Error Analysis of Tissue Resistivity Measurement

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Abstract—We identified the error sources in a system for measuring tissue resistivity at eight frequencies from 1 Hz to 1 MHz using the four-terminal method. We expressed the measured resistivity with an analytical formula containing all error terms. We conducted practical error measurements with in-vivo and bench-top experiments. We averaged errors at all frequencies for all measurements. The standard deviations of error of the quantization error of the 8-bit digital oscilloscope with voltage averaging, the nonideality of the circuit, the in-vivo motion artifact and electrical interference combined to yield an error of ±1.19%. The dimension error in measuring the syringe tube for measuring the reference saline resistivity added ±1.32% error. The estimation of the working probe constant by interpolating a set of probe constants measured in reference saline solutions added ±0.48% error. The difference in the current magnitudes used during the probe calibration and that during the tissue resistivity measurement caused ±0.14% error. Variation of the electrode spacing, alignment, and electrode surface property due to the insertion of electrodes into the tissue caused ±0.61% error. We combined the above errors to yield an overall standard deviation error of the measured tissue resistivity of ±1.96%.

Index Terms—Calibration, electrode, error analysis, error measurement, four terminals, noise measurement, probe constant, tissue resistivity.

I. INTRODUCTION

Researchers have used various methods to measure tissue resistivity. Among them, the four-terminal technique has become the most popular because of its ability to minimize the error caused by the electrode-tissue interface impedance in the measured resistivity. Normally, with the four-terminal method, a resistivity probe comprises two current electrodes for delivering and collecting the current flowing through the tissue and two voltage electrodes between the two current electrodes for sensing the voltage drop due to the current flow through the tissue. The tissue resistivity is calculated as

\[ \rho = R_a \times K_p \]  

(1)

where \( R_a \), the apparent resistance, is the ratio of the voltage difference sensed by the two voltage electrodes and the current delivered and collected by the two current electrodes and \( K_p \), the probe constant, is the ratio of the tissue resistivity divided by the apparent resistance. The probe constant is determined by the electrode configuration of the resistivity probe and is measured by calibration with a reference material.

Due to the differences in the purpose of measurement, accuracy requirement, animal species, tissue types, and the availability of instruments for resistivity measurement systems, there have been variations of the resistivity probe design, the instrumentation, and how the measurement is conducted.

Van Oosterom et al. [1] measured the intramural resistivity of canine hearts in-vivo from 10 Hz to 5 kHz using a four-terminal configuration with four consecutive electrodes of the 20 0.05 mm² electrode contacts along a shaft. They detected the voltage difference between the voltage electrodes with a differential amplifier and then utilized the phase of the current source to control the operation of two sample and hold circuits to detect the phase difference between the voltage and the current with another differential amplifier. Zheng et al. [2] measured the excised skeletal muscle resistivity of five mammals and chickens from 1 Hz to 1 MHz with a four-electrode cylindrical Plexiglas tube. They detected voltage difference between the two inner electrodes with a differential amplifier and the current through the muscle with a current-to-voltage converter and measured the output of the circuits with analog oscilloscopes. Ackmann [3] designed a four-terminal system for measuring complex bioelectric impedance from 5 Hz to 100 kHz with a Princeton Applied Research (PAR) Model 5404 lock-in amplifier and from 100 kHz to 1 MHz with a PAR Model 5402 lock-in amplifier. In his configuration, the amplitude and phase of the voltage difference between the two voltage electrodes were measured with a phase-sensitive detection technique using a signal in phase with the current as the reference. Steendijk et al. [4] measured canine myocardial resistivity in-vivo with a resistivity probe embedded in a suction cup, which attached the resistivity probe onto the epicardial surface. The resistivity probe had two perpendicular arrays of four point surface electrodes. They measured the myocardial resistivity from 5 kHz to 60 kHz with a modified analog-signal conditioner-processor (Leycom Sigma-5, CardioDynamics, Rijnsburg, The Netherlands), which was originally developed for intraventricular volume-conduc-
tance measurement. The voltage difference was sensed by a differential amplifier and then detected with phase-sensitive detection for the impedance measurement. Rigaud et al. [5] measured resistivity of sheep’s muscle, liver, lung, spleen, and intestine in a four-electrode cell in-vitro from 100 Hz to 10 MHz with a Solartron 1260 impedance/gain-phase analyzer. They used the two differential amplifiers in the Solartron 1260 to measure the voltage difference and the current magnitude, respectively. Bragós et al. [6] measured swine myocardial resistivity in-vivo from 1 kHz to 1 MHz with a four-terminal plunge probe with the left anterior descending coronary artery occluded. They used an HP 4192A impedance analyzer to supply the current and to measure the current magnitude and the output voltage of a differential amplifier, which detected the voltage difference between the two voltage electrodes. Blad and Baldetorp [7] measured complex impedance of excised normal liver tissue and tumor tissue ex-vivo from 1.5 kHz to 700 kHz in a cylindrical chamber with four stainless-steel needles penetrating the measured tissue as the current and voltage electrodes. They used two current generators controlled by a sine wave generator. The voltage difference between the voltage electrodes was detected by a differential amplifier with two input buffers. They used two synchronous demodulators, two low-pass filters, and two digital voltmeters to measure the real and imaginary parts of the tissue impedance.

Several researchers have analyzed the error of the four-terminal measurement. Swanson and Webster [8] analyzed the errors in four-electrode impedance plethysmography due to inadequate instrumentation, improper electrode application, and physiological changes. Ackmann [3] evaluated the phase error of four-terminal measurement by experimentally manipulating the cable length and extra resistors to emulate the imbalance between the two voltage electrodes in the cable stray capacitance and the electrode-tissue interface impedance. The result was consistent with his computer simulation of an equivalent circuit model. He further measured the phase error of the lock-in amplifier, the current source, and the preamplifiers to obtain a phase calibration for the overall system. Riu et al. [9] used an equivalent circuit of a resistivity measurement system to analyze the requirements for limiting the magnitude of the coupling capacitance, common-mode voltage, and the voltage-measuring circuit’s input impedance on the system design to attain a measurement error below 1%. Boone and Holder [10] studied the problem of attaining a high-precision electrical impedance tomodiagram (EIT) due to errors caused by common-mode voltage, electrode-tissue interface impedance, input impedance of the differential amplifiers, and output impedance of current sources. Al-Hatib [11] derived analytical formulas using an equivalent circuit of the patient-instrument connection for a four-terminal measurement configuration to describe the errors in the measured bioimpedance and in the overall common-mode rejection ratio (CMRR) induced by the cable capacitance in combination with the skin-electrode impedance and the bioimpedance to be measured. He further constructed an experimental circuit to demonstrate the error caused by the stray capacitance of various cable types and different values of bioimpedance and to compare the performances of a single-ended version and a symmetrical version of the current source. Scharfetter et al. [12] modeled with the PSpice program a whole body or segmental bioimpedance spectroscopy system including the stray capacitance between cables, electrode leads, different body segments and earth, and signal ground and earth. They conducted simulation from 5 kHz to 1 MHz and found the latter two types of stray capacitance cause a significant spurious dispersion in the measured impedance spectra at frequencies higher than 500 kHz.

We have not found literature that leads to the evaluation or estimation of the overall error in the measured resistivity. In this study, we utilize a typical four-terminal resistivity measurement system as an example to demonstrate a systematic method for analyzing the measurement error of tissue resistivity.

II. THEORY

We have described our system and procedure for measuring tissue resistivity [13]. In this error analysis paper, we use the same equations as in that paper except that we will add error terms wherever appropriate.

Our measurement system is shown in [13, Fig. 1]. During a frequency sweep, the computer calculates the tissue resistivity at each frequency by

\[
(\rho_r + \varepsilon_r) = \left( \frac{V_2}{V_1} \right) (K_0 + \varepsilon_0)(K_c + \varepsilon_c)(K_w + \varepsilon_w)(K_p + \varepsilon_p),
\]

(2)

In (2), \((\rho_r + \varepsilon_r)\) is the measured tissue resistivity with \(\rho_r\) the real tissue resistivity and \(\varepsilon_r\) the measurement error. \(V_1\) and \(V_2\) are the output voltage values of channel 1 and channel 2, respectively, of the digital oscilloscope. \((K_0 + \varepsilon_0)\) is the oscilloscope constant defined as \((V_0/V_2)/(V_1/V_2)\), the inverse response of the oscilloscope when measuring a voltage ratio, where \(V_0\) is the output voltage of the differential amplifier and \(V_2\) the current-to-voltage converter, respectively.

The ideal oscilloscope constant \(K_0\) equals one and the oscilloscope error \(\varepsilon_0\) accounts for the difference between the two channels of the digital oscilloscope mostly due to the quantization error.

\[(K_c + \varepsilon_c)\] is the circuit constant defined as \((V_c/I_c)/(V_e/I_e)\), the inverse response of the tissue resistance-detecting circuit, where \(V_c\) is the input voltage of the differential amplifier and \(I_c\) the input current of the current-to-voltage converter. The ideal circuit constant \(K_c\) equals the ratio of the gain of the current-to-voltage converter divided by that of the differential amplifier. The circuit error \(\varepsilon_c\) is due to circuitry nonidealities, such as insufficient CMRR of the differential amplifier, finite gain of operational amplifiers in the circuits, and leakage current.

\[(K_w + \varepsilon_w)\] is the wire constant defined as \((V_e/I_e)/(V_c/I_c)\), where \(V_e/I_e\) is the real apparent resistance of measuring the tissue with the resistivity probe and \(V_c/I_c\) is the apparent resistance sensed by the tissue resistance-detecting circuit. The ideal wire constant \(K_w\) is 1 and the wire error \(\varepsilon_w\) accounts for the loading effect of the stray capacitance on the signal wires between the electrodes and the circuit in series with the electrode-tissue interface impedance.

\[(K_p + \varepsilon_p)\] is the measured probe constant. It contains an error term \(\varepsilon_p\) due to the inaccuracy in measuring the probe constant. The real probe constant \(K_p\) equals the ratio of the real tissue resistivity \(\rho_t\) divided by the real apparent resistance \(V_e/I_e\), i.e., the inverse response of the resistivity probe, where \(V_e\) is the
voltage difference between the voltage electrodes and \( I_c \) is the current flowing through the current electrodes.

Equation (2) means that we obtain the tissue resistivity by multiplying the output ratio of the oscilloscope with the inverse responses of the oscilloscope, the circuits, the wires, and the resistivity probe. \((V_{1}/V_{2})/(K_{s}+\varepsilon_{s})(K_{w}+\varepsilon_{w})(K_{t}+\varepsilon_{t})\) in (2) is equivalent to the apparent resistance in (1). Note that, in practical measurement, it is usually not necessary to measure each of the four constants separately. In our measurement, we combine the oscilloscope constant and the circuit constant into an oscilloscope–circuit constant \((K_{o-c}+\varepsilon_{o-c})\) and the wire constant and the probe constant into a wire–probe constant \((K_{w-p}+\varepsilon_{w-p})\). We re-express the equation for the measured tissue resistivity as

\[
(p_{r}+\varepsilon_{r}) = \left(\frac{V_{1}}{V_{2}}\right)(K_{o-c}+\varepsilon_{o-c})(K_{w-p}+\varepsilon_{w-p}) \tag{3}
\]

Our procedure of tissue resistivity measurement includes several steps. The following describes and formulates the error sources in each step. Note that, in the following, the subscript next to a parenthesis represents the step number in which the quantity inside the parenthesis is measured or is pertinent to the configuration used.

In Step 1, the measurement of the oscilloscope–circuit constant, the practical value of the oscilloscope–circuit constant calculated by the computer can be expressed as

\[
(K_{o-c}+\varepsilon_{o-c})_{1} = \frac{(R_{f}+\varepsilon_{f})_{1}}{\left(\frac{V_{1}}{V_{2}}\right)_{1}} \tag{4}
\]

where \((R_{f}+\varepsilon_{f})_{1}\) is the resistance of the reference resistor measured with a multimeter. The error term \(\varepsilon_{f}\) is due to the accuracy limit of the multimeter. The exact value of \(R_{f}\) is unknown and the measured value \((R_{f}+\varepsilon_{f})_{1}\) is used in the calculation of the oscilloscope–circuit constant. The measured value \((V_{1}/V_{2})_{1}\) contains errors induced by the oscilloscope and the tissue resistance-detecting circuit. The exact value of the oscilloscope–circuit constant \(K_{o-c}\) is also unknown and the value \((K_{o-c}+\varepsilon_{o-c})_{1}\) measured in this step will be used in the calculation in the subsequent steps.

In Step 2, the measurement of saline resistivity, the measured value of the saline resistivity is

\[
(p_{s}+\varepsilon_{s})_{2} = \left(\frac{V_{1}}{V_{2}}\right)_{2}(K_{o-c}+\varepsilon_{o-c})_{1}(A_{d}/A)_{2} \tag{5}
\]

where \((A_{d}/A)(K_{A/d}+\varepsilon_{A/d})_{2}\) is the value of the wire–probe constant of the syringe tube, where \(A\) is the inner cross-sectional area of the syringe tube and \(d\) is the voltage electrode distance. The error term \(\varepsilon_{A/d}\) is due to the inaccuracy of measuring \(A\) and \(d\) with a ruler or a caliper. The wire error at this step is less than 0.1% with 15 cm short wires between the electrodes and the circuit, 3 cm distance between the voltage electrodes, and 1 cm distance between a current electrode and its adjacent voltage electrode.

We use the oscilloscope–circuit constant \((K_{o-c}+\varepsilon_{o-c})_{1}\) measured in Step 1 in this equation since we used the same circuit and the same oscilloscope in all steps.

In Step 3, the measurement of the wire–probe constant, the computer calculated the wire–probe constant \((K_{w-p}+\varepsilon_{w-p})_{3}\) as \((p_{s}+\varepsilon_{s})_{2} ([V_{1}/V_{2}](K_{o-c}+\varepsilon_{o-c})_{1}])^{-1}\) according to (2). However, we noticed that the temperature in a laboratory could be different from location to location and from time to time, and the heat emitted from the instruments could heat up the ambient temperature of the measurement site by more than 1°C. Hence, the real value of the measured wire–probe constant is

\[
(K_{w-p}+\varepsilon_{w-p})_{3} = \frac{(p_{s}+\varepsilon_{s})_{2}(K_{T}+\varepsilon_{T})_{3}}{\left(\frac{V_{1}}{V_{2}}\right)_{3}(K_{o-c}+\varepsilon_{o-c})_{1}} \tag{6}
\]

where \((K_{T}+\varepsilon_{T})_{3}\) represents the change of saline resistivity due to the change of the temperature in the saline solution from Step 2 to Step 3. and \(K_{T}\) equals one. Suppose that \(T_{2}\) is the saline temperature at \(T_{2}\) and \(T_{3}\) at \(T_{3}\) respectively. and \(k_{s}\) is the temperature coefficient of the saline resistivity, then \((K_{T}+\varepsilon_{T})_{3}\) equals \((1+k_{s})^{(T_{3}-T_{2})}\). In a preliminary experiment, we measured the resistivity of saline solutions of various salt concentrations ranging from 0.1% to 1.1% at temperatures from 20°C to 40°C. We found that the saline resistivity decreased by 2% as the temperature increased by 1°C. Hence, \(k_{s} = -2\%\).

We minimized the temperature error by measuring the saline temperatures \(T_{2}\) and \(T_{3}\) at Step 2 and Step 3, respectively, with a thermometer and by dividing the measured wire–probe constant by the temperature constant \((K_{T}+\varepsilon_{T})_{3}\).

In an in-vivo animal experiment for myocardial resistivity measurement, such as the one described in [13], there were movement artifact [15] and electrical interference coming from the measured animal, the equipment, such as the ventilator, and the power lines running through the walls around the laboratory room. We use a noise constant \((K_{n}+\varepsilon_{n})_{3}\), to represent the effect of the noise. The ideal noise constant \(K_{n}\) is 1 and the noise error \(\varepsilon_{n}\) represents the error caused by the motion artifact and the electrical interference in the in-vivo measurement. The noise constant is implicit in the expression of \((V_{1}/V_{2})_{3}\) as \((K_{n}+\varepsilon_{n})_{3} (V_{e}/I_{e})_{3} / (K_{o-c}+\varepsilon_{o-c})_{3}\).

In Step 4, the measurement of the tissue resistivity, according to (2), the computer calculated the tissue resistivity as the voltage ratio output of the digital oscilloscope multiplied by the oscilloscope–circuit constant obtained in Step 1 and the wire–probe constant obtained in Step 3. However, because the wire–probe constant could possibly change in this step, the resistivity obtained with the computer calculation is

\[
(p_{r}+\varepsilon_{r})_{4} = \left(\frac{V_{1}}{V_{2}}\right)_{4}(K_{o-c}+\varepsilon_{o-c})_{1}(K_{w-p}+\varepsilon_{w-p})_{3} \times (K_{g}+\varepsilon_{g})_{4} (K_{i}+\varepsilon_{i})_{4} (K_{e}+\varepsilon_{e})_{4} \tag{7}
\]

where \((K_{g}+\varepsilon_{g})_{4}\) is called the interpolation constant, \((K_{i}+\varepsilon_{i})_{4}\) the current constant, and \((K_{e}+\varepsilon_{e})_{4}\) the electrode constant, with all their ideal values \(K_{g}\), \(K_{i}\), and \(K_{e}\) being 1. The interpolation error \(\varepsilon_{g}\) represents the error of estimating the wire–probe constant for a measurement from a set of wire–probe constants measured by calibrating the plunge probe in each of a set of saline solutions with various resistivity values. The current error \(\varepsilon_{i}\) represents the deviation of the wire–probe constant due to the difference of the current magnitude during tissue resistivity measurement.
measurement and that during probe calibration. It arose due to the fact that the interface impedance between the electrode and the tissue or the saline solution changes the distribution of the current density flowing through the interface [14], [15]. The electrode error accounts for the change of the wire–probe constant caused by the action of inserting the electrodes into the tissue, which may change the electrical property of the electrode surface or alter the electrode spacing or alignment.

Just as in Step 3, there was also in-vivo motion artifact and electrical interference in this step. The noise constant (\(K_{nr} + \varepsilon_{nr}\)) is implicit in the measured voltage ratio \((\frac{V_1}{V_2})_d\), i.e., \((\frac{V_1}{V_2})_d = (K_{nr} + \varepsilon_{nr})_d(V_c/I_c)_d/(K_{oc-c} + \varepsilon_{oc-c})_d\).

By substituting the detailed expression for \((K_{nr} + \varepsilon_{nr})_d\) in (7) and substituting \((V_c/I_c)_d/(K_{oc-c} + \varepsilon_{oc-c})_d\) for \((V_1/V_2)_d\) and \((K_{nr} + \varepsilon_{nr})_j(V_c/I_c)_j/(K_{oc-c} + \varepsilon_{oc-c})_j\) for \((V_1/V_2)_j\), \(j = 3\) to 4, based on the assumption that \((V_c/I_c) = (V_c/I_c)_d\) in all steps because of using short signal wires, we obtain

\[
(\rho_r + \varepsilon_r)_d = \left(\frac{V_1}{I_c}\right)_d^2 \frac{V_1^2}{I_c^2} (K_{oc-c} + \varepsilon_{oc-c})_d \times \left(\frac{A}{d}\right) (K_{A/d} + \varepsilon_{A/d})_2 (K_T + \varepsilon_T)_3 \times (K_g + \varepsilon_g)_j (K_i + \varepsilon_i)_j (K_e + \varepsilon_e)_4
\]

where \((K_{oc-c} + \varepsilon_{oc-c})_d\) is called the overall oscilloscope–circuit–noise constant and is defined as \([(K_{oc-c} + \varepsilon_{oc-c})_j](K_{oc-c} + \varepsilon_{oc-c})_d(K_{nr} + \varepsilon_{nr})_d)/(K_{oc-c} + \varepsilon_{oc-c})_d(K_{nr} + \varepsilon_{nr})_d). The reason for combining the oscilloscope–circuit and the noise constants is because of the difficulty of measuring them separately [see (12)]. Note that \((K_{oc-c} + \varepsilon_{oc-c})_j\) is a known value that has been measured in Step 1 while \((K_{oc-c} + \varepsilon_{oc-c})_j\) and \((K_{nr} + \varepsilon_{nr})_j\), \(j = 3\) to 4, are implicit values contained in the measured \((V_1/V_2)_j\) after the subtraction of the oscilloscope–circuit terms are different because they are obtained from distinct measurements.

According to (8), we express the statistics of the overall error of the measured tissue resistivity \((\rho_r + \varepsilon_r)_d\) as [16]

\[
\frac{\varepsilon_{m_r}}{\rho_r} = \left(\frac{(K_{oc-c} + \varepsilon_{oc-c})_n}{K_{oc-c}_n}\right) \left(\frac{(K_{A/d} + \varepsilon_{A/d})_2}{K_{A/d}_2}\right) \times \left(\frac{(K_T + \varepsilon_T)_3}{K_T}_3\right) \times \left(\frac{(K_i + \varepsilon_i)_4}{K_i}_4\right) \frac{1}{\left(\frac{V_c}{I_c}\right)_d^2} \left(\frac{V_1}{I_c}\right)_d^2 (K_{oc-c} + \varepsilon_{oc-c})_d \times \left(\frac{A}{d}\right) (K_{A/d} + \varepsilon_{A/d})_2 (K_T + \varepsilon_T)_3 \times (K_g + \varepsilon_g)_j (K_i + \varepsilon_i)_j (K_e + \varepsilon_e)_4
\]

\[
\frac{\varepsilon_{s_r}}{\rho_r} = \left(\frac{(K_{oc-c} + \varepsilon_{oc-c})_n}{K_{oc-c}_n}\right) \left(\frac{(K_{A/d} + \varepsilon_{A/d})_2}{K_{A/d}_2}\right) \times \left(\frac{(K_T + \varepsilon_T)_3}{K_T}_3\right) \times \left(\frac{(K_i + \varepsilon_i)_4}{K_i}_4\right) \frac{1}{\left(\frac{V_c}{I_c}\right)_d^2} \left(\frac{V_1}{I_c}\right)_d^2 (K_{oc-c} + \varepsilon_{oc-c})_d \times \left(\frac{A}{d}\right) (K_{A/d} + \varepsilon_{A/d})_2 (K_T + \varepsilon_T)_3 \times (K_g + \varepsilon_g)_j (K_i + \varepsilon_i)_j (K_e + \varepsilon_e)_4
\]

In summary, by conducting Step 5 and Step 6, we obtained a quantity that characterized the combined statistic of the interference noise \(\varepsilon_{nr}\) and the oscilloscope–circuit error \(\varepsilon_{oc-c}\) at Step 5. Fig. 1(b) shows 152 samples of \((\frac{V_1}{V_2})_5\) along with their mean and standard deviation.

In order to evaluate the error of the measured tissue resistivity with (9) and (10), we performed a series of experiments to measure the individual error terms separately at 1 Hz, 10 Hz, 100 Hz, 1 kHz, 10 kHz, 100 kHz, 500 kHz, and 1 MHz.

To evaluate the statistics of the overall oscilloscope–circuit–noise constant \((K_{oc-c} + \varepsilon_{oc-c})_d\), we first measured the individual oscilloscope–circuit constants \((K_{oc-c} + \varepsilon_{oc-c})_d\) with reference resistors of nominal resistance 50, 100, 150, 200, and 250 \(\Omega\), which encompassed possible apparent resistance when measuring tissue resistivity. We measured the reference resistance with an HP 34401A multimeter. We added two other resistors to the two ends of each reference resistor to form a series combination of 1:3:1 resistance ratio. For each reference resistor we took eight measurements of the oscilloscope–circuit constant \((K_{oc-c} + \varepsilon_{oc-c})_d\), which was calculated as \((R_f + \varepsilon_f)/(V_1/V_2)_d\). Fig. 1(a) shows the 40 measured oscilloscope–circuit constants along with their mean and standard deviation.

Next, we measured the noise constant in an in-vivo condition. When an open-chest pig was alive, we put a cup of saline on top of the beating heart and immersed the plunge probe in the saline. We assumed that the noise from the pig, the ventilator, and the wall has the same effect on the measuring signals as that when we inserted the plunge electrodes in the pig’s myocardium. The computer accessed the noise-contaminated voltage ratio \(V_1/V_2\) from the digital oscilloscope. We designated this step as Step 5.

After the pig died and the heart stopped beating, we turned off the ventilator and measured the voltage ratio \(V_1/V_2\) with the plunge probe immersed in the saline solution and with the same voltage at the output of the function generator as that in the previous step. At this time, the waveform of \(V_c\) and \(V_t\) looked clean. There was little in-vivo noise in the measured signal and, hence, the noise error \(\varepsilon_{nr}\) was virtually zero. We accessed the voltage ratio with a 401-time averaging and, hence, the oscilloscope–circuit error \(\varepsilon_{oc-c}\) was divided by 20 and could be neglected. We call this Step 6.

Assuming negligible wire error, i.e., assuming \((V_c/I_c)_j = (V_c/I_c)_d\), \(j = 5\) or 6, the ratio of the two measured voltage ratios will be

\[
\frac{(\frac{V_1}{V_2})_5}{(\frac{V_1}{V_2})_6} = \left(\frac{(\frac{V_c}{I_c})}{(\frac{V_c}{I_c})_d}\right) \left(\frac{(\frac{V_c}{I_c})_d}{(\frac{V_c}{I_c})_d}\right) (K_{oc-c} + \varepsilon_{oc-c})_d
\]

Because \((V_c/I_c)_d = (V_c/I_c)_6\) and \(K_n = 1\), (11) can be reduced to

\[
\frac{(\frac{V_1}{V_2})_5}{(\frac{V_1}{V_2})_6} = \left(\frac{(\frac{V_c}{I_c})}{(\frac{V_c}{I_c})_d}\right) \left(\frac{(\frac{V_c}{I_c})_d}{(\frac{V_c}{I_c})_d}\right) (K_{oc-c} + \varepsilon_{oc-c})_d
\]
To simulate the statistic of the overall oscilloscope–circuit–noise constant \( (K_{\alpha_{c-n}} + \varepsilon_{\alpha_{c-n}}) \), we randomly select two from the measured oscilloscope–circuit constants shown in Fig. 1(a) and substitute them for \( (K_{\sigma_{c-c}} + \varepsilon_{\sigma_{c-c}}) \) and \( (K_{\sigma_{c-c}} + \varepsilon_{\sigma_{c-c}}) \) terms in the expression of \( (K_{\alpha_{c-n}} + \varepsilon_{\alpha_{c-n}}) \), i.e., \( ((K_{\sigma_{c-c}} + \varepsilon_{\sigma_{c-c}})/\varepsilon_{\sigma_{c-c}})(K_{\sigma_{c-c}} + \varepsilon_{\sigma_{c-c}})/(K_{\sigma_{c-c}} + \varepsilon_{\sigma_{c-c}}) \). We assume that \( (K_{\sigma_{c-c}} + \varepsilon_{\sigma_{c-c}})/\varepsilon_{\sigma_{c-c}} \) and \( (K_{\sigma_{c-c}} + \varepsilon_{\sigma_{c-c}})/\varepsilon_{\sigma_{c-c}} \) have the same statistics as \( (K_{\sigma_{c-n}} + \varepsilon_{\sigma_{c-n}})/\varepsilon_{\sigma_{c-n}} \). Hence, we randomly select two samples from the results shown in Fig. 1(b) and substitute them for \( (K_{\sigma_{c-c}} + \varepsilon_{\sigma_{c-c}})/\varepsilon_{\sigma_{c-c}} \) and \( (K_{\sigma_{c-c}} + \varepsilon_{\sigma_{c-c}})/\varepsilon_{\sigma_{c-c}} \) to calculate \( (K_{\sigma_{c-c}} + \varepsilon_{\sigma_{c-c}})/(K_{\sigma_{c-c}} + \varepsilon_{\sigma_{c-c}}) \). Fig. 1(c) shows 1000 samples of the overall oscilloscope–circuit–noise constant generated this way. Fig. 2(a) shows the mean and standard deviation of \( \varepsilon_{\alpha_{c-n}}/K_{\alpha_{c-n}} \). Table I also shows the standard deviation.

The variation in the measured oscilloscope–circuit–constant \( (K_{\alpha_{c-n}} + \varepsilon_{\alpha_{c-n}}) \) is primarily due to the quantization error of the digital oscilloscope. The HP54600B Digital Oscilloscope provides 8-bit resolution with ±1.9% accuracy as described in its User and Service Guide. To measure the variation of the oscilloscope constant, we connected the probes of the two channels of the oscilloscope together to the output of the function generator and measured the voltage ratio \( (V_1/V_2) \), which equals the value of \( (K_{\alpha_{c-n}} + \varepsilon_{\alpha_{c-n}}) \) in this configuration. We used various output amplitudes of the function generator and various vertical scales of the digital oscilloscope. We found that the variation of the measured oscilloscope constant is commensurate with that of the oscilloscope–circuit constant.

For the error \( \varepsilon_{A/d} \) of measuring the dimension of the syringe tube used to measure the saline resistivity in Step 2, we used a caliper with 0.01-mm resolution to measure the inner diameter.
Fig. 2. The individual and overall errors of the measured myocardial resistivity. In (a) and (f), the asterisks represent the errors in the scenario of our in-vivo measurement, where we averaged 100 repeated measurements in Step 1 and averaged ten repeated measurements in Step 2 and in Step 3. The triangular symbols represent the errors in measurements without averaging. (a) $\varepsilon_{m_{\text{area}}}/K_{\text{area}}$ and $(\varepsilon_{m_{\text{area}}} \pm \varepsilon_{s_{\text{area}}})/K_{\text{area}}$, (b) $\varepsilon_{m_{\text{A/d}}}/K_{\text{A/d}}$ and $(\varepsilon_{m_{\text{A/d}}} \pm \varepsilon_{s_{\text{A/d}}})/K_{\text{A/d}}$, (c) $\varepsilon_{m_{\text{g}}}/K_{\text{g}}$ and $(\varepsilon_{m_{\text{g}}} \pm \varepsilon_{s_{\text{g}}})/K_{\text{g}}$, (d) $\varepsilon_{m_{\text{e}}}/K_{\text{e}}$ and $(\varepsilon_{m_{\text{e}}} \pm \varepsilon_{s_{\text{e}}})/K_{\text{e}}$, and (f) $\varepsilon_{m_{\text{r}}}/K_{\text{r}}$.

TABLE I
ERROR STATISTICS OF THE MEASURED TISSUE RESISTIVITY. (STANDARD DEVIATION, IN PERCENTILE)

<table>
<thead>
<tr>
<th>Frequency (Hz)</th>
<th>1</th>
<th>10</th>
<th>100</th>
<th>1 k</th>
<th>100 k</th>
<th>500 k</th>
<th>M</th>
<th>Average</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\varepsilon_{\text{A/d}}/K_{\text{A/d}}$ (Note 1)</td>
<td>$\pm 2.37$</td>
<td>$\pm 2.73$</td>
<td>$\pm 2.34$</td>
<td>$\pm 2.60$</td>
<td>$\pm 1.95$</td>
<td>$\pm 1.66$</td>
<td>$\pm 1.54$</td>
<td>$\pm 2.01$</td>
</tr>
<tr>
<td>$\varepsilon_{\text{A/d}}/K_{\text{A/d}}$ (Note 2)</td>
<td>$\pm 1.31$</td>
<td>$\pm 1.52$</td>
<td>$\pm 1.30$</td>
<td>$\pm 1.44$</td>
<td>$\pm 1.08$</td>
<td>$\pm 0.92$</td>
<td>$\pm 0.86$</td>
<td>$\pm 1.12$</td>
</tr>
<tr>
<td>$\varepsilon_{\text{A/d}}/K_{\text{A/d}}$</td>
<td>$\pm 1.32$</td>
<td>$\pm 1.32$</td>
<td>$\pm 1.32$</td>
<td>$\pm 1.32$</td>
<td>$\pm 1.32$</td>
<td>$\pm 1.32$</td>
<td>$\pm 1.32$</td>
<td>$\pm 1.32$</td>
</tr>
<tr>
<td>$\varepsilon_{\text{r}}/K_{\text{r}}$ (Note 3)</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>$\varepsilon_{\text{g}}/K_{\text{g}}$</td>
<td>$\pm 0.42$</td>
<td>$\pm 0.63$</td>
<td>$\pm 0.41$</td>
<td>$\pm 0.46$</td>
<td>$\pm 0.45$</td>
<td>$\pm 0.44$</td>
<td>$\pm 0.44$</td>
<td>$\pm 0.63$</td>
</tr>
<tr>
<td>$\varepsilon_{\text{e}}/K_{\text{e}}$</td>
<td>$\pm 0.18$</td>
<td>$\pm 0.23$</td>
<td>$\pm 0.15$</td>
<td>$\pm 0.15$</td>
<td>$\pm 0.11$</td>
<td>$\pm 0.09$</td>
<td>$\pm 0.09$</td>
<td>$\pm 0.14$</td>
</tr>
<tr>
<td>$\varepsilon_{\text{r}}/K_{\text{r}}$</td>
<td>$\pm 0.59$</td>
<td>$\pm 0.65$</td>
<td>$\pm 0.79$</td>
<td>$\pm 0.62$</td>
<td>$\pm 0.49$</td>
<td>$\pm 0.28$</td>
<td>$\pm 0.51$</td>
<td>$\pm 0.49$</td>
</tr>
<tr>
<td>$\varepsilon_{\text{r}}/K_{\text{r}}$ (Note 1)</td>
<td>$\pm 2.81$</td>
<td>$\pm 3.17$</td>
<td>$\pm 2.86$</td>
<td>$\pm 3.02$</td>
<td>$\pm 2.47$</td>
<td>$\pm 2.22$</td>
<td>$\pm 2.19$</td>
<td>$\pm 2.53$</td>
</tr>
<tr>
<td>$\varepsilon_{\text{r}}/K_{\text{r}}$ (Note 2)</td>
<td>$\pm 2.01$</td>
<td>$\pm 2.20$</td>
<td>$\pm 2.10$</td>
<td>$\pm 2.11$</td>
<td>$\pm 1.87$</td>
<td>$\pm 1.74$</td>
<td>$\pm 1.77$</td>
<td>$\pm 1.89$</td>
</tr>
</tbody>
</table>

Note 1: Without any averaging in any step.
Note 2: With 101-time, 10-time, 10-time, and no averaging in Step 1, 2, 3, and 4, respectively.
Note 3: The temperature error became negligible after being corrected with a $-2\%/{ }^\circ\text{C}$ temperature coefficient.

of the syringe and the voltage electrodes distance 15 times. The calculated value for the standard deviation of $A/d$ is 1.32%. We assumed the mean value of the error of measuring $A/d$ to be zero. Fig. 2(b) shows the mean and standard deviation of $\varepsilon_{A/d}/K_{A/d}$. Table I also shows the standard deviation.

The temperature error $\varepsilon_T$ became less than 0.2% after being minimized in Step 3 by correcting the measured wire–probe constant with the temperature coefficient of the saline solution.

To evaluate the interpolation error $\varepsilon_{\text{interp}}$, we made a first set of eight saline solutions of approximately 100, 150, 200, 250, 300, 350, 400, and 500 $\Omega \cdot \text{cm}$, which covered the practical range of in-vivo tissue resistivity, and we measured the set of wire–probe constants in the saline solutions, just as we normally did in Step 2 and Step 3 for calibrating the resistivity probe. Then we prepared a second set of seventeen saline solutions with resistivity approximately uniformly distributed between 100 and 500 $\Omega \cdot \text{cm}$. On the one hand, we estimated the plunge probe’s wire–probe constants in each of the second set of saline solutions with the same method we used in Step 4 to estimate the wire–probe constant of the plunge probe in the tissue by interpolating the measured wire–probe constants in the first set of saline solutions. On the other hand, we repeated Step 2 and Step 3 to practically measure the wire–probe constants in the second set of saline solutions. Fig. 3 shows the distribution of the measured and the estimated wire–probe constants in the second set of saline solutions, as well as the wire–probe constants mea-
fig. 3. the wire–probe constants $K_{wp}$ of the two sets of saline solution used to evaluate the interpolation error $\varepsilon_\varepsilon$. only the results at 1 MHz are shown here. in a practical in-vivo myocardial resistivity measurement, only the wire–probe constants of the first saline set are measured, and those of the myocardium (represented by the second saline set) are estimated from those of the first saline set. for the evaluation of the interpolation error, the wire–probe constants of the second saline set were also measured and the difference between the estimated and the measured wire–probe constants of the second saline set constituted the interpolation error.

measured in the first set of saline solutions, at 1 MHz. the interpolation error $\varepsilon_\varepsilon$ accounts for the difference between the estimated and the measured wire–probe constant in each of the second set of saline solutions. we further summate the interpolation error at each frequency with different weighting for different saline resistivity according to the distribution of the swine myocardial resistivity measured in-vivo [13]. the result is shown in Table I and Fig. 2(c).

To evaluate the current error $\varepsilon_\varepsilon$, we first reviewed the current magnitude we used in Step 3 and Step 4 for our in-vivo measurement of swine myocardial resistivity [13] and found that the current magnitude at each frequency varied less than ±20% from Step 3 to Step 4. we repeated Step 3 to measure the wire–probe constants of the plunge probe in a set of saline solutions. for each saline solution, at each frequency, we measured the wire–probe constant with various current magnitudes from 80%, 84%, ..., 116%, to 120% with respect to the magnitude we normally used to calibrate the plunge probe in each of the saline solutions. we calculated the current error $\varepsilon_\varepsilon$, as the change of the measured wire–probe constants calibrated with non-100% current magnitude with respect to that calibrated with 100% current magnitude. we further summated the current error at each frequency with different weighting for different saline resistivity according to the distribution of the swine myocardial resistivity measured in-vivo [13]. Fig. 4 shows the wire–probe constants measured with various current magnitudes along with their mean value and standard deviation calculated with the above-mentioned weighting. Table I and Fig. 2(d) show the statistics of the current error.

We evaluated the electrode error $\varepsilon_\varepsilon$ by observing the change of apparent resistance of a saline solution measured with the plunge probe before the first insertion into the myocardium of an open-chest pig and after each subsequent insertion. we repeated this experiment on three pigs when we were doing in-vivo resistivity measurement in two directions at three fixed sites of
from insertion of the plunge probe into the myocardium of open-chest pigs’ beating hearts. The apparent resistance of a saline solution was measured before the first insertion and after each insertion. The cumulative electrode error is the deviation of the apparent resistance measured after each insertion with respect to that measured before the first insertion. The electrode error shown in Table I and Fig. 5(e) is the average of the cumulative electrode error after every insertion from the three pig experiments.

each pig’s left ventricle [13]. We actually had seven insertions in each of the three pigs with the extra one for re-insertion when we found the electrodes were not inserted well or the orientation of the electrode array was not in the right direction. The process included inserting the electrodes into the swine myocardium, rinsing them in a first saline solution, measuring the apparent resistance in a second saline solution with the same concentration as the first saline solution. We observed the difference between the apparent resistance measured in the second saline solution after the electrodes were pulled out and the apparent resistance measured before the first insertion. Fig. 5 shows the cumulative electrode error at each frequency after each insertion. We calculated the electrode error $\varepsilon_e$ as the average of the drift of the measured apparent resistance after every insertion with respect to the preinsertion value of the three in-vivo pig experiments.

Note that, during the measurements of $\varepsilon_g$ and $\varepsilon_e$, the saline temperature could change considerably. To ensure the accuracy of the error evaluation, first we kept the saline temperature as constant as possible. Second, we measured the saline temperature with a thermistor and corrected the error caused by temperature variation with the $-2\,^\circ\text{C}$ temperature coefficient.

After obtaining the statistics of all the individual errors, we estimated the overall error with (9) and (10). Table I and Fig. 2(f) show the statistics of the overall error of the measured tissue resistivity. The mean value is systematic error, we know it, and we correct for it when we report the final results [13]. Therefore, it is not shown in Table I, which shows only the standard deviation of the random error.

IV. DISCUSSION

Our system is a low-cost instrument for tissue resistivity measurement. The components used for building the differential amplifier, the current-to-voltage converter, and the temperature-measuring circuit cost less than US $20. The year 2000 list prices of the HP54600B digital oscilloscope, HP54659B Measurement/storage module, and the HP33120A function generator are US $2,550, US $495, and US $1,795, respectively. Instruments of compatible functions are readily available in most laboratories as well as the personal computer and the LabVIEW tools. In contrast, the HP4192A impedance analyzer used by Bragós [6] costs US $25,500, the PAR 5204 lock-in amplifier and PAR 5202 lock-in amplifier used by Ackmann [3] costs US $10,000, the Solartron 1260 impedance analyzer used by Rigaud [5] costs US $25,500.

It is possible to reduce the overall error in the measured resistivity by reducing individual errors. $N$-time averaging of a repeated uncorrelated measurements divides the standard deviation of measurement result by a factor of $(N - 1)^{1/2}$ [16]. The digital oscilloscope provides an averaging function to reduce the quantization error. However, consistent triggering for averaging is not easy at low frequencies. We programmed our LabVIEW virtual instrument to fulfill the averaging function so that we can average even at low frequencies such as 1 Hz. In our in-vivo measurement, where we averaged 101 repeated measurements in Steps 1 and ten repeated measurements in Steps 2 and 3, the standard deviation of $\varepsilon_{\text{error}}$ is theoretically divided by a factor of 1.8. The price paid is large measurement time. In our system, a resistivity measurement at eight frequencies from 1 Hz to 1 MHz takes 90 s without averaging. A resistivity measurement with ten-time averaging and 101-time averaging take 150 s and 12 min, respectively. In some situations, such as a postmortem measurement, it is not appropriate to take averaging because of the fast changes in tissue resistivity.

We can reduce the oscilloscope error by using a digital oscilloscope with higher resolution such as 14-bit resolution.

In Fig. 3, the dependence of the wire–probe constant on the apparent resistance was due to the stray capacitance between the measured animal and the ground [13]. In [13], we demonstrated how the in-vivo calibration alleviated the error caused by this stray capacitance. However, there was still residual error, the interpolation error $\varepsilon_g$, because we had to obtain the wire–probe constant for calculating the tissue resistivity by interpolating the finite set of the wire–probe constants calibrated with a set of saline solutions. It is possible to reduce the interpolation error $\varepsilon_g$ by improving the interpolation algorithm. It is also possible to further reduce the error by calibration with more saline solutions within the target resistivity range so that the interpolation is more accurate.
In [13], we controlled the output of the function generator at constant voltage. The current magnitude during in-vivo tissue measurement varied less than ±20% from that during probe calibration. The current error εc can be minimized by using same current magnitude for calibrating the tissue resistivity probe in Step 3 and for measuring the tissue resistivity in Step 4. Nevertheless, the current error δc shown in Fig. 4 has been quite small compared with other errors. Further improvement might not make much sense. The constant voltage method used in [13] is good enough, although our LabVIEW virtual instrument also provides current control modes.

Our experiment showed that the change of electrode condition after successive insertions caused a larger error than the change of a single insertion. We, thus, suggest frequent calibration during multiple insertions to reduce the electrode error εc.

In all four steps in our measurement procedure, we used signal wires shorter than 15 cm, which were short enough so that the error caused by the stray capacitance combined with the electrode-tissue interface impedance was less than 0.1%. In case long wires are used, driven shields [17] can minimize the error from stray capacitance. We also can minimize the effect of stray capacitance by placing the buffer stage of the tissue resistance-detecting circuit close to the electrodes.

In our analysis and experiments, we have tried to include all error sources in a typical plunge probe resistivity measurement. However, depending on the situation of the measured tissue, the electrodes configuration, and the circuits and the instruments used, there are possibly other errors. For example, when measuring myocardial resistivity with the plunge probe, there will be error due to anisotropic resistivity distribution with respect to the fiber direction; the myocardial fiber direction changes transmurally about 160° from epicardial layer to endocardial layer [18]. In an in-vivo measurement on an open-chest pig’s ventricle, we tried to minimize the influence of the fiber direction by making a myocardial flap 1 mm wide, 1 mm thick, and 15 mm long along the local fiber direction with one end still connected to the heart. We laid the flap in a four-electrode rectangular tube with the same cross-sectional area as the flap to measure the myocardial resistivity along the fiber direction. Repeated measurements showed considerable change and variation of myocardial resistivity in the rectangular tube. We speculate that ischemia caused deviation of the myocardial resistivity [6], [19], [20]. As another error source, the in-chamber blood, which has different resistivity than myocardium, can perturb the myocardial resistivity measurement if the interelectrode distance or the insertion depth of the plunge electrodes is large compared with the myocardial thickness [21]. Our in-situ measurements with plunge probes [13] showed that the swine myocardial resistivity at low frequencies changed considerably after the pig’s heart stopped beating. The resistivity below 1 kHz could increase by 50% within 60 min and it could double after 100 min. The resistivity at 500 kHz and 1 MHz increased by less than 15% within 6 h after the pig died. The results of in-vitro measurements also depend on time as well as how the tissue has been treated before measurement. These time factors add to overall resistivity error when comparing tissue resistivity measured at different times.

We made several surface resistivity probes where the four electrodes were lying on the surface of an epoxy base. Repeated measurements on myocardium or agar phantoms with these surface probes showed possibly over 10% variations of measured resistivity. We suspect the variation came from the variation of the contact between the electrodes and the measured media. Similar error could occur with the surface point probe used by Steendijk et al. [4] if good contact could not be ensured. Steendijk et al. fixed the resistivity probe on the epicardial surface of the beating heart with a flexible silicone rubber suction cup whose inner space was maintained at a slight vacuum with a pump. We repeated this measurement with our own suction cup, but found vacuum-induced bruise on the epicardium under the cup, which has an unknown effect on the myocardial resistivity. Accordingly, we recommend not using surface probes. Instead, in our in-vivo measurement [13], we used plunge probes to cope with the contact variation error of surface probes.

When we measured the noise constant at Step 5, we assumed same statistical characteristics of the noise signals measured by inserting the plunge electrodes into the myocardium and by immersing the electrodes in a saline solution contained in a metal cup on the top of the open-chest animal’s heart. This assumption was based on the fact that there was no removing or adding of noise sources between the two conditions and the fact that the observed amplitude and time characteristics of the noise waveforms in the two conditions were similar. Note that this assumption is valid only when the noise remains stationary, which is a condition that cannot always be guaranteed in a clinical environment.

Our system is capable of measuring the phase angle as well as the magnitude of the tissue resistivity. However, in our in-vivo measurement of swine myocardial resistivity, we found that the digital oscilloscope sometimes failed to provide stable phase measurement if the interference noise became too large compared with the signal. When the phase measurement of tissue resistivity is desired in a noisy measurement situation, we suggest the use of extra circuits to filter out the noise or even the adoption of the more costly phase-sensitive detection technique used by Ackmann [3], instead of our low-cost measurement method, for more stable phase measurement. Nevertheless, similar procedures and formulas as those presented in this paper can be used to evaluate the error of the measured phase angle of the tissue resistivity.

V. CONCLUSION
We analyzed the performance of a commonly used tissue resistivity measurement configuration in terms of the measurement error of the tissue resistivity. We identified error sources and derived the analytical equation of the measurement error step by step following the tissue resistivity measurement procedure. We evaluated the error terms in the analytical equation by in-vivo or bench-top experiments. The total error in the measured resistivity varies depending on the accuracy of the instruments, the characteristic of the circuit, the stray capacitance of the signal wires, the temperature change during the measurement procedure, the tissue resistivity, the current magnitude during measurement, the electrode bending and surface condition of the four-terminal resistivity probe, and the stray capacitance between the measured object and ground. Our error
evaluation method remains applicable for different circuits and instruments, or different resistivity probe configurations.

REFERENCES


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