ApplicationNOTE



EIS of Organic Coatings and Paints

Introduction

All My Impedance Spectra Look the Same!

"I'm an experienced polymer chemist. I'm trying to use Electrochemical Impedance Spectroscopy (EIS) to predict the corrosion resistant properties of paints. I've recorded many EIS spectra on painted metal samples immersed in electrolyte. All (or most) of these spectra look the same, regardless of the changes that I makes in my paint formulation. I obviously cannot use the results to evaluate paint performance. What's going on?"

This very common complaint has at least two causes:

- 1) You have a very high quality paint with very reproducible EIS spectra.
- 2) You are attempting to make measurements that are beyond the capabilities of the potentiostat in your EIS system.

The second cause is more common. The spectra look the same because you are measuring the characteristics of your potentiostat, not those of the paint.

The rest of this application note describes the effects that a potentiostat can have on EIS measurements of coated metals. Gamry Instrument potentiostats will be used as examples, but the discussion will apply to EIS systems built around any potentiostat.

This note concludes with some specific recommendations for getting meaningful EIS spectra on coating systems that are difficult to measure.

Background

If you are not very knowledgeable concerning electrochemical instrumentation, look at Gamry's Applications Note entitled "Primer on Potentiostats". This Primer will introduce the terminology used in talking about potentiostats and some potentiostat concepts that are used in the remainder of this application note. You will also need a basic understanding of EIS to follow the discussion in this note. Experienced EIS users should have no problems with the level of the discussion. If you are less experienced or you want to brush up on your basics, look at Gamry's Application Note entitled "Basics of Electrochemical Impedance Spectroscopy".

Coating Capacitance

A capacitor is formed when a non-conducting medium, called the dielectric, separates two conducting plates. The value of the capacitance depends on the size of the plates, the distance between the plates and the properties of the dielectric. In the case of a coated metal immersed in electrolyte, the metal is one plate, the coating is the dielectric, and the electrolyte is the second plate.

The capacitance relationship is:

$$C = \frac{\varepsilon_0 \varepsilon_r A}{d}$$

With,

C = the capacitance

 ε_{0} = electrical permittivity

 ε_r = relative electrical permittivity (dielectric

A = surface of one plate

d = distances between two plates

Whereas the electrical permittivity is a physical constant, the relative electrical permittivity (dielectric constant) depends on the material. Table 1 gives you a few useful $\epsilon_r\,$ values.

Material	ε _r
vacuum	1
water	80.1 (20° C)
organic coating	2 - 7

Table 1. Typical Dielectric Constants (Relative Electrical Permittivity)

Notice the large difference between the dielectric constant of water and that of an organic coating. The capacitance of a coated substrate changes as it absorbs water. EIS can be used to measure that change.

Notice that the capacitance of a coating increases when the area of the coating increases and when the coating thickness decreases.

Equivalent Circuit Model - Perfect Coating

A metal covered with an undamaged coating generally has very high impedance. The equivalent circuit for this situation is in Figure 1.

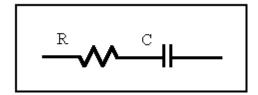


Figure 1. Purely Capacitive Coating

The model includes a resistor (due to electrolyte resistance) and the coating capacitance in series.

A Nyquist plot for this model is shown in Figure 2. In making this plot, the following values were assigned:

- $R \qquad = 500 \; \Omega \; (realistic \; only \; for \; a \; very \; poorly \; conductive \\ solution)$
- $C \qquad = 200 \ \text{pF} \ (\text{realistic for a 1 cm2 sample, a 25} \ \mu\text{M} \\ \text{coating, and} \ \epsilon_r = 6)$
- $F_i = 0.1 \text{ Hz}$ (lowest scan frequency -- a bit higher than typical)
- $F_f = 100 \text{ kHz}$ (highest scan frequency)

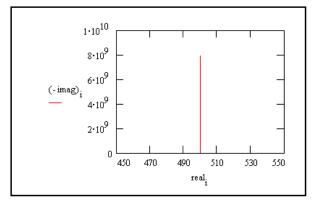


Figure 2. Typical Nyquist Plot for an Excellent Coating

The value of the capacitance cannot be determined from the Nyquist plot. It can be determined by a curve fit or from an examination of the data points. Notice that the intercept of the curve with the real axis gives an estimate of the solution resistance.

The highest impedance on this graph is close to 10^{10} W. This is close to the limit of measurement of many EIS systems.

The same data are shown in a Bode plot in Figure 3. Notice that the capacitance can be estimated from the graph but the solution resistance value does not appear on the chart. Even at 100 kHz, the impedance of the coating is higher than the solution resistance.

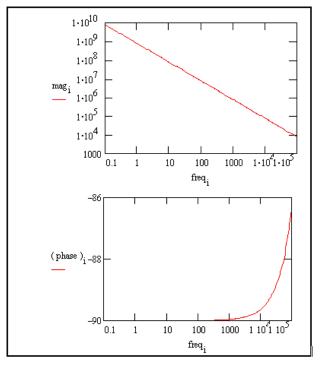


Figure 3. Typical Bode Plot for an Excellent Coating

Equivalent Circuit Model - Real Coating

The impedance behavior of a purely capacitive coating was discussed above. Most coatings degrade with time, resulting in more complex behavior.

After a certain amount of time, water penetrates into the coating and forms a new liquid/metal interface under the coating. Corrosion phenomena can occur at this new interface.

The impedance of coated metals has been very heavily studied. The interpretation of impedance data from failed coatings can be very complicated. Only the simple equivalent circuit shown in Figure 4 will be discussed here.

Even this simple model has been the cause of some controversy in the literature. Most researchers agree that this model can be used to evaluate the quality of a coating. However, they do not agree on the physical processes that create the equivalent circuit elements. The discussion below is therefore only one of several interpretations of this model.

Cc represents the capacitance of the intact coating. Its value is much smaller than a typical double layer capacitance. Its units are pF or nF, not μ F. Rpo (pore resistance) is the resistance of ion conducting paths that develop in the coating. These paths may not be physical pores filled with electrolyte.

On the metal side of the pore, we assume that an area of the coating has delaminated and a pocket filled with an electrolyte solution has formed. This electrolyte solution can be very different from the bulk solution outside of the coating. The interface between this pocket of solution and the bare metal is modeled as a double layer capacity in parallel with a kinetically controlled charge transfer reaction.

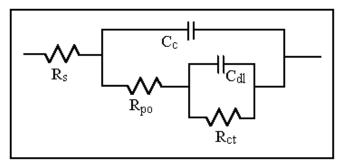


Figure 4. Equivalent Circuit for a Damaged Coating

When you use EIS to test a coating, you fit a data curve to this type of model. The fit estimates values for the model's parameters, such as the pore resistance or the double layer capacitance. You then use these parameters to evaluate the degree to which the coating has failed.

In order to show a realistic data curve, we need to do this operation in reverse. Assume that we have a 10 cm2 sample of metal coated with a 12 μ M film and that we have five delaminated areas. 1% of the total metal area is delaminated. The pores in the film that access these delaminated areas are represented as solution filled cylinders with a 30 μ M diameter.

The parameters used to develop the curves are shown below:

- C_c = 4 nF (Calculated for 10 cm² area , ϵ_r = 6 and 12 μ M thickness)
- $R_{po} = 3400 \Omega$ (Calculated assuming k = 0.01 S/cm)
- $R_s = 20 \Omega$ (Assumed)
- C_{dl} = 4 μ F (Calculated for 1% of 10 cm² area and assuming 40 μ F/cm²)
- $R_{ct} = 2500 \ \Omega$ (Calculated for 1% of 10 cm² area using Polarization Resistance at 1 mm/year and assumed constants)

With these parameters, the Nyquist plot for this model is shown in Figure 5. Notice that there are two welldefined time constants in this plot.

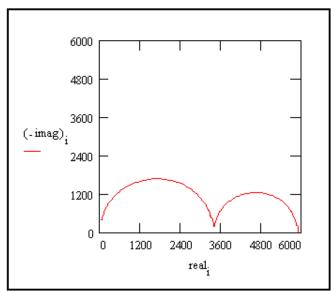


Figure 5. Nyquist Plot for a Damaged Coating

The Bode plot of the same data is shown in Figure 6. The two time constants are visible but less pronounced on this plot.

The Bode plot does not go high enough in frequency to measure the solution resistance. In practice this is not a problem, because the solution resistance is a property of the test solution and the test cell geometry, not a property of the coating. Therefore, it is usually not very interesting when you are testing coatings.

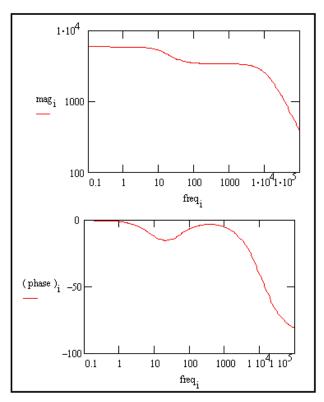


Figure 6. Bode Plot for a Damaged Coating

Problems in Measurement of Small Signals

Thick, high quality coatings characteristically have almost infinite resistance and very low capacitance.

It is obvious that their high resistance results in very small currents, especially at low frequencies where resistive elements in the models dominate. On a more subtle level, their low capacitance results in small AC currents. For example:

The impedance of a 10 nF capacitor at 1 kHz is 16 k Ω . With a 10 mV excitation at this frequency a potentiostat measures 630 nA.

The impedance of a 10 pF capacitor (often representative of a thick coating) at 1 kHz is 16 M Ω . With a 10 mV excitation, the potentiostat has to measure 630 pA.

Basic physics and the realities of electronics design and construction make it difficult to measure small currents. The problem is especially severe for small AC currents at high frequencies.

The result of these limitations is discussed in the following sections.

EIS – Open Lead Experiment

There is a very simple test you can run to test the limits of your potentiostat and its associated EIS system. Record an EIS spectrum with no cell attached. We call this test the "Open Lead Experiment". Gamry's Applications Note entitled "Quick Check of EIS System Performance"" discusses the methods used to produce this plot.

The EIS spectrum recorded in an Open Lead Experiment using a Gamry Instruments' PC4/300 with a 10 mV excitation voltage is seen in Figure 7.

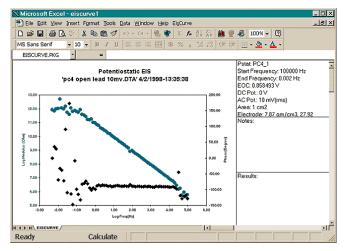


Figure 7. Open Lead EIS Bode Plot – PC4 with 10 mV Excitation

The open lead Bode plot looks like a noisy spectrum for a parallel RC network. This shape is seen in the open lead spectrum of every EIS system that we have tested. The diagonal line in the magnitude plot corresponds to a capacitor. The horizontal line at low frequency in the magnitude plot is equivalent to a resistor.

The open lead spectrum is fairly repeatable for a given hardware/software system. However, different potentiostats (with the same model number) may show variations in the spectrum, especially in the low frequency region. Differences of one half decade in impedance are not uncommon.

You cannot measure impedances that lie above the open lead spectrum. The EIS300/PC4 cannot measure $10^9 \Omega$ at 10 kHz since the system with no cell measures only $10^7 \Omega$ at this frequency.

The open lead spectrum is dependent on a number of factors. Shielding and grounding, excitation amplitude and DC level are some of the most important.

Accuracy Contour Plot – PC4/300 Potentiostat

NOTE: Achieving these specifications in your experiment may require very careful shielding of the cell and special cell design.

The results of the open lead experiment and a few additional tests can be used to generate a very useful graph that we call the "Accuracy Contour Plot". The accuracy of your impedance measurements can be predicted from this graph. An accuracy contour plot for the PC4 Potentiostat and the EIS300 EIS software can be seen in Figure 8.

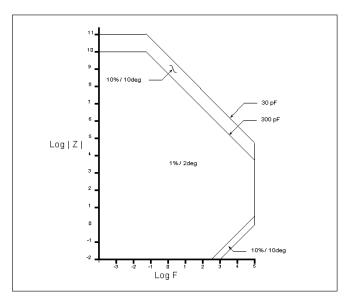


Figure 8. Accuracy Contour Map – EIS300/PC4 with 10 mV Excitation

This map applies when the EIS300 is operating with an PC4 Potentiostat in controlled potential mode. The AC excitation is 10 mV and a high quality Faraday shield surrounds a cell isolated from earth ground.

On the contour plot, each impedance measurement is a single point, defined by the frequency and the measured impedance at that frequency. Notice that the enclosed areas in the map are labeled with two numbers. They are the maximum error in percent of reading and maximum phase error at any point in the labeled region. For example, at 100 mHz this system can measure 10^9 Ω with errors of less than 1% in magnitude and 2° in phase. At $5 \times 10^{10} \Omega$, still at 100 mHz, the errors can be larger – 10% and 10°. Above $10^{11} \Omega$, the accuracy is unspecified, even though the instrument may function.

Notice the diagonal boundary lines that are labeled with equivalent capacitor values. You cannot measure a capacitor smaller than 30 pF, unless you are willing to accept errors of greater than 10% and 10°.

The accuracy contour plot shown above only applies to isolated cells. It does not apply when the potentiostat is used to make measurements on earth grounded "real world" systems such as highway bridges or pipeline probes. All cells with a connection to earth ground will severely degrade the system performance. Degradation of two orders of magnitude in impedance is common.

An accuracy contour map is valid for only one excitation (AC) amplitude. The map in Figure 8 applies at 10 mV excitation amplitude. In most cases, increasing the amplitude shifts the limits on the map upwards. At 100 mV excitation, the boundaries of the 10%, 10 degree region are 20 pF and $5 \times 10^{11} \Omega$. The minimum capacitor limitation is almost independent of amplitude. The low frequency resistance is a stronger function of excitation amplitude.

You may have noticed that the accuracy contour map has additional diagonal lines in its lower right hand corner. This region is generally not of interest in EIS on coatings, so it will not be discussed here. Contact us if you encounter this limitation in your work.

Accuracy Contour Map – Other EIS Systems

The accuracy contour plot shown in Figure 8 only apples to the Gamry Instruments EIS300 running a PCI4/300 Potentiostat with 10 mV of potential excitation. If you have a different system, you need a different map. Gamry's Application note entitled "Accuracy Contour Plots" includes instructions for estimation (or measurement) of the accuracy contour map for your EIS system.

If you don't want to go to the trouble of measuring the accuracy contour plot, we can offer some guidelines. Older potentiostats, such as the Gamry Instruments PC3 or the Ametek PAR 273 will be worse than the PC4 by one or two decades in impedance. Their DC limit at 10% accuracy and 4 degrees of phase error will be around $10^9 \Omega$ and their capacitor limit will be at about 300 pF (or worse).

Note: Some EIS manufacturers quote system performance with high excitation amplitudes and high error bands. Be careful, a system may claim to measure $10^{12} \Omega$, but the manufacturer may not tell you that that measurement requires 5 volts of excitation and/or allows 50% error.

Specialized potentiostats optimized for low current measurements are available. The Gamry Reference 600 is one example. It offers a DC limit of $10^{12} \Omega$ at 10% accuracy and a capacitive limit of 16 pF.

Good Measurements Aren't Easy

Careful experiment design is required if you expect optimal performance from your EIS system. The following hints may prove helpful.

Faraday Shield

A Faraday shield surrounding your cell is mandatory for very low level measurements. It reduces both current noise picked up directly on the working electrode and voltage noise picked up by the reference electrode.

A Faraday shield is a conductive enclosure that surrounds the cell. The shield can be constructed from sheet metal, fine mesh wire screen, or even conductive plastic. It must be continuous and completely surround the cell. Don't forget the areas above and below the cell. All parts of the shield must be electrically connected.

The shield must be electrically connected to the potentiostat's ground terminal.

Avoid External Noise Sources

Try to keep your system away from electrical noise sources. Some of the worst are:

- Fluorescent lights
- Motors
- Radio transmitters
- Computers and Computer Monitors

Avoid AC powered or computerized apparatus within your Faraday shield.

Cell Lead Length and Construction

Your cell leads must have a resistance higher than that of the impedance you are trying to measure. If you use coaxial cable we recommend a virgin Teflon dielectric. Long leads can severely degrade the AC response of your potentiostat.

Lead Placement

Many coating tests involve cells with capacitances so small that the capacitance between the potentiostat's leads can result in an error. Alligator clips can have 10 pF or more of mutual capacitance if they are run alongside each other.

If you wish to avoid excessive capacitance due to lead placement,

- Place the Working (and Work Sense) leads as far as possible from the other cell connections.
- Have the leads approach the cell from different directions.
- Remove alligator clips from the leads. In extreme cases you can replace banana plugs and pin jacks with smaller connectors.

The cell leads must not be moved during an experiment which measures small currents. Both microphonic and triboelectric effects can create spurious results when the cell cables are moved.

Cell Construction

Make sure that your cell construction does not limit your response. A cell where the resistance of the insulating material between the electrodes is only $10^{10} \Omega$ cannot be used to measure $10^{12} \Omega$ impedances. In general, glass and Teflon are the preferred cell construction materials.

You also must worry about shunt capacitance. Make the "inactive" portion of your electrodes as small as possible. Avoid placing electrodes close together and parallel with each other.

How to Measure an Impossible System

What can you do if you are faced with a sample that will produce data outside the defined regions of the accuracy contour map? These suggestions may help.

AC Amplitude

Larger AC amplitude may help you make difficult measurements. As discussed above, increasing the amplitude can move the low frequency limits in the accuracy contour map upwards. It has less effect on the minimum capacitance.

One concern is that the electrical field created by the excitation will cause failure in the paint. A five volt excitation across a 25 micron coating creates a field of 200 kV/meter. Most bulk plastics (PVC is an exception) claim dielectric strengths in excess of 12 MV/meter. Assuming coatings are one tenth as good as bulk plastics, dielectric breakdown should not be a factor unless the coating thickness is less than 5 microns.

Electrode Area

Electrode area is a critical experimental parameter. In general, EIS measurements on coatings should use as large an area as possible. Increasing the area has several beneficial effects:

- The capacitance of a point film is directly proportional to the sample area. If one cm² of a paint has an un-measureable capacitance (for example ten pF), 100 cm² of the same film will have an easily measureable 1 nF capacitance.
- If the paint has a uniform resistance, the resistance of a sample is inversely proportional to the sample area. Make the sample 100 times bigger and the resistance falls by a factor of 100.
- Some paints have only a few, widely separated, defects. Increasing the area increases the chance that a defect will be present in the sample.

Get a Low Current Potentiostat

If the suggestions above are impracticable, your only choice is get a potentiostat optimized for low current measurement. Some newer potentiostat can measure capacitances as low as three pF and low frequency resistances as high as $10^{13} \Omega$ (with one volt excitation).

The Gamry Instruments Reference 600 offers this type of performance at a very reasonable price.

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