Is there a significant transmural gradient in repolarization time in the intact heart?

Cellular Basis of the T Wave
A Century of Controversy

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The ECG is one of the oldest and most versatile noninvasive cardiac diagnostic tests. It has remained in use essentially in its original form despite dramatic advances in cardiac electrophysiology. In May 1887, Augustus Desiré Waller recorded the first human Electrogram using a primitive instrument called a Libbmann capillary electrometer. It had 2 deflections corresponding to ventricular depolarization and repolarization.1 In 1903, Willem Einthoven invented the String Galvanometer—a more sophisticated voltage recording instrument and recorded an Elektrokardiogramm with 5 deflections that he named PQRST.2

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Since its initial invention, the body surface ECG has become a commonly used and extremely valuable test for the diagnosis of a variety of cardiac conditions. Despite a century of prolific use and intensive investigation, the cellular basis of ECG waveforms, particularly the T wave, remains a matter of debate.

Anecdote of the T wave
The saga of the T wave began in 1856, when 2 German physiologists Kölliker and Müller attempted to explore the electric activity of the heart using frog sciatic nerve attached to gastrocnemius muscle as a voltage recording instrument and observed 2 contractions (see review by Noble and Cohen3). In retrospect, the second “contraction” probably reflected a voltage gradient related to the T wave of Einthoven. In 1883, Burdon-Sanderson and Page4 were the first to demonstrate that in the frog’s heart, the wave of excitation spreads from the base to the apex of the ventricle. The record was diphasic, with the first positive (R) wave followed by a negative (T) wave. They also demonstrated that the T wave corresponds to repolarization of the ventricle. Similar series of experiments by Bayliss and Starling in 18925 in the canine heart showed that the T waves are usually upright in mammals. This was followed by Mines6 on frog’s heart using Einthoven’s String Galvanometer in 1913. Mines stated that the T wave is an electric expression of asymmetrical passing off of excitation (repolarization) in different parts of the ventricles and that the polarity of the T wave can be predicted by changing the duration of the action potential in a controlled manner. In 1931, Wilson et al7 put forward the hypothesis that the concordance of the T wave with the R wave on the surface ECG can be explained by assuming that the depolarization and the repolarization waves must travel in opposite directions at least in some parts of the ventricle. The
The development of the left ventricular wedge technique has greatly advanced our understanding of the role of electric heterogeneity in the genesis of the T wave. In 1959, Reynolds and Vander Ark demonstrated that in the canine heart, earlier repolarization of epicardium as compared with endocardium was associated with a positive T wave and vice versa. In 1972 Burgess et al showed that the ventricular functional refractory periods are longer on the endocardium than epicardium and apex as compared with the base. Using the glass microelectrode technique, Solberg et al in 1974 recorded transmembrane action potentials from tissue slices obtained from the canine papillary muscle. They demonstrated a transmural gradient of repolarization with the subendocardial cells having the longest duration of action potential followed by the endocardial and epicardial cells. Similar evidence that earlier repolarization occurs in the canine epicardium was obtained by Spach and Barr and Abildskov in 1975.

In an elegant experimental model of an open chest canine heart, in 1984, Higuchi and Nakaya et al recorded monophasic action potentials from the endocardial and the epicardial surface of the LV with a simultaneous recording of an epicardial-unipolar ECG in close proximity. They found a high degree of correlation between the transmural APD gradient and the polarity of T wave. In their experiments, warming the epicardial surface by dropping warm physiological saline solution led to abbreviation of epicardial APD with a reversal of transmural repolarization gradient, manifested as negative T waves.

This classic work was followed by a study by Franz et al in the human heart. He recorded monophasic action potential from the endocardial and epicardial surfaces of the human heart and showed that epicardial repolarization occurs earlier than the endocardium leading to a transmural repolarization gradient.

**M Cell and the Left Ventricular Wedge Preparation**

The discovery of the M cell (Masonic, Midmyocardial, Moe cell) has significantly advanced our understanding of the transmural voltage gradient. An increasing number of studies have shown that the ventricular myocardium is not a homogeneous structure, but it is comprised of 3 distinct cell types, ie, epicardial, subendocardial and endocardial cells, with clearly distinct electrophysiological and pharmacological properties. The M cells are located in deep subendocardium of the canine LV. The hallmark of M cells is the ability of its action potential to prolong more than that of the epicardium or endocardium in response to slowing of the rate or in response to agents that prolong APD.

The development of the canine and rabbit left ventricular wedge models has greatly advanced our understanding of the role of electric heterogeneity in the genesis of the repolarization waves of the ECG. This experimental preparation provides us with the unique ability to record transmembrane action potentials from 3 different myocardial cell types simultaneously.
together with a pseudo-ECG. Because the pseudo-ECG is recorded by placing 2 electrodes in the Tyrode's solution (ie, a homogenous volume conductor) bathing the wedge preparation 1 to 1.5 cm from the epicardial and endocardial surfaces, the electric field of the preparation as a whole is measured. We use the pseudo-ECG to distinguish it from the body surface ECG, although both recordings are principally similar. The simultaneous recordings of transmembrane action potentials and the pseudo-ECG provides us with ability to correlate changes in APD with those in ECG morphology. Using the wedge preparation, we have demonstrated that differences in the time course of repolarization of the 3 predominant myocardial cell types contribute to electrocardiographic inscription of the T wave in pseudo-ECG.13 As shown in Figure 1, the different time course of repolarization of phases 2 and 3 of the 3 principle myocardial cell types gives rise to opposing voltage gradients on either side of the subendocardial M cells, leading to the inscription of the T wave. In the case of an upright T wave, the epicardial cells repolarize earliest and the M cells repolarize last. Final repolarization of the epicardial cells coincides with the peak of the T wave and final repolarization of the M cells coincides with the end of the T wave.25,26 Thus, the M cell APD approximates the QT interval and the epicardial APD approximates the QTpeak interval. The interval from the peak of T wave to the end of T wave (Tp-e) correlates closely with transmural dispersion of repolarization (TDR). Also, the polarity of the T wave can be changed by changing the APD in a controlled manner.

**Is there an Apico-Basal Voltage Gradient in the LV?**

It has been argued that the canine left ventricular wedge is too small a sample and does not reflect what might happen in an intact heart.27 This prompted us to develop an experimental model where we use the intact rabbit LV instead of a wedge. In this model, we tested the relative role of the transmural versus the apico-basal voltage gradient in the genesis of the T wave. We elected to use the rabbit LV instead of the dog for several reasons. First, as shown in Figure 2A, the isolated rabbit LV is perfused by cannulating the left main coronary artery that allows us to use the complete left ventricular tissue in the experiment and it ensures that the preparation is well perfused. Because it includes the whole ventricle it allows us to record the transmural and the apico-basal voltage gradient simultaneously. Secondly, cells with electric resemblance to the M cells are located in the endocardium of the rabbit LV,28,29 so that it is technically easier to simultaneously record action potentials from the endocardium and epicar-

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**Figure 1.** A. The cellular basis of the T wave in the canine left ventricular wedge preparation. Simultaneous recordings of action potential with of 3 different cell type with a pseudo-ECG are shown. Note that the opposing voltage gradient on either side of the M region leads to inscription of the T wave. The peak of the T wave coincides with the end of repolarization of the epicardial cell and the end of the T wave coincides with the end of repolarization of the M cells. B. The cellular basis of the T wave in the rabbit left ventricular wedge. In the rabbit, the entire endocardium has the same electrophysiological properties as M cells. Therefore, the end of the T wave coincides with the end of repolarization of the endocardium in the rabbit. Epicardial pacing leads to earlier activation of epicardial cells with shorter APD and delays the activation of endocardial cells with a longer APD, leading to an increase in QT and Tp-e interval. Epi, Epicardium; Endo, Endocardium; AP, action potential. Modified and reproduced with permission from Circulation. 1998;98:1928–1936.
The action potential recordings obtained from intact epicardial and endocardial surface eliminates the possibility of injury and uncoupling due to dissection. Some have proposed that transmural dispersion of repolarization seen in the small canine wedge preparation may be related to injury and uncoupling of the cells because the action potential is recorded from a transected subendocardial surface.³⁰,³¹ Realistically the cut surface constitutes only an extremely small fraction of the entire wedge preparation that should have little influence on the pseudo-ECG. In any case, the arterially-perfused rabbit LV is devoid of such a confounding factor.

The arterially perfused rabbit LV was prepared in a similar way as the rabbit left ventricular wedge.²⁸,³²–³⁴ To accomplish this, as shown in Figure 2B, we recorded the endocardial and the epicardial transmembrane action potentials simultaneously from the apical and the basal areas of the rabbit LV. Two pseudo-ECGs were recorded: one recorded transmurally and another across the apico-basal axis of the left ventricular wall. It is important again to emphasize that the pseudo-ECGs were recorded in the volume conductor in which the electrodes were placed at least 2 cm from the epicardial and endocardial surfaces, or from the apex and base of the isolated LV. After control recording at a basic cycle length of 1000 ms, 0.1 μmol/L cisapride at was infused to determine if it has any preferential effects on epicardial/endocardial or apical/basal cell action potential.

Representative recordings from these experiments are shown in Figure 3A. Under control conditions, the APD of apical cells (both endocardial and epicardial surfaces) was slightly longer than the APD of the respective basal cells; however, the difference was not statistically significant. On the other hand, both at the apical and the basal recording sites, the APD of the endocardial cells were consistently longer than the epicardial cells and the results were statistically significant. This difference in the time course of repolarization of the epicardial and endocardial cells, ie, the transmural dispersion of repolarization, caused the inscription of the T wave in the pseudo-ECG recorded across the transmural axis. In both apical and basal recording sites, the peak of the T wave was coincident with the end of repolarization of the epicardium and the end of T wave was coincident with the end of repolarization of the endocardium. In contrast, the pseudo-ECG recorded along the apico-basal axis had T waves that were either flat or negative.

After infusion of the IKr blocker cisapride, the QT interval prolonged with a preferential prolongation of APD of the endocardial cells as compared with the epicardial cells across the entire LV, leading to an increase in TDR and a corresponding increase in the duration of descending limb of the T wave, ie, the Tp-e interval. Cisapride increased the APD of the apical cells slightly more than the basal cells; however the observed difference was not statistically significant. Once again, the apico-basal pseudo-ECG showed nonspecific T wave morphologies and we were unable to identify any correlation between the morphological changes in action potential and the pseudo-ECG. Findings were reproduced in 4 similar experiments. Composite data are shown in Figure 3B.

The canine left ventricular wedge spans 5 cm of the LV across the apico-basal axis. A decade of experience with this preparation confirmed that there is no significant dispersion of repolarization across the apico-basal axis in such a small segment of LV.¹³ Does this hold true in an entire canine LV? This issue was elegantly addressed in recent work by Rosenbaum and coworkers in canine hearts using optical mapping.
In this study investigating the physiological basis of T wave memory, 3 wedge preparations were used from anterior, lateral, and posterior surface of canine LV, respectively. Action potentials were recorded with optical mapping technique from all 3 wedge preparations. The dispersion of repolarization was calculated across the transmural axis of each wedge (ie, TDR) and also between 2 LV wedges (ie, segmental dispersion of repolarization). The transmural pseudo-ECG was calculated by subtracting the action potentials from epicardial cells and M cells. The segmental ECG was calculated by subtracting action potentials of M cells recorded from 2 different LV wedges/segments after accounting for known activation time between segments. The results showed the presence of TDR in all 3 LV wedges, but there was no significant segmental dispersion of repolarization. Both the measured and the calculated transmural Pseudo-ECGs showed upright T wave that matched the polarity of in vivo ECG. However, the calculated segmental ECG failed to register any T wave. These observations outlined above, clearly demonstrate the prominent role of TDR in the electrocardiographic inscription of the T wave, both in the rabbit and canine LV.

![Figure 3](http://circep.ahajournals.org/)

**Figure 3.** A, A representative recording from the rabbit LV showing well-defined T waves in pseudo-ECGs recorded across the transmural axis under control conditions and after infusion of cisapride. Note that the pseudo-ECG recorded across the apico-basal axis fails to reveal any T waves, and there is no correlation between changes in APD and ECG morphology. B, Composite data showing similar findings in 4 similar experiments. Note that APD was slightly longer at the apex as compared with the base; however, the difference is not as pronounced as seen between the endocardium and the epicardium. AP, action potential; Epi, epicardium; Endo, endocardium; APD\(_{90}\), APD at 90% repolarization. **P<0.05 compared between endocardium vs epicardium, ##P<0.05 compared between apex vs base.
Clinical Evidence for Transmural Dispersion of Repolarization

Figure 4 presents a mathematical model in which the ECG was calculated first from a transmural linear array of cells in which repolarization is assumed to be homogeneous and then from a more physiologically-relevant model in which repolarization is heterogeneous. In the homogeneous myocardium, the R wave and T wave would be always of opposite polarity and reversal of direction of activation does not have any effect on the QT interval, QRS duration or Tp-e interval. In such a case, the T wave is merely a reflection of the transmural conduction time. In contrast, a transmural repolarization gradient in the heterogeneous myocardium would produce a positive T wave which has the same polarity as the R wave.

However, clinical observations do not agree with the result of a homogeneous model. Fish et al studied the effect of reversing the direction of activation of the left ventricular wall in patients with heart failure (Figure 5). The epicardium of the left ventricular free wall was paced at cycle length of 600 ms through the epicardial electrode of a resynchronization device followed by pacing from the endocardial aspect of the LV free wall by an endovascular...
catheter at the same cycle length. The effects of epicardial and endocardial pacing on the QT interval and T\textsubscript{p-e} interval was measured in lead V6 on body surface ECG. Lead V6 was selected because it was closest to the site of pacing and because it can record the voltage across the transmural axis of the left ventricular wall. A representative recording from such an experiment is shown in Figure 5A. If the assumption that the myocardium is electrically homogeneous is correct, the R and the T wave should always be of opposite polarity and the reversal of direction of activation of the ventricle should not have any effect on QT and T\textsubscript{p-e} interval. This is clearly not the case (Figure 5A).\textsuperscript{36} Reversing the direction of activation of the left ventricular wall by shifting from endocardial to epicardial stimulation caused an increase in the amplitude and width of the T wave, a prolongation of QT interval and a significant increase on T\textsubscript{p-e}.\textsuperscript{33,37} This behavior is also observed in the canine and rabbit left ventricular wedge preparation\textsuperscript{33,37} (Figure 1) and is predicted by the mathematical model illustrated in Figure 4 to be the result of transmural heterogeneities of repolarization time course.

Do the above findings hold true for the apico-basal voltage gradient? In other words, if apico-basal dispersion of repolarization contributes importantly to the genesis of the T wave, then pacing the ventricle from the apex followed by the base should affect the QT and the T\textsubscript{p-e} interval in a predictable fashion. To answer this question, we analyzed the ECGs obtained during electrophysiological (EP) study in 5 patients in whom the right ventricle was paced from the apex (RVA) and the outflow tract (RVOT) at a basic cycle length of 600 ms. The QT and T\textsubscript{p-e} intervals were calculated during RVA and RVOT pacing. We selected chest lead V2 and limb lead II for evaluation. Chest lead V2 was selected since it measures transmural gradient closest to the site of pacing, and limb lead II was selected because it samples across the apico-basal axis of the heart. As shown in Figure 5B, changing the pacing site from RVA to RVOT did not have any effect on the QT and the T\textsubscript{p-e} interval in lead V2 or lead II. The composite data of 5 patients showed that the QT interval remained unchanged (385±11006\textsubscript{16.4} RVA pacing versus 384±11006\textsubscript{13.7} RVOT pacing, p=NS) as did the T\textsubscript{p-e} interval (85±3.5 RVA pacing versus 84±3.7 RVOT pacing, p=NS). Changing the pacing site from the RVA to RVOT led to the reversal of the polarity of the T wave in the apico-basal axis in lead II. In contrast, the polarity of the T wave in transmural axis lead V2 remained unchanged.

These observations demonstrate the prominent role of TDR over apico-basal dispersion of repolarization in the genesis of T wave. It is important to emphasize that these data do not exclude the presence of apico-basal dispersion of repolarization under some conditions. Several lines of evidence support
apico-basal heterogeneity of repolarization. For example, Bauer et al have demonstrated that in the canine heart in vivo, the effective refractory period is longer in the epicardial muscle layer of the apex than in the base and on administration of dofetilide apico-basal dispersion of repolarization becomes more pronounced with preferential increase in the refractory period at the apex. Similarly, Janse et al reported shortest repolarization times in anterobasal areas and longest repolarization times in posteroapical regions of intact canine LV using unipolar electrograms. Also in enzymatically isolated myocytes from rabbit heart, APD is longer in myocytes isolated from the base as compared with the apex.

Although the apico-basal dispersion of repolarization does exist, such dispersion is much smaller as compared with transmural dispersion. Also, apico-basal dispersion occurs over a much longer distance as compared with transmural dispersion. It is the steepness of the repolarization gradient that is thought to serve as a key factor for arrhythmogenesis and not the total magnitude of dispersion. That is, a small amount of dispersion of repolarization over a long distance will yield a less steep repolarization gradient as in the case of apico-basal repolarization gradient and is less likely to be arrhythmogenic. The transmural dispersion of repolarization is not only more pronounced as compared with apico-basal dispersion, but it also occurs over a much shorter distance resulting in a very steep electric gradient that in our opinion is likely to influence not only the inscription of the T wave but also contribute to arrhythmogenesis.

Conclusions

The presence of an important transmural repolarization gradient in ventricle has been well established in a variety of species, including man. Although apico-basal voltage gradients may contribute, we believe that the transmural voltage gradient plays a predominant role in the genesis of the T wave. The wedge preparation, like any animal model, has limitations, but it has a remarkable ability to reproduce the electrocardiographic and arrhythmic manifestations of a wide variety of syndromes observed in clinical practice. This model coupled with in vivo data from humans and mathematical simulations of the myocardium provide compelling evidence in support of a predominant role for the transmural voltage gradient in the genesis of the T wave. Dispersion of repolarization, irrespective of its precise geometry, has been shown to serve as the substrate for reentry under a variety of conditions. It is clear that the T wave corresponds to repolarization of the ventricle and it is an electric manifestation of a normally asymmetrical repolarization process. An important electrocardiographic parameter obtained from body surface ECG, the T_{pe} interval, has been shown to be a predictor of arrhythmias under a variety of clinical conditions.

Although the exact mechanism for the genesis of the T_{pe} interval on the surface ECG remains a matter of some controversy, its clinical utility as an index of arrhythmogenesis is gaining recognition.

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Disclosure

None.

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Response to Patel et al

Tobias Ophoth, PhD, Ruben Coronel, MD, PhD, Michiel J. Janse, MD, PhD

Reductionist models are justified when smaller preparations reproduce the characteristics of the intact organ. The wedge preparation of the left ventricular free wall described by Patel et al is a good example of the pitfalls of reductionism. Although it is tailored to study transmural differences (because it uncovers a plane that is normally not accessible), it reduces other heterogeneities. Figure 3 in the article by Patel et al shows pseudotransmural and apico-basal ECGs of a rabbit wedge preparation in relation to apico-basal and transmural gradients in repolarization. The authors emphasize that the endocardial-epicardial gradient in repolarization translates into the $T_{p-e}$ interval of the transmural ECG but not of the apico-basal ECG. The latter, however, is clinically more relevant, and its T wave does correlate with the apico-basal repolarization gradient. Thus, even when a relevant endocardial-epicardial gradient in repolarization would be present, which is not the case in the whole heart (see Figures 1 and 2 in our article), it is simply invisible in a clinically relevant ECG. We agree with our colleagues that the $T_{p-e}$ interval relates to arrhythmogenesis, as would probably each segment of a prolonged T wave, but differ in opinion about what the $T_{p-e}$ interval reflects. Figure 1 (right panel) in our article shows that $T_p$ correlates with the yellow area, whereas $T_e$ correlates with the blue area. Therefore, the $T_{p-e}$ interval indicates dispersion in repolarization of the left ventricle as a whole.

KEY WORDS: T wave ■ cellular basis ■ transmural dispersion of repolarization ■ apico-basal dispersion of repolarization ■ controversy
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