1 Design Goal

This lab is not a design lab but a measurement and modeling lab. The goal is to provide equivalent circuits for two different types of electrodes in salt water: stainless steel and silver/silver-chloride.

You will also learn how to make your own silver/silver-chloride electrodes out of fine silver wire.

2 Background

Read the “Electrodes” chapter of Wikipedia book for the course:

Polarizable and nonpolarizable electrodes

http://en.wikipedia.org/wiki/Ideally_polarizable_electrode

A non-polarizable electrode has no polarization, current flows freely, and the electrode-electrolyte interface behaves like a resistor. A platinum-hydrogen electrode is considered a good non-polarizable electrode, but platinum in an NaCl solution (the usual case for bio-electrodes) is highly polarizable. A non-polarizable electrode that can be used in salt solutions is the silver/silver-chloride electrode, which we’ll look at in this lab.

A polarizable electrode ideally has no DC current flow between the electrode and the surrounding electrolyte. The electrode/electrolyte interface behaves like a capacitor. Platinum in NaCl is highly polarizable, but is too expensive for us—we’ll look at stainless steel electrodes, which appear to be moderately polarizable.

Real electrodes have characteristics that don’t match either ideal, and we’ll try to characterize that non-ideal behavior in this lab. Later on this quarter, we’ll use Ag/AgCl electrodes for the EKG.

A more comprehensive source is Bioimpedance and bioelectricity basics by Sverre Grimnes and Ørjan Grottem Martinsen, particularly Chapters 2 (Electrolytics) and 3 (Dielectrics).

Silver/silver chloride

The electrodes most commonly used in the nanopipette and nanopore labs are Ag/AgCl electrodes. These are relatively cheap to make and are non-polarizing. The reaction is $Ag \rightarrow Ag^+ + e^-; Ag^+ + Cl^- \rightarrow AgCl$. Note that both the silver and the silver chloride have low solubility, and so remain on the electrode—the flow of current (electrons in the wire and chloride ions in the solution) affects the ratio of silver to silver chloride on the surface of the electrode.
You will be making your own silver/silver-chloride electrodes by electroplating AgCl onto fine silver wires. Note: “fine silver” refers to silver that is at least 99.9% pure silver—there are many other grades of silver, such as “sterling silver” that is 92.5% silver.

Stainless steel

Surgical steel (such as type 316L stainless steel) is commonly used for implantable devices, because of its strength, its resistance to corrosion in the body, and very low toxicity. It is not commonly used for electrodes, though, and we’ll try to see why in this lab.

Because stainless steel is a complicated alloy, it is not clear what the electrochemical reactions are in salt water. The resistance to corrosion that gives it the “stainless” name comes mainly from a layer of chromium oxide on the surface of the steel. This layer forms naturally in oxidizing conditions, but in reducing conditions it may not form, and the steel behaves differently, corroding much more readily. (The two states are referred to as “passive” and “active” in discussions of the corrosion of stainless steel, and the half-cell potentials are different.)

The most important reactions for stainless steel as an electrode are probably $Fe + 2Cl^- \rightarrow FeCl_2 + 2e^-$, $Fe \rightarrow Fe^{2+} + 2e^-$, and $2H_2O + 2e^- \rightarrow 2H_2 + H_2O$, but I’m not certain of that.

http://en.wikipedia.org/wiki/Surgical_steel

3 Pre-lab assignment

Review your electrochemistry and read up on Ag/AgCl electrodes. Try to find some useful on-line sources other than those mentioned in the background section of this lab handout—be sure to cite them properly in your lab writeup.

Compute the surface area of the electrodes you will use.

- The stainless electrodes are about 2.7cm long to the plastic spacer and 1/8” (3.18mm) in diameter. The spacer should hold them 2cm apart. (I’ll bring calipers and a micrometer to lab, if anyone wants to do more careful measurements.)

- The fine silver wire in your kits is 24-gauge AWG (look up the diameter), and you will be folding it around a wire holder that is 4.4mm thick and you will be immersing the wires to a depth of 1cm, 2cm, or 3cm (depending which calibration line you use on the electrode holder), so the length will be 2.44cm, 4.44cm, or 6.44cm.

Make a gnuplot function for computing the magnitude (absolute value) of impedance for a pair of electrodes as a function of frequency and the three parameters $R_1, R_2, C_1$. Gnuplot is perfectly capable of dealing with complex numbers, but you might want to define $j = \text{sqrt}(-1.)$ to make the formulas easier to write. Figure 2 has some gnuplot commands that may be useful.

Each electrode/electrolyte interface could be modeled with the circuit show in Figure 1. The two electrode-electrolyte interfaces are in series, back-to-back, so the half-cell potentials cancel (assuming the same electrolyte concentration at both electrodes). Because we are cancelling out the half-cell potentials, we can lump the two electrode-electrolyte interfaces together, and just model the whole electrode-electrolyte-electrode system as a resistor in series with a parallel arrangement of a capacitor and resistor. This is the standard model used for EKG measurements, for example.

Make another gnuplot function that models the magnitude of impedance with a power law: $|Z| = af^b$. You will use these two functions for fitting models to your data.
Figure 1: Simplified equivalent circuit for a polarizable electrode. The half-cell potential is uncertain—different sources give different potentials for stainless steel in salt water, depending on the measurement conditions and what reference half cell is used. Luckily, for our application the half-cell potentials cancel. $R_1$ and $C_1$ correspond to the electrode/electrolyte interface, and $R_2$ to the bulk resistance of the electrolyte—it should decrease roughly inversely to the concentration of the ions in solution.

$$j = \sqrt{-1}$$

# impedance of two impedances in parallel
$\text{parallel}(z1,z2) = z1*z2/(z1+z2)$
# note: there is no need for a "series" function, since that is just
# $z1+z2$

# impedance of a capacitor of C Farads at f Hz
$Z_C(C,f) = 1./(j*2*\pi*f*C)$

# impedance of an inductor of L Henries at f Hz
$Z_L(L,f) = j*2*\pi*f*L$

# There is no need of a function for resistors, since $Z_R(R,f)=R$ and
# the resistance can just be used directly.

Figure 2: Some useful assignments and functions for building circuit models in gnuplot.
Do some plotting of the functions with different parameters, so that you get a feel for what each parameter does to the shape of the curve. You will probably have to give gnuplot some initial values for fitting your data, so it would be a good idea to have some feeling for what the parameters mean.

4 Parts, tools, and equipment needed

Parts for this lab from kit:

- alligator clip leads (made in lab 1)
- resistors
- 1 foot fine silver wire 24 gauge
- 2 EKG electrodes

Parts students need to provide on their own:
None

Tools for this lab:
None

Equipment in lab:

- function generator
- two multimeters
- electrode holders for silver wire
- cups for salt water (3 per station)
- secondary containment tubs
- sponges in case of spills (there’d better not be any!)
- stock NaCl solutions (1M, 0.1M, 0.01M) (about 75ml/setup, or 1l of each)
- NaCl solution for electroplating (1l of 1M)
- paper towels
- (optional) calipers and micrometer

5 Procedures

Warning: salt water and expensive electronics equipment do not play nicely together. All liquids must be kept contained in the secondary containment tubs throughout the lab. Do not remove the cups from the tubs!
Characterizing stainless steel electrodes

Immerse the electrodes in 1M NaCl solution, up to the plastic spacer, as shown in Figure 3, so that you have a known area of electrode submerged. You can rest the alligator clips on the rim of the cup to help stabilize the electrodes.

First, measure the DC resistance of the electrodes with the ohmmeter.

Next, connect a voltmeter across the electrodes, an ammeter in series, and the whole thing to the function generator, so that you can simultaneously measure the RMS AC voltage and current. Actually, the ammeters we have only measure RMS current up to 5kHz, so you’ll get better results if you replace the ammeter with a 1Ω or 10Ω resistor and a voltmeter, as the voltmeter function is pretty accurate to 50kHz, and usable up to 1MHz.

Measure the RMS voltage and RMS current for several frequencies from 3Hz to 300kHz. You should measure at several frequencies per decade, roughly uniformly spaced in log space (say, 3, 5, 7.5, 10, 15, 20, 30, …, 3E5). Sorry, you’ll have to record this by hand, as we don’t have the time to set up the equipment with LabView and automate the test. It may help to have one person making the measurements and calling out the numbers to their partner who records the numbers in a file.

Plot the magnitude of the impedance (V/I) as a function of frequency (use a log scale for frequency!). Fit both your models to the data. Which is the better fit?

Repeat the measurements for the other two salt concentrations, and make one plot with the three sets of data points and 6 curves (both models for each of the three concentrations).

Electroplating silver wire with AgCl

Wrap the silver wire around the electrode holder as shown in Figure 4. Use alligator clip leads to connect the electrodes through an ammeter to a power supply.

Put the electrode holder upright in a cup within the secondary containment tub, and carefully add 1M NaCl solution until it is past the shoulders but not deep enough for the alligator clips to
Figure 4: Holder showing wire wrapped around it. Cut the silver wire into two pieces 6” long. Wipe any finger grease off of it with a paper towel. Wrap one wire around the left ear, through the left armpit, and back up around the ear again. Do the same with the other wire on the right ear. (Note: the wire in this picture is about 7” long—you’ll have no loose ends with a 6” wire.)

touch the solution. Turn on the power with a very low voltage, and gradually increase the voltage until you have a current of about 1mA for each square centimeter of area for one wire (you did precompute the area of the wire in the pre-lab, didn’t you?). After a minute or two you should have a fairly uniform grey coating of AgCl on the positive wire.

Reverse the polarity and plate the other wire also. You may want to reverse the polarity and re-plate the first wire for about half as long, to make the coatings more consistent.

If your coatings are mottled, try wiping finger grease off with a paper towel and plating some more. If you were doing this lab with real biological samples, the wires would probably be cleaned with acetone or other organic solvent, and rinsed with deionized water, both before and after plating.

There is another method for producing the AgCl layer, by immersing the fine silver wires in chlorine bleach, but bleach is a bit too hazardous for a lab that has no sink and no eyewash station nearby. The electroplating method supposedly produces a thicker, more uniform coating of AgCl anyway.

**Characterizing Ag/AgCl electrodes**

Immerse the plated electrodes in 1M NaCl (preferably not the same solution as used for electroplating, which may be somewhat depleted of Cl\(^-\) ions), up to one of the calibrated lines on the electrode holder (1cm, 2cm, 3cm at the shoulders). It may be easiest to hold the holder upright in the cup first, and gradually pour in liquid.

Do the same characterization experiments with the Ag/AgCl electrodes that you did with the stainless steel electrodes.
Characterizing EKG electrodes

Take two of the disposable EKG electrodes and stick them together face-to-face, trying to seal them together all the way around. Clip alligator leads onto them and try to characterize the EKG electrodes the same way you did the home-made Ag/AgCl electrodes.

6 Demo and writeup

Your report should include the schematic diagram of the equivalent circuit for each type of electrode, the plots and fits of the magnitude of electrode impedance, and your interpretation of the results. The writeup should be understandable by another bioengineer who has not read this assignment.

Some things to discuss:

- Which models fit the data best?
- What electrodes would you use and what frequency would you use if you were trying to measure the concentration of salt water electronically?
- How would you modify the measurement technique to get the full complex impedance, rather than just the magnitude of it?

7 Design Hints

Since this is a measurement lab, rather than a design lab, the main things to think about are how to make and record the large number of measurements efficiently.