# **BENG 186B Principles of Bioinstrumentation**

### Week 7 Review

## **Exercises**

Selections from:

2015 Homework 5 2015 Homework 6

#### **BENG 186B Winter 2015 HW #5**

Due Thursday February 26 at the beginning of class

1. One of the major problems during dialysis after bleeding out is the introduction of air bubbles as those can lead to embolism, which can result in cardiac arrest depending on where the embolism lodges. A safety check within the machine is to incorporate a bubble detector to warn clinicians on staff if bubbles have formed. As an engineer testing such a detection system, you receive the following signals from the system driven with a signal generator to mechanically stimulate test samples under two conditions: with and without bubbles. In each case the mechanical actuation by the signal generator couples to a membrane with compliance  $C_d = 0.001 \text{ Pa}^{-1}$  in fluidic contact with the test sample.



- (a) Assuming that the amplitude of the signal generator voltage is kept the same in both tests with and without bubbles, and the signal generator is tuned to operate at the natural frequency in each case, determine which of the signals A and B correspond to the test with or without bubbles respectively. Explain why.
- (b) Determine the compliance of bubbles  $(C_b)$  with the parameters from the given plot. Assuming that  $C_b$  is directly proportional to the total volume of air inside the test system, explain how you can use the bubble detector signal to calculate the bubble density.

2. For many biological problems, cell sorting is an important task of differentiating and categorizing different cell types within a heterogeneous mixture. Utilizing the Hall Effect phenomena, you design a system where you can separate cells of interest by using charged antibodies that attach specifically to the antigens of your cells. Since you have two cell types to separate, you design negatively and positively charged antibodies.

Due to the separation of charges, the tube through which the cells flow acts like a parallel plate capacitor with a distance of 5 mm. The cells are flowing at 100 mL/s through the tube with a relative permittivity of 80. You induce the Hall Effect after generating a magnetic field of 0.25 T.

- (a) Draw the system with the corresponding vectors for the generated magnetic field, flow of cells, and electric field.
- (b) Assuming each cell to have a valance of either +1 or -1, calculate the concentration of cells of either type covering the walls of the tube.
- 3. You have a device that can measure blood velocity using ultrasonic pulses. This requires the user to position the source and receiver correctly.
  - (a) Given the following different setups, determine the expected frequency shifts when using an ultrasound sound system to measure the blood velocity.



(b) To calculate the Doppler shift, you need to assume the direction of the blood velocity (shown directed to the right in the above examples). If your assumption is wrong, how will that reflect in your calculated Doppler shift?

4. **Design problem**: Design an automated sphygmomanometer that includes a digital readout of systolic and diastolic pressure and heart rate.

You have these components available:

- (a) Cuff with arbitrary digital control over applied pressure;
- (b) Sonic transducer contacting skin over artery inside the cuff, with digital readout of the acquired sound waveform;
- (c) Alphanumeric display;
- (d) Programmable microcontroller;
- (e) Start button and sleep button;
- (f) Power supply with power on/off switch.

Sketch a diagram of the system, and outline the algorithm running on the microcontroller to control the cuff, acquire the sound signal, and compute and display the systolic pressure, diastolic pressure, and heart rate.

You do not need to write actual code, but be quantitative in your description with sufficient information for a programmer to be able to complete the design. You may use timing diagrams and equations to define the sequence of control variables, the corresponding measured quantities, and the computation of the output variables.

Define important parameters in the algorithm, and give numerical values based on the physiological range that you expect.

#### **BENG 186B Winter 2015 HW #6** Due *Thursday March 12 at the beginning of class*

- 1. The lab technicians are running a routine checkup on a patient's blood sample using some electrodes connected to an aterial gas meter. The meter measures the electrodes and calculates pH, PCO<sub>2</sub>, and [O<sub>2</sub>]. Unfortunately, the meters are broken. Fortunately, you come in to save the day as the biomedical engineer with a multimeter to measure the electrodes directly.
  - (a) You measure a voltage of -50 mV across the pH probe. The probe uses a saturated HCl concentration of 100 mM. What is the blood sample's pH?
  - (b) You calibrate the PCO<sub>2</sub> electrode with a standard (pH = 7.4, PCO<sub>2</sub> = 40 mmHg). Then, you put a sample of the blood from part (a) in the PCO<sub>2</sub> electrode with 25 mM NaHCO<sub>3</sub>. What is the sample's PCO<sub>2</sub>?
  - (c) For measuring  $PO_2$ , you construct a transimpedance amplifier to convert the current output from the Clark electrode to a voltage:



You set  $R_1 = 1 \text{ M}\Omega$ ,  $R_2 = 50 \text{ k}\Omega$ , and  $R_3 = 5 \text{ M}\Omega$ . Then, you flow the blood sample through a Clark electrode at 50 mL/s which has 0.7 V applied across it. You measure a voltage of 5.5 V at  $V_{\text{out}}$ . What is the sample's  $[O_2]$ ?

- 2. Pulse oximeters work by passing red light (660 nm) and infrared light (805 nm) through a body part (such as a finger). The absorbance is measured using a photodiode and an amplifier. The SpO<sub>2</sub> can be calculated from the ratio of the absorbances.
  - (a) Why do pulse oximeters use two different wavelengths to measure  $SpO_2$ ? Be sure to mention the isobestic wavelength's role in computing  $SpO_2$ .
  - (b) What is the origin of the DC component of the photodiode signal? What is the origin of the AC component?
  - (c) Shown below is a sample of the raw pulse oximeter data:



The  $SpO_2$  can be estimated by:

$$\operatorname{SpO}_{2} = \left(110 - 25 \times \frac{\operatorname{Abs}_{660\,\mathrm{nm}}}{\operatorname{Abs}_{805\,\mathrm{nm}}}\right)\%$$

What is the patient's  $SpO_2$ , according to the raw pulse oximeter data? *Hint:* Normalize the AC components against their corresponding DC components first. The AC component is the peak-to-peak voltage and the DC component is the average voltage.

(d) What is the patient's heart rate, according to the raw pulse oximeter data?

- 3. Cardiomyocytes have a typical membrane time constant of 2 ms and starts to fibrillate at above 100 mA.
  - (a) Suppose an unfortunate person gets struck with 200 kV lightning. The current enters and exits the body through skin ( $R_{skin} = 100 \text{ k}\Omega$ ). How much current will flow through the victim? Assume the internal body resistance is negligible compared with the skin resistance.
  - (b) If 20% of the lightning strike's current mentioned in part (a) passes through the heart, how long can the victim withstand the strike before suffering cardiac arrest?
  - (c) Lightning strikes aren't the only electrical hazard to worry about. Ground faults can make medical equipment dangerous to use without any outward signs of a problem.

You are designing a 12–lead ECG powered by a 120 VAC supply. Because you are a competent engineer, you add resistors in series with each ECG electrode lead wire as part of the device's overall safety design. Using the graph below, specify what resistor value you should use so that any ground fault currents are imperceptible.



AC-1 imperceptible

AC-2 perceptible but no muscle reaction

AC-3 muscle contraction with reversible effects

AC-4 possible irreversible effects

AC-4.1 up to 5% probability of heart fibrillation

- AC-4.2 5-50% probability of fibrillation
- AC-4.3 over 50% probability of fibrillation

Graph source: International Electrotechnical Commission (IEC) standard 60479

4. **Design problem:** A group of athletes would like to know their metabolic rates during exercise. A non-invasive way to estimate this is to measure the CO<sub>2</sub> and O<sub>2</sub> concentration in their breath as they train.

Your task is to design a circuit to interface a Severinghaus electrode (measures  $PCO_2$ ) and a Clark electrode (measures  $PO_2$ ) to a microcontroller. This microcontroller is already loaded with software that computes the metabolic rate based on  $PCO_2$  and  $PO_2$ . You have:

- The Severinghaus electrode. Its usable output range is -250 mV to 250 mV.
- The Clark electrode. Its usable output current is  $0 \,\mu A$  to  $100 \,\mu A$ . This electrode needs  $0.7 \,V$  applied to it.
- Op-amps, and any resistors/capacitors as needed.
- A stable  $\pm 5$  V power supply.
- A microcontroller with a built-in ADC. The ADC can only measure positive voltage.

The circuit needs to filter out signals above 10 Hz from both sensors. The ADC should read 0 V to 5 V linearly with respect to each electrode's usable output. You must include numerical values in your design. **BONUS**: Design your circuit in such a way that a - 5 V supply is not required.