Assignment 1 Theoretical Neuroscience [Gatsby]

TAs:

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Due: 31 January, 2020

1. The Hodgkin-Huxley neuron

Numerically integrate the Hodgkin-Huxley equations with matlab (or your favorite package). If you're using matlab, it's a good idea to use the Matlab ode45 function, or if you're using Python, scipy.solve_ivp. The equations are:

$$C\frac{dV}{dt} = -\overline{g}_{Na}m^{3}h(V - E_{Na}) - \overline{g}_{K}n^{4}(V - E_{K}) - \overline{g}_{L}(V - E_{L}) + I_{stim}$$
(1)

- $\frac{dx}{dt} = \alpha_x (1-x) \beta_x x \quad \text{where } x \text{ is } m, n \text{ or } h$ $\alpha_n (V) = 0.01 (V+55) / [1 \exp(-(V+55)/10)]$ (2)
 - (3)
- $\beta_n(V) = 0.125 \exp(-(V+65)/80)$ (4)

$$\alpha_m(V) = 0.1(V+40)/[1 - \exp(-(V+40)/10)]$$
(5)

$$\beta_m(V) = 4 \exp(-(V+65)/18) \tag{6}$$

$$\alpha_h(V) = 0.07 \exp(-(V+65)/20) \tag{7}$$

$$\beta_h(V) = 1/\left[\exp(-(V+35)/10) + 1\right]$$
(8)

 $\mathrm{Let}\ C = 10\ \mathrm{nF}/\mathrm{mm^2}, \overline{g}_L = .003\ \mathrm{mS}/\mathrm{mm^2}, \overline{g}_K = 0.36\ \mathrm{mS}/\mathrm{mm^2}, \overline{g}_{Na} = 1.2\ \mathrm{mS}/\mathrm{mm^2}, E_K = -77\ \mathrm{mV}, E_L = -77\ \mathrm{$ -54.387 mV, and $E_{Na} = 50$ mV. Use an integration time step of 0.1 ms.

Remember to keep your units consistent. F/S = Farad/Siemens = 1 second.

- (a) Run the simulations with $I_{stim} = 200 \text{ nA/mm}^2$. Plot the membrane potential (V) and gating variables (m, h, and n) versus time.
- (b) Write down expressions for the equilibrium values of the gating variables $(m_{\infty}, h_{\infty}, \text{and } n_{\infty})$, and plot them versus voltage.
- (c) Plot the firing rate versus I_{stim} , up to a firing rate of 50 Hz. The firing rate should jump suddenly from zero to a non-zero value. This is called a type II behavior. Type I behavior is when the firing rate begins at zero and increases continuously without any jumps.
- (d) What happens to the plot of firing rate versus I_{stim} as you decrease \overline{g}_K ?
- (e) Spikes are initiated at the axon hillock, where the axon meets the soma. This is because \overline{g}_{Na} is very high there. What happens to the plot of firing rate versus I_{stim} as you increase \overline{g}_{Na} ?

2. The linear integrate and fire neuron

An approximate treatment of spiking neurons is to think of them as passively integrating input and, when the voltage crosses threshold, emitting a spike. This leads to the linear integrate and fire neuron (sometimes called the leaky integrate and fire neuron, and often abbreviated LIF), which obeys the equation

$$C\frac{dV}{dt} = -g_L(V - \mathcal{E}_L) + I_0 \,.$$

This is just the "linear integrate" part. To incorporate spikes, when the voltage gets to threshold (V_t) , the neuron emits a spike and the voltage is reset to rest (V_r) .

(a) Compute the firing rate of the neuron as a function of I_0 . This firing rate will be parameterized by three numbers: \mathcal{E}_L , V_t , and V_r .

Hint #1: The firing rate is the inverse of the time it takes to go from V_r to V_t . Hint # 2: Changing variables, and defining new quantities, almost always makes life easier. For example, you might let $v = V - \mathcal{E}_L$ and define $V_0 \equiv I_0/g_L$ and $\tau \equiv C/g_L$.

- (b) Let $I(t) = g_L V_0 \sin(\omega t)$, $V_r = \mathcal{E}_L$, $V_t = \mathcal{E}_L + \Delta V$, and define $C/g_L \equiv \tau$. Start with $V_0 = 0$ and integrate for a long enough time that the neuron equilibrates. Then increase V_0 very slowly compared to the time constant, τ . Show that the neuron will start spiking repetitively when $V_0 > (1 + \tau^2 \omega^2)^{1/2} \Delta V$.
- 3. Warmup nullclines. Consider a model that is bound to come up again, in one form or another,

$$\tau_x \frac{dx}{dt} = -x + \tanh\left(\beta(x-y)\right)$$

$$\tau_y \frac{dy}{dt} = -y + \alpha x.$$

For all questions, assume $\alpha > 0$ and $\beta > 1$.

- (a) Draw the nullclines for an α and β of your choice.
- (b) What are the conditions on α and β for there to be three fixed points?
- (c) Assume α and β are such that there are three fixed points. Determine the stability of each of them. Draw trajectories starting near x = y = 0.
- (d) Assume α and β are such that there is one fixed point. Determine its stability. Draw trajectories starting near x = y = 0.
- 4. Hodgkin-Huxley nullclines. Consider a simplified Hodgkin-Huxley type model,

$$\tau \frac{dV}{dt} = -(V - \mathcal{E}_L) - hm(V)V$$

$$\tau_h \frac{dh}{dt} = h_\infty(V) - h$$

$$m(V) = \frac{1}{1 + \exp(-(V - V_t)/\epsilon_m)}$$

$$h_\infty(V) = \frac{1}{1 + \exp(+(V - V_h)/\epsilon_h)}$$

with parameters

$$\begin{aligned} \mathcal{E}_L &= -65 \text{ mV} \\ V_t &= -50 \text{ mV} \\ \epsilon_h &= 10 \text{ mV} \\ \epsilon_m &\ll 1 \text{ mV}. \end{aligned}$$

The remaining parameter, V_h , will be specified as needed (it will take on a range of values).

- (a) Sketch the nullclines in V-h space for $V_h = -60, -50$ and -40 mV. Put voltage on the x-axis and h on the y-axis. For each equilibrium, tell us whether it is stable or unstable, or hard to tell without a detailed stability analysis.
- (b) Find the condition on V_h that guarantees more than one equilibrium.
- (c) For a value of V_h (which you choose) such that there is more than one equilibrium, sketch the trajectories starting at V slightly greater than V_t and h = 1.

5. The passive cable equations.

Consider a passive cable with radius a, as shown here,



This is a bare-bones schematic; in addition to what's shown, there is an external current, $I_e(x,t)$, and a current associated with channels, $I_m(x,t)$.

We want to derive the cable equation, which we'll eventually restrict to the passive cable equation. We'll start with the equation for the membrane potential, V(x,t),

$$C \frac{\partial V(x,t)}{\partial t} = I(x - dx/2) - I(x + dx/2) - I_m(x,t) + I_e(x,t)$$
(50j)

where C is the capacitance of the piece of dendrite between the dotted lines. Next is the equation for the current,

$$I(x - dx/2) = \frac{V(x - dx) - V(x)}{R}$$
(50k)

where R is the resistance along the dendrite, between x - dx and x. A virtually identical expression holds for I(x + dx). Inserting these into the equation for the voltage yields

$$C \frac{\partial V(x,t)}{\partial t} = \frac{V(x-dx) - 2V(x) + V(x+dx)}{R} - I_m(x,t) + I_e(x,t).$$
(501)

(a) Verify that when you Taylor expand the voltage terms on the right hand side to lowest nonvanishing order, this becomes

$$C \frac{\partial V(x,t)}{\partial t} = \frac{dx^2}{R} \frac{\partial^2 V(x)}{\partial x^2} - I_m(x,t) + I_e(x,t).$$
(50m)

That's the cable equation! However, we want sensible answers in the limit $dx \to 0$. For that we need to know how C and R scale with dx.

(b) First, resistance. You may have learned in our physics class that resistance is proportional to length and inversely proportional to area – something that follows (with a little work) from I = V/R. It thus makes sense to define the resistivity of a material, here denoted r_L , via $R = r_L \times \text{length/area}$. For our setup (remember that the cylinder has radius *a*), this means

$$R = r_L \frac{dx}{\pi a^2}.$$
(50n)

Next, the capacitance. That scales with area: the more area for a given voltage, the more the charge. Thus, it makes sense to define the specific capacitance via $C = c_m \times \text{area}$. For our setup, the relevant voltage is across the dendritic walls, so the relevant area is $2\pi a dx$. We thus have

$$C = c_m 2\pi a dx. \tag{500}$$

Insert these into the equation for the membrane potential, and show that

$$c_m \frac{\partial V(x,t)}{\partial t} = \frac{a}{2r_L} \frac{\partial^2 V(x)}{\partial x^2} - \frac{I_m(x,t)}{2\pi a dx} + \frac{I_e(x,t)}{2\pi a dx} \,. \tag{50p}$$

(c) There's still a dependence on dx; to get rid of that we define the current densities

$$i_m(x,t) \equiv \frac{I_m(x,t)}{2\pi a dx} \tag{50qa}$$

$$i_e(x,t) \equiv \frac{I_e(x,t)}{2\pi a dx} \,. \tag{50qb}$$

Inserting these into the above equation almost gives us the passive cable equation. The last thing we need to do is write down an expression for i_m in terms of the voltage. We could use Hodgkin-Huxley type equations, but here we'll stick to passive channels. For that we'll write, as usual,

$$I_m = \frac{V - \mathcal{E}}{R_m} \tag{50r}$$

where \mathcal{E} is the reversal potential. Note that R_m is the resistance across the membrane. As usual, resistance is proportional to distance divided by area. However, we're mainly interested in the area dependence; distance is the thickness of the membrane, which is pretty much constant. We'll thus define

$$R_m = \frac{r_m}{2\pi a dx} \,. \tag{50s}$$

Here r_m depends on the membrane, but it's about the same for dendrites and neurons. Combining this with the equation for I_m , and taking into account the definition of i_m , we have

$$i_m = \frac{V - \mathcal{E}}{r_m} \,. \tag{50t}$$

Show that when you insert this into our current version of the cable equation, and multiply by r_m , you end up with the standard cable equation.