Homework 2- Single AdEx Neuron

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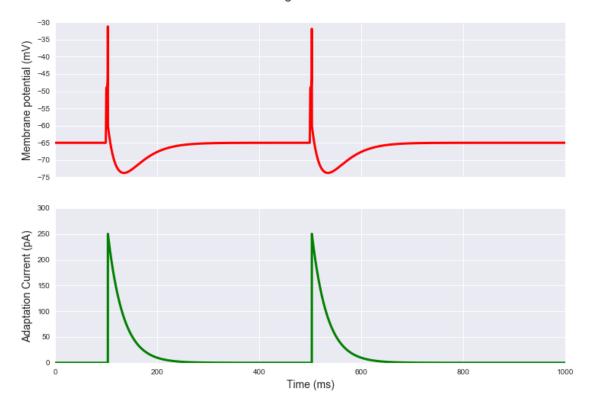
1 Question 1

```
In [2]: # Set parameters
        R = 100 * br.Mohm # membrane resistance
        tau_m = 24 * br.ms # membrane time constant
        u_rest = -65 * br.mV # resting potential
        u_r = -60 * br.mV # reset potential
        theta_reset = 40 * br.mV # reset threshold
        vartheta_rh = -52 * br.mV # rheobase threshold
        delta_T = 1 * br.mV # sharpness of ap upswing
        tau_w = 30 * br.ms # adaptation time constant
        a = 0 * br.nS # subthreshold adaptation constant
        b = 250 * br.pamp # adaptation jump after a spike
        eqs = '''
        du/dt = (-(u - u_rest) + delta_T*exp((u-vartheta_rh)/delta_T) -
                R*w + R*I) / tau_m : volt
        dw/dt = (a*(u - u_rest) - w)/tau_w : amp
        I = input_current(t) : amp
        \mathbf{t} \cdot \mathbf{t} \cdot \mathbf{t}
```

1.1 Part A

```
In [3]: input_current = np.zeros((1000,))
        input_current[[99, 499]] = 3900
        input_current = br.TimedArray(input_current*br.pamp, dt=1*br.ms)
       br.start scope()
        AdEx = br.NeuronGroup(1, eqs, threshold='u>theta_reset',
                              reset='u=u_r; w+= b', method='euler')
        AdEx.u = u_rest
        AdEx.w = 0 # * br.pamp
        rec = br.StateMonitor(AdEx, ('u', 'w'), record=True)
       br.run(1*br.second)
In [4]: # Plot membrane potential and adaptation current against time
        # for AdEx neuron.
        fig, axes = plt.subplots(2, sharex=True)
        fig.suptitle('Q1A: A Single AdEx Neuron', fontsize=18)
        axes[0].plot(rec.t/br.ms, rec.u[0] \star1000,
                     label='Membrane Potential (mV)', lw=3., c='r')
        axes[1].plot(rec.t/br.ms, rec.w[0]*10**12,
                     label='Adaptation Current (pA)', lw=3., c='g')
        axes[1].set_xlabel('Time (ms)', fontsize=14)
        axes[0].set_ylabel('Membrane potential (mV)', fontsize=14)
        axes[1].set_ylabel('Adaptation Current (pA)', fontsize=14)
        plt.show(fig)
```

Q1A: A Single AdEx Neuron



The membrane potential (u) increases when input current is applied, reaching the reset threshold and spiking. After each spike, the membrane potential decreases to -75 mV in the refractory period before returning to the resting potential.

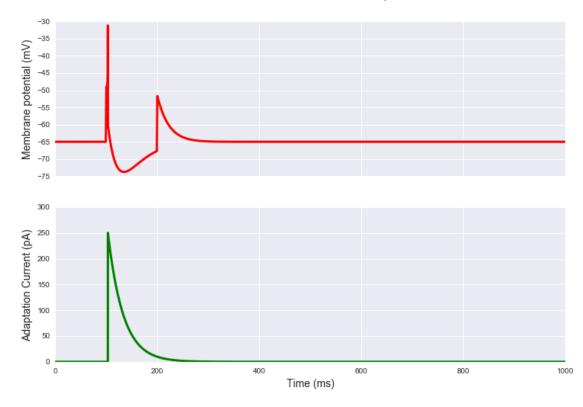
The adaptation current (w) peaks when a spike occurs and then degrades over time. The inputs are spaced out enough that the adaptation current drops back down to ~0 before the next input occurs, meaning that the adaptation did not affect the membrane potential significantly in this model.

The adaptation current influences the voltages only if inputs occur close enough together for the adaptation current not to drop to zero.

1.2 Part B

```
rec = br.StateMonitor(AdEx, ('u', 'w'), record=True)
br.run(1*br.second)
```

Q1B: AdEx With Shorter Gap

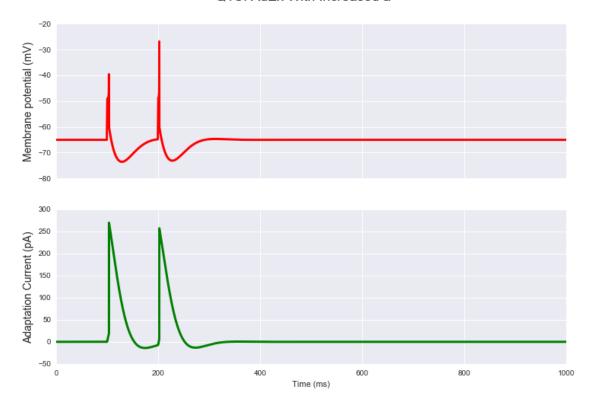


The decreased spacing between the inputs meant that the adaptation current had not dropped to zero. As a result, the adaptation current decreased the effect of the second input, preventing a spike. If we increased a or decreased tau_w , the adaptation current would decrease more quickly and a second spike would be possible so soon after the first spike.

1.3 Part C

```
In [7]: a = 10 * br.nS
       br.start_scope()
        AdEx = br.NeuronGroup(1, eqs, threshold='u>theta_reset',
                              reset='u=u r; w+= b', method='euler')
        AdEx.u = u rest
        AdEx.w = 0. * br.pamp
        rec = br.StateMonitor(AdEx, ('u', 'w'), record=True)
       br.run(1*br.second)
In [8]: # Plot membrane potential and adaptation current against time
        # for AdEx neuron.
        fig, axes = plt.subplots(2, sharex=True)
        fig.suptitle('Q1C: AdEx With Increased a', fontsize=18)
        axes[0].plot(rec.t/br.ms, rec.u[0]*1000,
                     label='Membrane Potential (mV)', lw=3, c='r')
        axes[1].plot(rec.t/br.ms, rec.w[0]*10**12,
                     label='Adaptation Current (pA)', lw=3., c='q')
        axes[1].set_xlabel('Time (ms)')
        axes[0].set_ylabel('Membrane potential (mV)', fontsize=14)
        axes[1].set_ylabel('Adaptation Current (pA)', fontsize=14)
        plt.show(fig)
```

Q1C: AdEx With Increased a



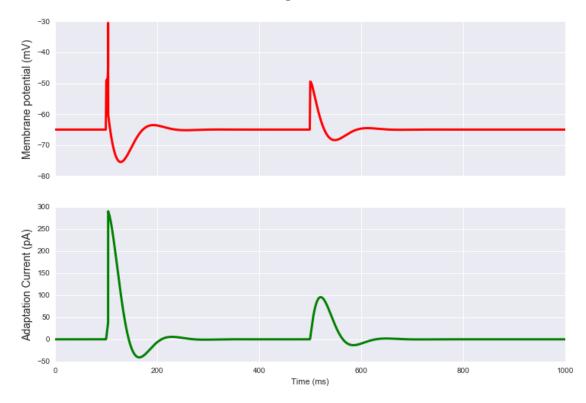
The positive value for *a* means that the adaptation current drops down to a negative value, unlike Q1B, and at a much faster rate than Q1B. As a result, the adaptation current is actually below zero, briefly decreasing adaptation, at the time of the second impulse current. The membrane potential is thus high enough at the time of the second impulse current to produce a spike, which has a greater amplitude than the first spike in spite of the fact that the impulse currents are the same.

2 Question 2

```
In [9]: # Set parameters
        R = 100 * br.Mohm # membrane resistance
        tau_m = 24 * br.ms # membrane time constant
        u rest = -65 * br.mV # resting potential
        u_r = -60 * br.mV # reset potential
        theta reset = 40 * br.mV # reset threshold
        vartheta_rh = -52 * br.mV # rheobase threshold
        delta_T = 1 * br.mV # sharpness of ap upswing
        tau_w = 50 * br.ms # adaptation time constant
        a = 30 * br.nS # subthreshold adaptation constant
        b = 250 * br.pamp # adaptation jump after a spike
        eas = '''
        du/dt = (-(u - u_rest) + delta_T*exp((u-vartheta_rh)/delta_T) -
                R*w + R*I) / tau_m : volt
        dw/dt = (a*(u - u_rest) - w)/tau_w : amp
        I = input_current(t) : amp
        \tau \cdot \tau \cdot \tau
```

2.1 Part A

Q2A: A Single AdEx Neuron



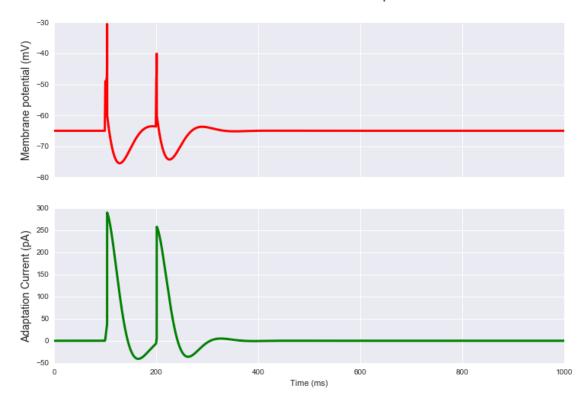
The second impulse current does not produce a spike (i.e., the membrane potential does not reach rheobase threshold), even though the adaptation current is zero at the time it is introduced. This means that the relationship between the neuron's resistance and firing threshold requires, at minimum, an impulse current >3800 pA and <= 3900 pA.

The increased value for tau_w results in a slower return of w to zero, while the increased value for a leads to a larger increase in the amplitude of w as a result of the impulse currents. The increased a also causes the adaptation current to decrease much faster after the impulse currents and to decrease below zero (causing a brief period of potentiation).

2.2 Part B

```
In [12]: # Run model
         input_current = np.zeros((1000,))
         input current [99] = 3900
         input_current[199] = 3800
         input current = br.TimedArray(input current*br.pamp, dt=1*br.ms)
         br.start_scope()
         AdEx = br.NeuronGroup(1, eqs, threshold='u>theta_reset',
                               reset='u=u_r; w+= b', method='euler')
         AdEx.u = u\_rest
         AdEx.w = 0 * br.pamp
         rec = br.StateMonitor(AdEx, ('u', 'w'), record=True)
         br.run(1*br.second)
In [13]: # Plot membrane potential and adaptation current against time
         # for AdEx neuron.
         fig, axes = plt.subplots(2, sharex=True)
         fig.suptitle('Q2B: AdEx With Shorter Gap', fontsize=18)
         axes[0].plot(rec.t/br.ms, rec.u[0]*1000,
                      label='Membrane Potential (mV)', lw=3., c='r')
         axes[1].plot(rec.t/br.ms, rec.w[0]*10**12,
                      label='Adaptation Current (pA)', lw=3., c='g')
         axes[1].set_xlabel('Time (ms)')
         axes[0].set_ylabel('Membrane potential (mV)', fontsize=14)
         axes[1].set_ylabel('Adaptation Current (pA)', fontsize=14)
         plt.show(fig)
```

Q2B: AdEx With Shorter Gap



A spike occurs after each impulse current. This happens because the second impulse current occurs while the adaptation current is negative (i.e, potentiating) as a result of the first impulse current. This brief negative adaptation current period occurs because *a* is positive.

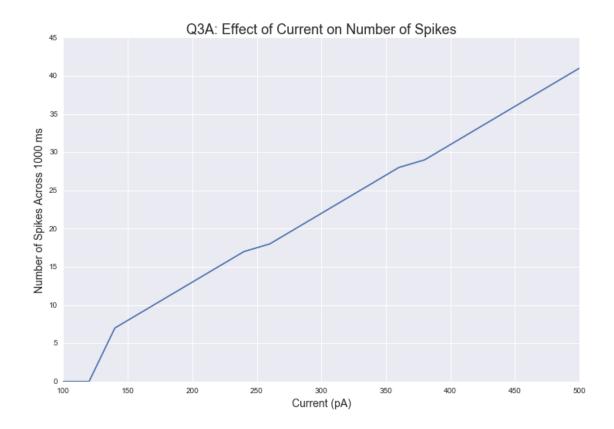
Conversely, in Q1B (where *a* equals zero), the second impulse current does not induce a spike, because the adaptation current is not negative at the time of the second impulse current. Thus, the adaptation current depresses in Q1B, but potentiates in Q2B.

3 Question 3

```
In [14]: # Set parameters
R = 100 * br.Mohm # membrane resistance
tau_m = 24 * br.ms # membrane time constant
u_rest = -65 * br.mV # resting potential
u_r = -60 * br.mV # reset potential
theta_reset = 40 * br.mV # reset threshold
vartheta_rh = -52 * br.mV # rheobase threshold
delta_T = 1 * br.mV # sharpness of ap upswing

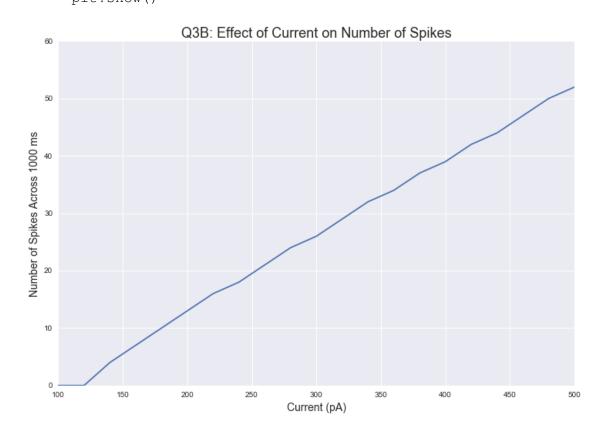
tau_w = 30 * br.ms # adaptation time constant
a = 0 * br.nS # subthreshold adaptation constant
b = 250 * br.pamp # adaptation jump after a spike
```

```
egs = '''
         du/dt = (-(u - u_rest) + delta_T * exp((u-vartheta_rh)/delta_T) -
                 R*w + R*I) / tau_m : volt
         dw/dt = (a*(u - u_rest) - w)/tau_w : amp
         I = input_current(t) : amp
         1.1.1
In [15]: # Track results
         names = {}
3.1 Part A
In [16]: # Run model
         current_range = range(100, 501, 20)
         n_spikes_a = []
         for current in current_range:
             input_current = np.ones((1000,)) * current
             input_current = br.TimedArray(input_current*br.pamp, dt=1*br.ms)
             br.start_scope()
             AdEx = br.NeuronGroup(1, eqs, threshold='u>theta_reset',
                                   reset='u=u_r; w+= b', method='euler')
             AdEx.u = u\_rest
             AdEx.w = 0 * br.pamp
             rec = br.SpikeMonitor(AdEx, ('u', 'w'), record=True)
             br.run(1*br.second)
             n_spikes_a.append(rec.count[0])
         names['A'] = {'n_spikes': n_spikes_a,
                       'descrip': r'AdEx: low $\tau_w$, high b, low $u_r$'}
In [17]: # Plot A
         fig, ax = plt.subplots()
         ax.set_xlabel('Current (pA)', fontsize=14)
         ax.set_ylabel('Number of Spikes Across 1000 ms', fontsize=14)
         ax.set_title('Q3A: Effect of Current on Number of Spikes', fontsize=18)
         ax.plot(current_range, n_spikes_a)
         plt.show()
```



3.2 Part B

```
In [18]: # Run model
         tau_w = 200 * br.ms
         b = 30 * br.pamp
         n_spikes_b = []
         for current in current_range:
             input_current = np.ones((1000,)) * current
             input_current = br.TimedArray(input_current*br.pamp, dt=1*br.ms)
             br.start_scope()
             AdEx = br.NeuronGroup(1, eqs, threshold='u>theta_reset',
                                   reset='u=u_r; w+= b', method='euler')
             AdEx.u = u\_rest
             AdEx.w = 0 * br.pamp
             rec = br.SpikeMonitor(AdEx, ('u', 'w'), record=True)
             br.run(1*br.second)
             n_spikes_b.append(rec.count[0])
         names['B'] = {'n_spikes': n_spikes_b,
                       'descrip': r'AdEx: high $\tau_w$, low b, low $u_r$'}
```

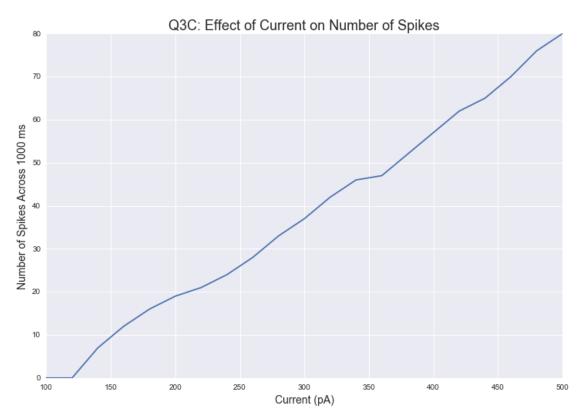


3.3 Part C

```
In [20]: # Run model
    tau_w = 200 * br.ms
    b = 30 * br.pamp
    u_r = -50 * br.mV

n_spikes_c = []
    for current in current_range:
        input_current = np.ones((1000,)) * current
        input_current = br.TimedArray(input_current*br.pamp, dt=1*br.ms)
        br.start_scope()
```

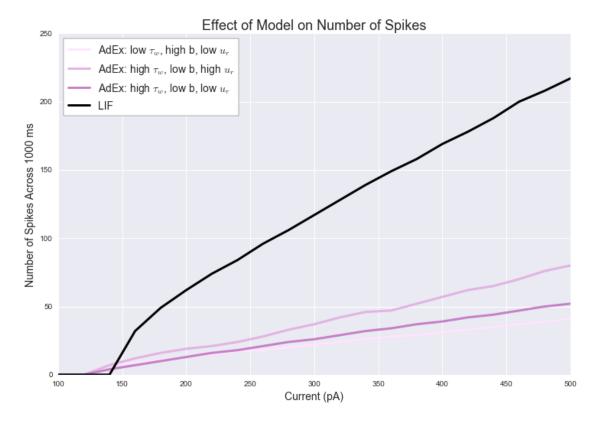
```
AdEx = br.NeuronGroup(1, eqs, threshold='u>theta_reset',
                                   reset='u=u_r; w+= b', method='euler')
             AdEx.u = u\_rest
             AdEx.w = 0 * br.pamp
             rec = br.SpikeMonitor(AdEx, ('u', 'w'), record=True)
             br.run(1*br.second)
             n_spikes_c.append(rec.count[0])
         names['C'] = {'n_spikes': n_spikes_c,
                       'descrip': r'AdEx: high $\tau_w$, low b, high $u_r$'}
In [21]: # Plot C
         fig, ax = plt.subplots()
         ax.set_xlabel('Current (pA)', fontsize=14)
         ax.set_ylabel('Number of Spikes Across 1000 ms', fontsize=14)
         ax.set_title('Q3C: Effect of Current on Number of Spikes', fontsize=18)
         ax.plot(current_range, n_spikes_c)
         plt.show()
```



3.4 Create LIF curve

```
In [22]: # Define parameters of the LIF model
         R = 100 * br.Mohm # membrane resistance
         tau = 10 * br.ms # membrane time constant
         thr = -50 * br.mV # spike threshold
         u rest = -65 * br.mV # resting potential
         u_r = -70 * br.mV # reset potential
         # Define equation for the LIF model
         eqs = '''
         du/dt = (-(u - u_rest) + R*I) / tau : volt
         I = input_current(t) : amp
         n_spikes_lif = []
         for current in current_range:
             br.start_scope()
             tmp = np.ones((1000,))
             tmp *= current
             input_current = br.TimedArray(tmp*br.pamp, dt=1*br.ms)
             LIF = br.NeuronGroup(N=1, model=eqs, method='linear',
                                  threshold='u>thr', reset='u=u_r')
             LIF.u = u_rest
             rec = br.SpikeMonitor(LIF, 'u', record=True)
             br.run(1*br.second)
             n_spikes_lif.append(rec.count[0])
         names['LIF'] = {'n_spikes': n_spikes_lif,
                         'descrip': 'LIF'}
In [23]: # Plot number of spikes against current value.
         fig, ax = plt.subplots()
         for i, model in enumerate(names.keys()):
             if model == 'LIF':
                 c = 'black'
             else:
                 c = palette[i]
             ax.plot(current_range, names[model]['n_spikes'],
                     1w=3., c=c.
                     label=names[model]['descrip'])
         legend = ax.legend(frameon=True, loc='upper left', fontsize=14)
         frame = legend.get_frame()
         frame.set_facecolor('white')
         frame.set_edgecolor('black')
         ax.set_xlabel('Current (pA)', fontsize=14)
         ax.set_ylabel('Number of Spikes Across 1000 ms', fontsize=14)
```

ax.set_title('Effect of Model on Number of Spikes', fontsize=18)
plt.show(fig)



All else held equal, an increase in u_r (e.g., Q3B vs. Q3C) increases the rate at which increases in current increase the number of spikes. As b decreases and tau_w increases (e.g., Q3A vs. Q3C), the rate at which increases in current increase the number of spikes increases. Based on these simulations, it is impossible to know whether b or tau_w drives this difference.

All of the AdEx models' spike rates increase far more slowly than the LIF neuron. As such, the input current value has a much greater effect on the LIF model's spike rate than on AdEx models. Also, the LIF model has a higher overall spike rate than the AdEx model. The underlying differences in the model dynamics do affect the I - f functions dramatically.