Effects of the Location of Myocardial Infarction on the Spectral Characteristics of Ventricular Fibrillation

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Background: The location of the myocardial infarction (MI) might modify the spectral characteristics of ventricular fibrillation (VF) in humans.

Objective: To evaluate the effect of the location of the infarcted area on the spectral parameters of VF.

Methods: Patients with chronic MI (29 anterior, 32 inferior) and induced VF during cardioverter defibrillator implant were retrospectively studied. Dominant frequency (f_d), organization index (OI), and power of the harmonic peaks were calculated in the device-stored electrograms (EGM) during sinus rhythm (SR) and VF.

Results: The f_d of the VF was not affected by the left ventricular ejection fraction (LVEF) or the MI location (anterior: 4.54 ± 0.74 Hz, inferior: 4.77 ± 0.48 Hz, n.s.). The OI was also similar in both groups. However, in patients with inferior MIs, normalized peak power at f_d was higher (118.3 ± 18.5 vs 100.6 ± 28.2 , $P < 0.01$) and the normalized peak power of the harmonics was lower than in the anterior MI group. The analysis of EGM during SR showed similar results. The size of the necrotic area and its distance to the recording electrode might partially explain these results.

Conclusion: In our series, the spectral characteristics of the EGMs during VF showed significant differences depending on the MI localization. A higher fraction of energy (in the low-frequency region) was seen in inferior MIs, whereas the peak power at the harmonics increased in anterior MIs. A similar effect was seen during SR and VF, suggesting that it is caused by local electrophysiology abnormalities induced by the MI rather than by different intrinsic characteristics of the VF. (PACE 2008; 31:660–665)

myocardial infarction, ventricular fibrillation, spectral analysis, signal analysis, ICD, electrogram.

Introduction

Electrical activity during ventricular fibrillation (VF) has been considered as a highly disorganized and random process. However, the spectral analysis of optical or electrical recordings during VF reveals a substantial degree of structure and organization, and it allows the estimation of several periodicity and regularity indexes.^{1–3} Spectral VF parameters have been related to the duration of the arrhythmia,⁴ the presence of ischemia,⁵ the action of antiarrhythmic drugs,⁶ or the outcome of shock therapy.⁷ The existence of a necrotic area in the myocardium has been shown to alter the frequency spectrum in a dog model of VF.⁸ Moreover, the maintenance and characteristics of the VF may be related to mother rotors anchored to

specific anatomic structures, such as the papillary muscles, that can be affected by infarcted areas.⁹ Very little information is available on the spectral characteristics of the VF in patients with chronic myocardial infarction (MI). Specifically, it is not known whether the spectrum of this VF depends on the infarct location. The objectives of this study are to describe the characteristics of the electrical signal recorded by an intracardiac electrode during induced VF in patients with chronic MI, and to evaluate the effect of the location of the infarcted area on the VF spectral features.

Methods

Consecutive patients with chronic MI > 3 months old undergoing an implantable cardioverter defibrillator (ICD) implant for clinical reasons between November 2001 and August 2003 in two tertiary hospitals were included in this retrospective study. Clinical data as well as the location of the infarct were obtained from the clinical reports. The left ventricular ejection fraction (LVEF) was categorized in four degrees, from normal (>0.55) to slightly (0.41 to 0.55), moderately

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(0.31 to 0.40), or severely depressed (<0.31). The implant protocol at the time of the study required the induction of at least two consecutive VF episodes in order to ensure a safety margin for defibrillation. The ICD was programmed to record intracardiac electrograms (EGMs) during VF in a pseudounipolar configuration using the defibrillation coil in the right ventricle and the can as the indifferent electrode. The recorded EGMs were saved in digital format (128 samples per second) for further analysis.

VF Signal Processing and Spectral Analysis

The onset of each VF episode was manually annotated, and the following 3 seconds were selected for processing. We decided to analyze the first 3 seconds because this period of time was deemed to be adequate to estimate the spectrum of the signal according to the sampling frequency and the spectral band of interest. An automatic cancellation of low-frequency baseline oscillations (trends) was implemented using a 250-ms median filter and spline interpolation.¹⁰ A spectral representation $P(f)$ was obtained for each EGM segment with a Welch periodogram, with Hamming window of 128 samples, 50% overlap, and spectral resolution of 1,024 samples. This spectral representation was normalized to unit area, which is denoted as $P_n(f)$. In many cases, spectral profile of the monopolar EGMs closely corresponded to a harmonic structure, which was taken into account for using conventional parameters in this setting. These spectral parameters and its meaning have been recently reviewed,¹¹ and they are briefly summarized next (see Fig. 1):

- **Dominant Frequency (f_d):** It is the frequency at which the absolute spectral maximum occurs.
- **Fundamental Frequency (f_0):** For periodic signals, it is the inverse of the cycle length, that is, the inverse of the minimum repetition period.
- **Harmonics and Power Peaks:** For periodic signals, harmonics are the frequencies corresponding to the integer multiples of fundamental frequency f_0 . We considered up to five harmonics, given by $f_2 = 2 f_0$ to $f_5 = 5 f_0$. When the power peak did not happen exactly at the harmonic frequency due to noise, it was adjusted to correspond to the nearest relative maximum, within a tolerance of about 1.5 Hz. Accordingly, we measured the peaks of the normalized power spectrum at each frequency, denoted by $P_n(f_d)$, $P_n(f_0)$, and $P_n(f_2)$ to $P_n(f_5)$. The obtained values were multiplied by a factor of 1,000 in order to make the numerical expressions easier.
- **Organization Index:** We defined the Organization Index (OI) as follows. Given that the EGM spectra were mostly contained in the band from 4

Figure 1. An ICD-stored EGM VF example. (A) Spectral parameters measured from the power spectrum and spectral envelope. (B) Monopolar EGM.

to 30 Hz, we first obtained the total power in this band. Then, for the spectral peaks corresponding to fundamental frequency f_0 and to every harmonic peak within this band, their power in a bandwidth given by the 75% decrease in amplitude in each peak was obtained and summed. Accordingly, the OI was calculated as the ratio between the power in the bandwidth of f_0 and its spectral harmonics, and the total power in the band from 4 to 30 Hz.¹²

- **Spectral Envelope:** In order to yield an independent from periodicity spectral characterization, an auto-regressive model (Yule-Walker method, fourth order) was fitted for each EGM signal during VF, and the squared spectral envelope was estimated, thus yielding a spectral representation that has no harmonic structure, and hence it contains no explicit information about the periodicity, but rather about the underlying

physiological phenomena and about the acquisition characteristics.¹³

SR Signal Processing and Spectral Analysis

In order to evaluate whether the differences in the VF spectrum between the study groups represent a specific behavior of the VF for each infarct location or they just reflect intrinsic local electrophysiologic properties of the myocardium related to the infarcted area, the frequency content of the EGM during sinus rhythm (SR) was also analyzed. For this purpose, we obtained the ICD-stored EGM of SR for each patient. Spectral envelope was also estimated for each SR recording.

Simulations

To reproduce and to explain the results obtained in the previous analysis, a simulation study was conducted. The model has been described in detail elsewhere.¹⁴ In brief, it consists of a cellular automaton model, with activation being governed by a static conventional restitution curve. Additionally, an EGM recording model is given by the volume conductor equation in a homogeneous medium, and it was tuned for modeling a simple unipolar electrode configuration. As shown in Figure 2, a 1×2 -cm cellular tissue sheet was discretized (80 cell groups per cm). We compared homogeneous tissue with the presence of an infarcted, electrically silent region. Simulated EGMs were obtained at two different locations, as described in Figure 2. The infarcted region was considered either to the left, according to the figure, or to the right of the sheet, allowing us to analyze the effect of proximity between the infarct and the electrode. Unipolar EGMs were recorded at $x =$

2.4 cm and $y = 0.5$ cm, at a vertical distance to the plane of 0.2 cm. In order to analyze the effect of distance, an electrode was also modeled at $x = 4$ cm and $y = 0.5$ cm at the same vertical distance.

Results

Sixty-one patients were included in the study, 29 with anterior MI (47.5%, group A) and 32 with inferior MI (52.5%, group B). Age (63.0 ± 9.9 years for group A, 65.2 ± 8.6 years for group B) and gender (one female in each group) were not significantly different between groups. An LVEF $< 30\%$ was observed in 22/29 (75.9%) patients with anterior MI and in 11/32 (34.4%) patients of group B ($P = 0.001$). The spectral parameters were not affected by the degree of left ventricular dysfunction. Specifically, the dominant frequency did not show significant differences between patients with LVEF < 0.30 (4.61 ± 0.69 Hz) and patients with LVEF > 0.30 (4.71 ± 0.55 Hz, n.s.) and similar results were obtained for $P_n(f_d)$ (0.12 ± 0.03 vs 0.11 ± 0.02) and OI (0.79 ± 0.09 vs 0.78 ± 0.07 , respectively, both nonsignificant).

Low correlation coefficients were obtained for the spectral variables (R of 0.47 for f_d , 0.70 for f_0 , 0.01 for $P_n(f_d)$, 0.01 for $P_n(f_2)$, 0.18 for $P_n(f_3)$, 0.22 for $P_n(f_4)$, 0.15 for $P_n(f_5)$, and 0.19 for OI). Therefore, each measured spectral parameter was averaged in the two consecutive VF episodes, when available.

VF Signal Processing and Spectral Analysis

The f_d was similar for patients with anterior and inferior MIs (4.54 ± 0.75 Hz and 4.77 ± 0.49 Hz, respectively, n.s.). The differences in f_0 and OI were also nonsignificant. In contrast, the spectral pattern in groups A and B was significantly different (see Table I). In patients with inferior MI, normalized power at the f_0 peak was significantly higher (0.10 ± 0.02 vs 0.11 ± 0.01 ,

Figure 2. Simulation model. Grey zone represents the infarcted region, and it was considered either to the left, according to the figure, or to the right of the sheet, to analyze the effect of proximity between the infarct and the electrode. Unipolar EGMs were recorded at $x = 2.4$ cm and $y = 0.5$ cm, at a vertical distance to the plane of 0.2 cm. To analyze the effect of distance, an electrode was also modeled at $x = 4$ cm.

Table I.

Spectral Characteristics of the VF Signal in Anterior and Inferior Infarcts

	Anterior	Inferior	P Value
f_d	4.54 ± 0.74	4.77 ± 0.48	n.s.
f_0	4.54 ± 0.75	4.76 ± 0.47	n.s.
$P_n(f_d)$	100.6 ± 28.2	118.3 ± 18.5	0.006
$P_n(f_2)$	20.0 ± 10.3	13.7 ± 7.9	0.01
$P_n(f_3)$	5.4 ± 3.8	2.9 ± 2.5	0.004
$P_n(f_4)$	1.8 ± 1.7	0.9 ± 0.9	0.012
$P_n(f_5)$	0.8 ± 0.9	0.3 ± 0.2	0.003
OI	0.79 ± 0.12	0.80 ± 0.07	n.s.

SPECTRUM OF MYOCARDIAL INFARCTION DURING VF

Figure 3. Averaged spectral envelopes in logarithmic units and crossover about 10 Hz. (A) VF in ICD-stored EGM. (B) SR in ICD-stored EGM. (C) Simulation of changes with distance. (D) Simulations of changes with proximity between electrode and infarction.

$P < 0.01$), whereas the normalized spectral power at the harmonics was higher in patients with anterior MI. The same analysis did not yield any significant difference when made on tip-to-coil configuration available in the ICD-stored EGM, and hence, those results are not included here for shortness. Also, the analysis of the first 4 or 5 seconds gave similar results (not included). The averaged spectra in logarithmic units for both groups are depicted in Figure 3. A crossover of the spectral envelopes is clearly seen, with greater fractional power below 10 Hz approximately in inferior infarctions, and showing a significant, opposite trend above this frequency. The spectral parameters were not affected by the degree of left ventricular dysfunction.

SR Signal Processing and Spectral Analysis

The averaged spectral envelopes in logarithmic units for both groups during SR are shown in Figure 3. A crossover between spectral envelopes at about 10 Hz is again apparent, showing a low-pass filtering effect for inferior MIs, very similar to the one observed in the preceding VF analysis.

This crossover was also obtained using instead an autoregressive spectral analysis (not shown) thus demonstrating that it was not a phenomenon dependent on the signal processing fitting procedure. The use of the same electrode configuration for the acquisition of the EGMs during VF and SR may contribute to explain the similarity of both spectral envelopes as observed in Figs. 3A and B.

Simulations

Possible reasons for this effect were assessed using a simulation model. As shown in Figure 3, the higher frequencies become attenuated as the electrically silent area and the recording electrode come closer, thus moving the electric source away from the electrode. This low-pass filtering effect is a consequence of the electric field characteristics in quasi-electrostatic conditions.¹⁵ The crossover effect, also reproduced by the simulations, is due to the normalization of the area under the spectral curve to one. The increment of the size of the infarcted region has also a low-pass filtering effect: the greater the size of the infarction (and hence, the lesser effective myocardial mass to contribute

to the EGM voltage), the lower the spectral content at high frequency in the unipolar EGM.

Discussion

Our study shows that the spectral characteristics of the electrical signal during induced VF recorded with an intracardiac electrode depend on the location of the infarct. In patients with inferior MIs, the contribution of the low-frequency band to the total energy of the spectrum is relatively higher, whereas the relative amplitude of the harmonics increases in patients with anterior infarcts. This effect may be due to an alteration of the electrophysiologic properties of the myocardium induced by the infarcted area and does not necessarily reflect different intrinsic characteristics of the VF. The activation mathematical model suggests that differences in the distance from the electrode to the infarcted region or in the size of this region could play a role in reducing the fractional energy of the high-frequency band in inferior MIs.

Several experimental studies have shown that the existence of an infarcted area may change the frequency content of local EGMs during VF. In a canine model of chronic MI, the cycle length of VF was longer and the degree of organization greater than in the control group.¹⁶ In a similar experimental preparation, Jacobson et al.⁸ found no significant changes in the f_d or in the fractional peak power at the f_d peak, but demonstrated a significant reduction of the total and peak power in the infarcted animals. Finally, frequency maps of the left ventricle in sheep with chronic anterior MI showed regions of high-frequency activation usually being located close to the interventricular septum, whereas a low f_d was recorded within the infarcted area.¹⁷ However, none of these studies analyze specifically the power distribution along the frequency axis. Moreover, the effect of the infarct location on the VF characteristics cannot be assessed since the infarct is always produced by occlusion of the proximal left anterior descending coronary artery. The distribution of the necrotic area in the myocardium might potentially alter the VF spectrum by several mechanisms. First, it has been suggested that the VF may be induced and maintained by rotors anchored in specific anatomic areas such as the papillary muscles,^{17,18} and the involvement of these areas in the necrotic process would depend on the location of the infarct. Second, the MI may induce changes in the basic electrophysiological properties of the myocardium that surrounds the recording electrode, and this effect could modify the VF spectral characteristics irrespective of the intrinsic properties of the arrhythmia. Our results support this mechanism, since the location of the infarct induces similar changes in the frequency distri-

bution of the EGMs during SR. Specifically, the effect on the power of the low and intermediate frequency bands is very close to the one observed during VF. The differences of the spectral envelope seen in the region above 30 Hz might be due to the higher conduction velocity of the electric impulse during sinus rhythm.

The effects of the location of the infarcted area on the VF spectral pattern have not been evaluated in humans. Our results suggest that patients with inferior MI show a greater fractional power at the f_d component. In contrast, the normalized power at the harmonic frequencies is lower when compared to the anterior MIs. The reasons for these differences are unknown. A similar effect is observed in our two-dimensional computer model of ventricular activation when the distance between the infarcted region and the electrode decreases, or when the size of the simulated infarcted area is increased. Inferior MIs can involve the right ventricle and might be closer to the ICD recording electrode than anterior infarcts, potentially explaining the obtained results. In contrast, a larger amount of electrically active myocardium would be expected in the inferior infarct group, since anterior MIs are more often associated with extensive scars. More precise information on the size and location of the infarcted areas as well as their anatomic relationship with the electrode position might be obtained with image techniques and could be useful to better assess this issue, but we lack this information due to the retrospective character of the study. A second limitation derives from the use of an ICD to obtain the EGMs, which reduces the available signal to a unique intracardiac recording. Multiple recordings from both ventricular chambers during sinus rhythm and VF might be useful to further delineate the influence of the MI location and size on the spectral characteristics of the arrhythmia. Finally, the two-dimensional mathematical model of propagation clearly oversimplifies the phenomena involved in the generation of the EGM in an infarcted human heart. It was used only to simulate the effects of the size of the necrotic region and its distance to the recording electrode on the spectrum of the electrical signal. A more realistic approach would require a substantial increase in the complexity of the model and lies beyond the scope of this study.

Conclusion

The spectral characteristics of the intracardiac signal during VF depend on the MI location. A higher fractional energy in the low-frequency region is seen in inferior MIs, whereas the peak power at the harmonics increases in anterior infarcts. A similar effect is seen during SR and VF, suggesting that it is caused by local

electrophysiologic abnormalities induced by the MI rather than by different intrinsic characteristics of the VF. The mathematical model analysis suggests that this effect could be due either to differences in the position of the electrode relative to the necrotic area or to their separation distance, but further studies considering detailed anatomical information are required to confirm this result. Overall, these results should be taken into account

when analyzing further clinical studies involving patients with infarctions in terms of spectral features of their presenting arrhythmias.

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