used the least-squares method by minimizing the following function

$$R(\overline{p}) = \frac{1}{2} \sum_{i=1}^{n} [F(\overline{p}) - V_m]^2$$

$$= \frac{1}{2} [F(\overline{p}) - V_m]^T [F(\overline{p}) - V_m]$$
(3)

In the above equation, the iterative formula for computing p follows

$$\overline{p}_{k+1} = \overline{p}_k + \Delta \overline{p}_k \tag{4}$$

where p_{k+1} and p_k are localization results at the (k+1)th and kth iterative computation respectively, and

$$\overline{\Delta p_k} = -[[F'(\overline{p_k})]^T [F'(\overline{p_k})] + \lambda I]^{-1} [F'(\overline{p_k})]^T [F(\overline{p_k}) - V_m]$$
 (5)

where λ is a regularization parameter determined from the Composite Residual and Smoothing Operator (CRESO) method [6], and I is the identy matrix. From an initial estimate of p, the iterative process above is executed until satisfying $\max(\Delta V) \le \varepsilon$, where $\Delta V = F(\overline{p}_k) - V_m$ and ε is a very small positive real number.

3. Results

Important anatomical structures were visualized in situ during ICE catheter pullback and without fluoroscopy. In the RA, ICE depicted features such as the appendage, crista terminalis, tricuspid annulus, interatrial septum, fossa ovalis, and coronary sinus ostium. In addition, ICE revealed anatomical structures outside the RA such as other cardiac cavities, pulmonary veins, and the aortic valve. Similarly, in the LV, ICE determined the orientation of the LV walls and depicted the anterior and posterior papillary muscles, the interventricular septum, the right ventricle, and the mitral annulus. Threedimensional anatomical images were successfully constructed based on multiple ICE images throughout the cardiac cycle, as illustrated in Figure 2.

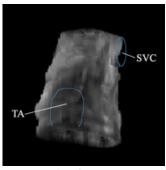


Figure 2. Example of RA geometry by ICE.

3.1. Comparison of measured and computed catheter locations

While injecting the location signal through the endocardial catheter tip, the electrical potential inside the cavity was sensed by the probe electrodes. According to our catheter localization method, all 10 catheter tipelectrodes were nonfluoroscopically localized within the three-dimensional geometry, and were correctly matched with corresponding endocardial anatomy. The difference in nonfluoroscopic tip-electrode locations compared to ICE was 6.3±3.4 mm. One example is shown in Figure 3.

3.2. Reproducibility of catheter location

We tested the reproducibility of our system by selecting two separate current signal cycles, and then

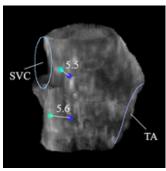


Figure 3. Representative figure depicting the difference between computed and measured locations (in mm).

localizing the catheter tips using each cycle information independently. The positioning error between the two cycles was 0.7 ± 1.0 mm.

4. Discussion

We previously developed a catheter system that combined noncontact cavitary mapping with ICE imaging. This system permitted virtual electrical-anatomical imaging of the heart. In the present study, we developed a method for localizing navigational electrode-catheters inside the cavity in concert with the electrical-anatomical imaging catheter. Unlike present mapping systems, our algorithm permitted localizing electrode-catheters within the confines of true images of the endocardial anatomy readily available by ICE. With this approach, details of the heartbeat can be related to underlying anatomy, and ablation therapy can be delivered at specific sites more effectively.

5. Conclusion

The present study demonstrated the feasibility of localizing electrode catheters within true and detailed three-dimensional images of cardiac anatomy during catheterization. Integrating this approach with noncontact electrical-anatomical imaging may facilitate diagnosis of arrhythmias and advance their therapy.

Acknowledgement

This study was supported by grant HL68768 from the National Heart, Lung, and Blood Institute, National Institutes of Health, Bethesda, Maryland.

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