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Dipartimento di Informatica Università di Pisa - Italy

Neural Modeling and Computational Neuroscience

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Neuroscience modeling

- Introduction to basic aspects of brain computation
- Introduction to neurophysiology
- Neural modeling:
 - Elements of neuronal dynamics
 - Elementary neuron models
 - Neuronal Coding
 - Biologically detailed models:

the Hodgkin-Huxley Model

- Spiking neuron models, spiking neural networks
- Izhikevich Model
- Introduction to Reservoir Computing and Liquid State Machines
- Introduction to glia and astrocyte cells, the role of astrocytes in a computational brain, modeling neuron-astrocyte interaction, neuronastrocyte networks,
- The role of computational neuroscience in neuro-biology and robotics applications.

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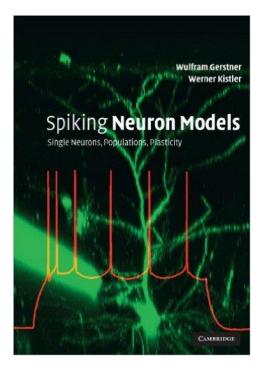
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References

W. Gerstner and W.M. Kistler, Spiking Neuron Models: Single Neurons, Population, Plasticity. Cambridge Univ. Press, 2002

on-line at: http://lcn.epfl.ch/~gerstner/SPNM/SPNM.html



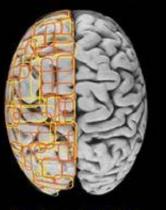
Chap. 2 – Sect. 2.1, 2.2 Chap. 4 – Sect. 4.1

References

P. Dayan and L.F. Abbott, Theoretical neuroscience. MIT Press, 2001.

THEORETICAL NEUROSCIENCE

Computational and Mathematical Modeling of Neural Systems

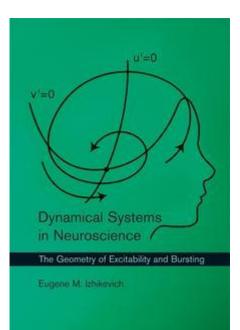


Peter Dayan and L. F. Abbott

Chap. 5 – Sect. 5.1, 5.2, 5.4, 5.5, 5.6

References

E.M. Izhikevich, Dynamical Systems in Neuroscience: The Geometry of Excitability and Bursting. The MIT press, 2007



Chap. 1 Chap. 2 Chap. 8 – Sect. 8.1, 8.2

Conductance-based Neuron Models

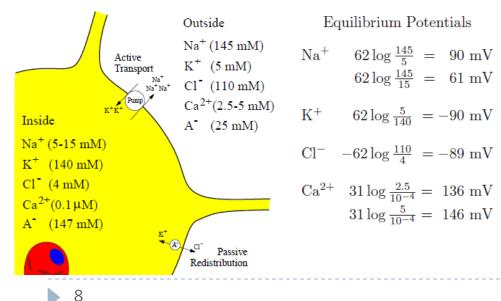
Reversal Potential (Repetita)

The reversal potential of an ion is its Nernst potential

$$E_{[ion]} = \frac{kT}{q_{[ion]}} \ln \frac{n_{out}}{n_{in}}$$

• If $\Delta u < E_{[ion]} \Rightarrow$ ions flow into the cell

• If $\Delta u > E_{\text{[ion]}} \Rightarrow$ ions flow out of the cell

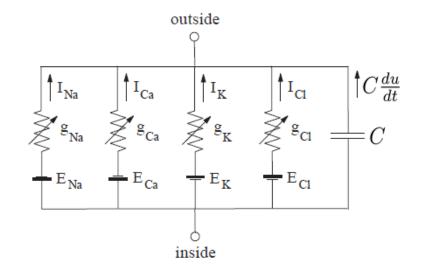


 $E_K < E_{Cl} < u_{rest} < E_{Na} < E_{Ca}$

- Ion channels: try to equilibrate the concentration of ions, i.e. try to meet the reversal potential
- Ion pumps: active pumps that balance the flow of ions

Equivalent Circuit (Repetita)

Electrical properties of neurons' membranes depicted in terms of the electrical circuit

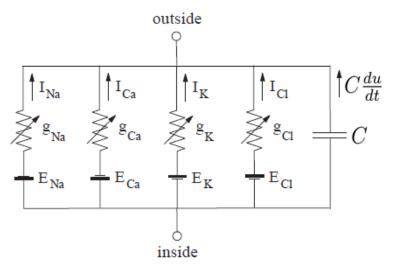


- $\uparrow C \frac{du}{dt}$ Membrane: capacitor = C Ions' channels: resistors + battery (reversed potentials) (reversal potentials)

$$I_{Na} = g_{Na}(u - E_{Na})$$
 $I_{Ca} = g_{Ca}(u - E_{Ca})$

$$I_K = g_K(u - E_K)$$
 $I_{Cl} = g_{Cl}(u - E_{Cl})$

Equivalent Circuit

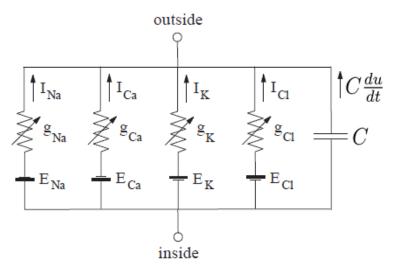


- Membrane: capacitor
- lons' channels: resistors + battery (reversal potentials)

$$E_K < E_{Cl} < u_{rest} < E_{Na} < E_{Ca}$$

$$I_{Na} = g_{Na}(u - E_{Na}) \qquad I_{Ca} = g_{Ca}(u - E_{Ca}) \qquad \text{inward current}$$
$$I_K = g_K(u - E_K) \qquad I_{Cl} = g_{Cl}(u - E_{Cl}) \qquad \text{outward current}$$

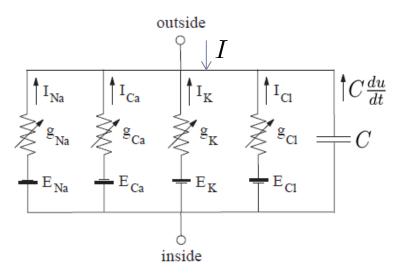
Equivalent Circuit



$$E_K < E_{Cl} < u_{rest} < E_{Na} < E_{Ca}$$

$$I_{Na} = g_{Na}(u - E_{Na}) \qquad I_{Ca} = g_{Ca}(u - E_{Ca}) \qquad \text{inward current}$$
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Equivalent Circuit

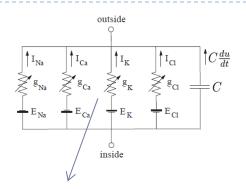


- Membrane: capacitor
- lons' channels: resistors + battery (reversal potentials)
- Applied current I

Using Kirchhoff's Current Law (KCL): $C = \frac{q}{u} \Rightarrow C\frac{du}{dt} = I_C$

$$C\frac{du}{dt} = I - I_{Na} - I_{Ca} - I_K - I_{Cl}$$
$$C\frac{du}{dt} = I - g_{Na}(u - E_{Na}) - g_{Ca}(u - E_{Ca}) - g_K(u - E_K) - g_{Cl}(u - E_{Cl})$$

Conductances



non-Ohmic currents (conductances are not constant)

Ion channels:



 Electrical conductance of individual channels is controlled by gates (gating particles)

Intracellular

Extracellular

membrane

inactivation

gate

- Gates can change the state of the channel: open/closed
- Gates can be sensitive to the <u>membrane potential (voltage-dependent</u> conductances), intracellular agents, neurotransmitters,

Na⁺

Open

(activated)

selectivity

Closed

(inactivated)

filte

voltage

sensor

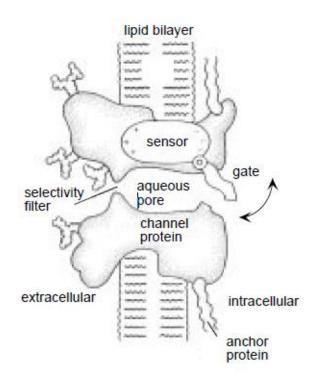
activation

gate

Closed

(not activated)

Persistent Conductances



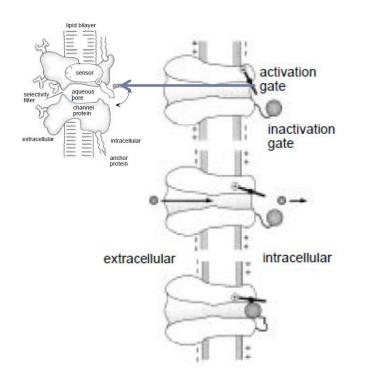
Voltage dependency:

depolarization of the membrane leads to increasing n

- a voltage sensor is connected to a swinging (activation) gate that can open or close the pore
- gate opening: activation of the conductance
- gate closing: de-activation of the conductance
- results in a persistent (or noninactivating) conductance
- Probability of the channel to be opened: $p = n^k$

gating variable: the probability that one of the k sub-units of the gate is opened

Transient Conductances



Voltage dependency:

Depolarization: increasing m, decreasing hHyper-polarization: decreasing m, increasing h

- Two gates regulates the channel:
 1 activation gate & 1 inactivation gate
- The activation gate is opened with probability m^k
- The inactivation gate (the ball) does not block the channel with probability h
- The channel is opened with probability m^kh
- The channel opens transiently while the membrane is depolarized

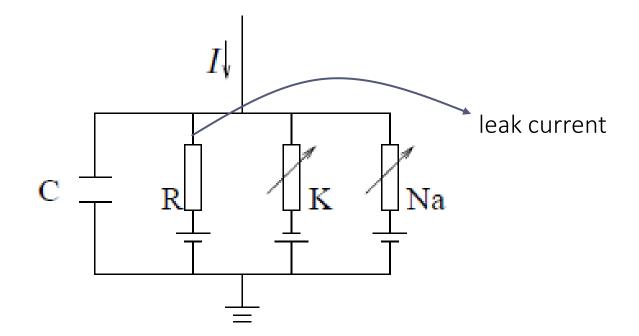
The Hodgkin-Huxley Model

The Hodgkin-Huxley Model

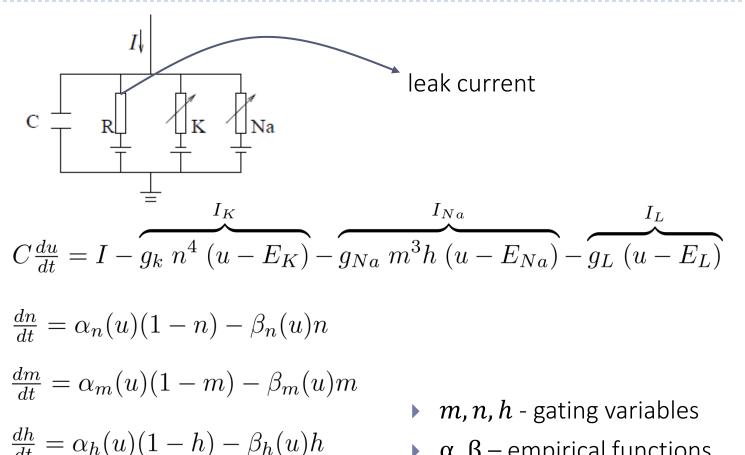
- One of the most important models in Computational Neuroscience
- Based on studies by Hodgkin and Huxley (in the 50s) on the squid axon
- > The squid axon has 3 major currents:
 - ▶ Voltage-gated persistent K⁺ current with 4 activation gates
 - Voltage-gated transient Na⁺ current with 3 activation gates and 1 inactivation gate
 - Ohmic leak current (all the other ions)

$$I_{[ion]} = g_{[ion]} p \left(u - E_{[ion]} \right)$$

Hodgkin-Huxley Model



Hodgkin-Huxley Model



α, β – empirical functions
 adjusted by Hodgkin and Huxley

Hodgkin-Huxley Model

$$C = I - g_k n^4 (u - E_K) - g_{Na} m^3 h (u - E_{Na}) - g_L (u - E_L)$$

The equations for the gating variables can be rewritten as

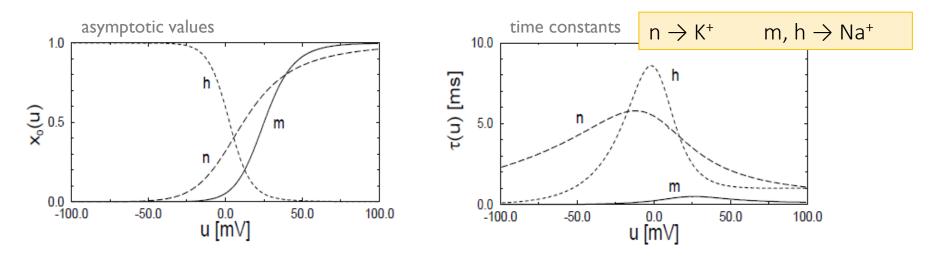
$$\frac{dn}{dt} = \frac{n_0(u) - u}{\tau_n} \qquad \frac{dm}{dt} = \frac{m_0(u) - u}{\tau_m} \qquad \frac{dh}{dt} = \frac{h_0(u) - u}{\tau_h}$$

where:

$$n_0(u) = \frac{\alpha_n(u)}{\alpha_n(u) + \beta_n(u)}, \tau_n(u) = \frac{1}{\alpha_n(u) + \beta_n(u)}$$
$$m_0(u) = \frac{\alpha_m(u)}{\alpha_m(u) + \beta_m(u)}, \tau_m(u) = \frac{1}{\alpha_m(u) + \beta_m(u)}$$
$$h_0(u) = \frac{\alpha_h(u)}{\alpha_h(u) + \beta_h(u)}, \quad \tau_h(u) = \frac{1}{\alpha_h(u) + \beta_h(u)}$$

- $n_0(t), m_0(t), h_0(t)$ asymptotic values
- $\tau_n(t), \tau_m(t), \tau_h(t)$ time constants

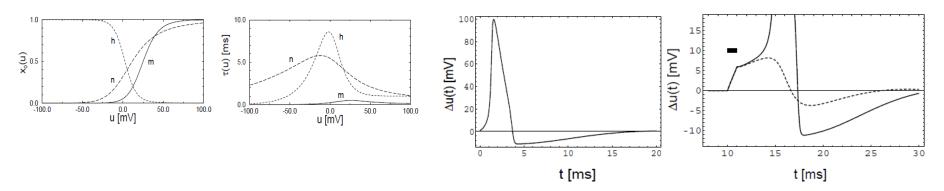
Hodgkin-Huxley Model – Dynamics



Sodium (Na⁺) – inward current:

- Activation increases for increasing membrane potential
- Inactivation increases for increasing membrane potential
- BUT: activation is faster than inactivation (transient current)
- ▶ Potassium (K⁺) outward current:
 - Activation increases for increasing membrane potential
 - BUT: activation is relatively slow (slower than activation of Na⁺)

Hodgkin-Huxley Model – Spike Generation



- An external input (e.g. an EPSP) leads to a depolarization (u increases)
- Conductance of Na⁺ increases rapidly, Na⁺ ions flow in the cell and u increases even further
- If the feedback is strong enough the action potential is initiated
- At high values of depolarization, the Na⁺ current is stopped by the inactivation gate $(h \rightarrow 0)$, conductance of K⁺ increases and K⁺ ions flow outside the cell
- The membrane is re-polarized, with a negative overshoot (refractoriness)
- Threshold behavior: if the stimulating input is below a certain amplitude the action potential is not initiated and the membrane is re-polarized

The Hodgkin-Huxley Model - Summary

$$C\frac{du}{dt} = I - \overbrace{g_k \ n^4 \ (u - E_K)}^{I_K} - \overbrace{g_{Na} \ m^3 h \ (u - E_{Na})}^{I_{Na}} - \overbrace{g_L \ (u - E_L)}^{I_L}$$

$$\frac{dn}{dt} = \alpha_n(u)(1 - n) - \beta_n(u)n$$

$$\frac{dm}{dt} = \alpha_m(u)(1 - m) - \beta_m(u)m$$

$$\frac{dh}{dt} = \alpha_h(u)(1 - h) - \beta_h(u)h$$

- Conductance-based neuron model
- Processes that regulate the voltage-dependent K⁺ and Na⁺ conductances well described
- Biophysical mechanisms responsible for action potentials explicitly included in the mathematical model
- Accurate biological realism, BUT slow and difficult to analyze.

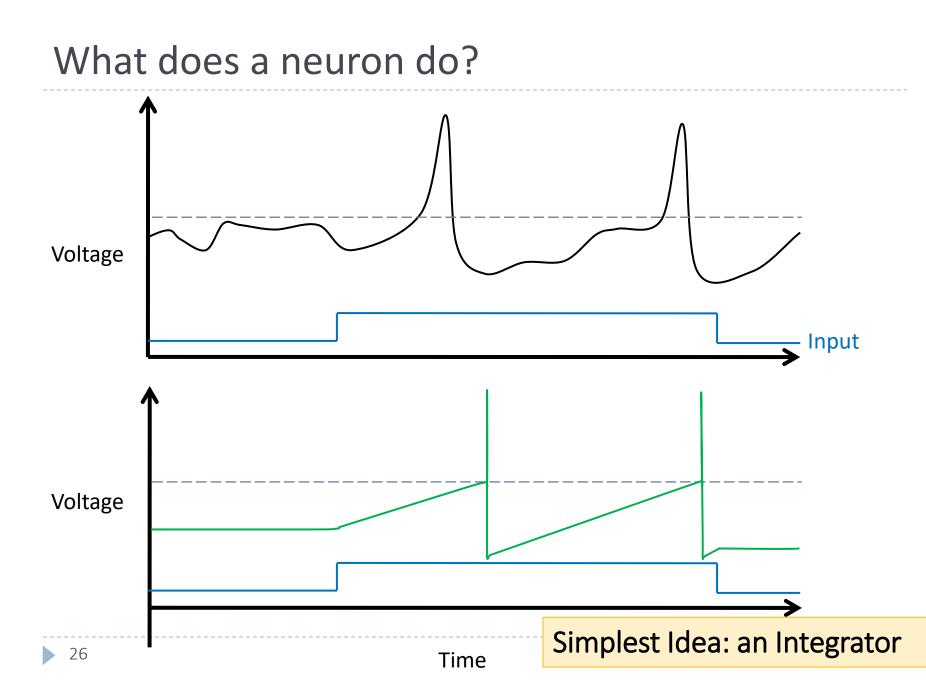
Formal Spiking Neuron Models

Phenomenological Spiking Neuron

- Neuron models can be simplified and simulations can be accelerated if the biophysical mechanisms of spikegeneration are not included explicitly in the model
- Formal threshold models of neuronal firing:
 - Spikes are stereotyped events that occur when the membrane potential crosses the threshold from below

$$t^{(f)}: \quad u(t^{(f)}) = \vartheta \quad \text{and} \quad \left. \frac{\mathrm{d}u(t)}{\mathrm{d}t} \right|_{t=t^{(f)}} > 0$$

- Spikes are fully characterized by their firing time
- Model only the sub-threshold dynamics



Integrate-and-Fire Model

- The most simple case: all membrane conductances are ignored
- The corresponding equivalent (simplified) circuit only contains a capacitor

From the definition of the capacity: C = \$\frac{q}{u}\$ \$\Rightarrow\$ C\frac{du}{dt}\$ = I_C
 KCL: C\$\frac{du}{dt}\$ = I(t)\$ \$\frac{du}{dt}\$ = \$\frac{I(t)}{C}\$

- Spikes are formal events characterized by the firing time $t^{(f)}: \ u(t^{(f)}) = \vartheta$
- After the spike the potential is reset to u_r $\lim_{t \to t^{(f)+}} u(t) = u_r$
- Absolute refractory period: after the spike, the integration is suspended for $\,\Delta^{abs}$

Integrate-and-Fire Model

Equations

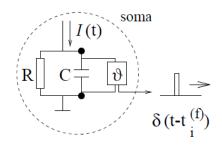
$$\begin{cases} \frac{du}{dt} = \frac{I(t)}{C} \\ t^{(f)}: \ u(t^{(f)}) = \vartheta \\ \lim_{t \to t^{(f)+}} u(t) = u_r \qquad u_r \text{ is often set to } 0 \end{cases}$$

• Suppose a constant input current I_0 is applied (e.g. an EPSP), and the last spike occurred at time $t^{(1)}$:

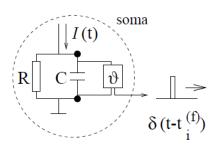
the time course of the membrane potential can be obtained by integration in the time interval $t^{(1)}$; t

$$u(t) = \int_{t^{(1)}}^{t} \frac{I_0}{C} ds = \frac{I_0}{C} (t - t^{(1)})$$

- The entire membrane conductance is modeled as a single leakage term
- Assumption: the conductances are all constant (true for small fluctuations around the resting membrane potential)
- Corresponding equivalent circuit: a capacitor in parallel with a resistor



- The entire membrane conductance is modeled as a single leakage term
- Assumption: the conductances are all constant (true for small fluctuations around the resting membrane potential)
- Corresponding equivalent circuit: a capacitor in parallel with a resistor



Ohm's Law + Kirchhoff's Voltage Law:

$$u(t) = I_R R \Rightarrow I_R = \frac{u(t)}{R}$$
KCL:

$$I_C + I_R = I(t) \Rightarrow C\frac{du}{dt} = -\frac{u(t)}{R} = I(t)$$

$$\tau_m \frac{du}{dt} = -u(t) + RI(t)$$

$$\tau_m = RC \qquad \text{membrane}$$
time constant

- Time course of the membrane potential?
- Suppose a constant input current I_0 is applied and the last spike occurred at time $t^{(1)}$
- u(t)???

$$\tau_m \frac{du}{dt} = -u(t) + RI(t)$$

First-Order linear differential equation (with initial condition $u(t^{(1)}) = u_r = 0$)

First-Order Linear Differential Equation

$$\begin{cases} y'(t) + a(t)y(t) = f(t) \\ y(t_0) = y_0 \end{cases}$$

solution: $y(t) = y_0 \ e^{-A(t)} + e^{-A(t)} \int_{t_0}^t f(s) e^{A(s)} ds$

where:

$$A(t) = \int_{t_0}^t a(s) ds$$

Also useful to remember:

$$e^{f(t)}f'(t)dt = e^{f(t)} + c$$

Euler Method

Numerical (approximate) method for solving ODEs

 $\begin{cases} y'(t) = f(t, y(t)) \\ y(t_0) = y_0 \end{cases}$

By discretizing the temporal variable t: $t_n = t_0 + nh$

h = dimension of the interval

The evolution of the system can be approximated by

$$y(t_{n+1}) = y(t_n) + hf(t_n, y(t_n))$$

- Time course of the membrane potential?
- Suppose a constant input current I_0 is applied and the last spike occurred at time $t^{(1)}$

• u(t)???

$$\tau_m \frac{du}{dt} = -u(t) + RI(t)$$

First-Order linear differential equation (with initial condition $u(t^{(1)}) = u_r = 0$)

$$u(t) = RI_0 \left(1 - e^{-\frac{t-t^{(1)}}{\tau_m}}\right)$$

(The membrane potential asymptotically approaches RI_0)

When will next spike occur?

$$u(t^{(2)}) = \vartheta = RI_0 \left(1 - e^{-\frac{T}{\tau_m}}\right) \qquad T = t^{(2)} - t^{(1)}$$

$$T = \Delta^{abs} + \tau_m \ln\left(\frac{RI_0}{RI_0 - \vartheta}\right)$$
Firing rate (with refractory period)
$$\nu = \left[\Delta^{abs} + \tau_m \ln\left(\frac{RI_0}{RI_0 - \vartheta}\right)\right]^{-1}$$

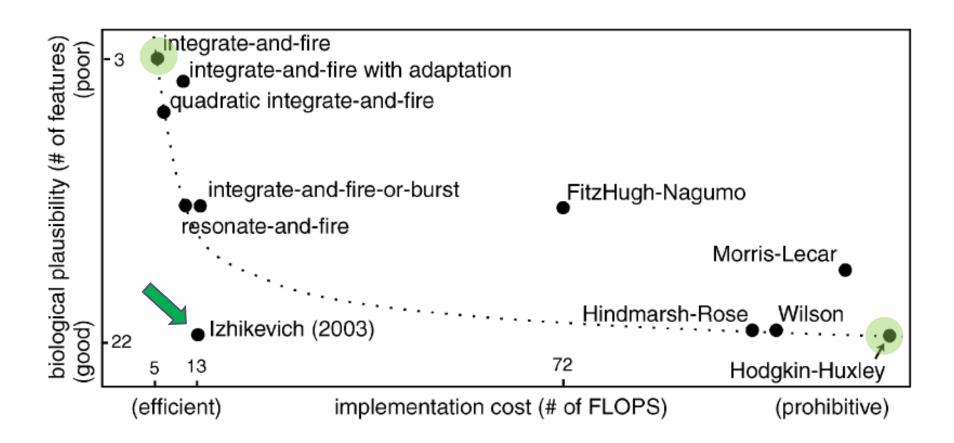
$$\underbrace{\Phi}_{\exists} 1.0 \left[\underbrace{\Phi}_{abs} + \frac{1}{20} \underbrace{\Phi}_{abs} + \frac{1}{20}$$

Izhikevich Model

Simple Spiking Models

- Modeling the dynamics of excitable neurons
 - Fast activation of Na⁺ channels
 - Slow inactivation of Na⁺/activation of K⁺
- Dynamical system with 2 variables
 - One variable for the fast voltage increase
 - One recovery variable for slow voltage decrease
- In many cases the sub-threshold dynamics leading to the action potential are more important than the shape of the action potential itself

Neuron Models – Biological Plausibility vs Cost



Izhikevich Model

Two dimensional system of ordinary differential equations

$$\frac{du}{dt} = 0.04 \ u(t)^2 + 5u(t) + 140 - r(t) + I$$
$$\frac{dr}{dt} = a(bu(t) - r(t))$$

 $|f u(t) \ge 30 \text{ mV}$

$$\begin{bmatrix} u = c \\ r = r + d \end{bmatrix}$$

- u is the membrane potential,
- r is a recovery variable
 (Na+ inactivation/K+ activation)
 provides negative feedback to u
- a, b, c, d are the parameters of the model
- ▶ *I* is the applied current

Izhikevich Model

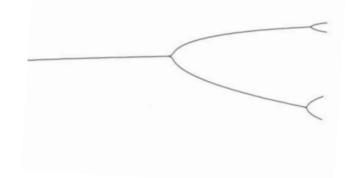
Two dimensional system of ordinary differential equations

```
v'= 0.04v<sup>2</sup>+5v +140 - u + l
u'= a(bv - u)
if v= 30 mV,
then v ← c, u ← u + d
```

Often in literature:

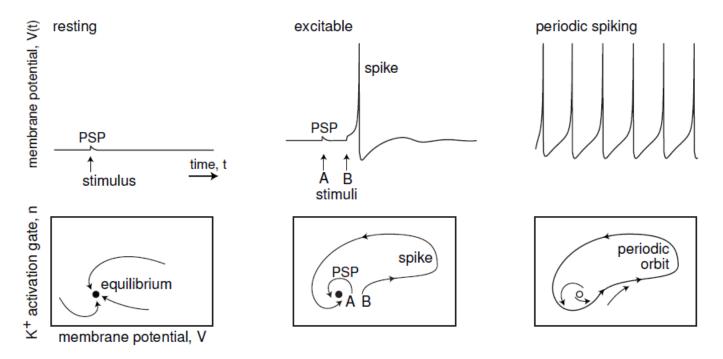
- v is the membrane potential
- u is the recovery variable

- The behavior of a neuron does not depends only on its electrophysiological properties
- Two neurons with the same electrophysiological properties can respond differently to the same input
- Neurons can be thought as dynamical systems
- Dynamical properties of the neurons have a major role Especially bifurcation dynamics



A bifurcation occurs when a small change to the parameter values of a system results in a sudden qualitative change in its behavior

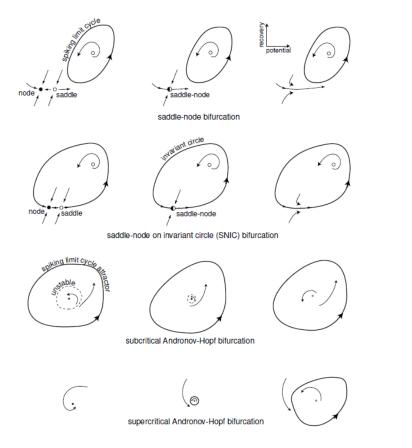
 Neurons are excitable because they are near a transition (bifurcation) between resting and sustained spiking activity



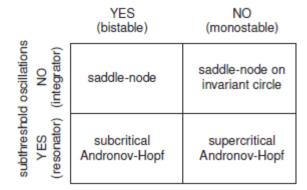
The system is excitable because its equilibrium is near a bifurcation

D

Four generic bifurcations

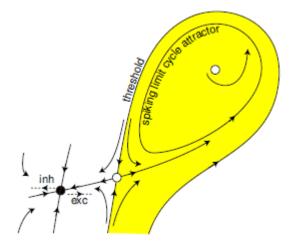


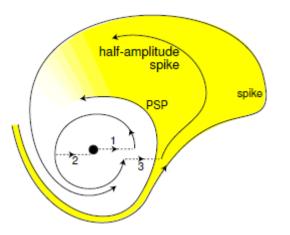
co-existance of resting and spiking states



- Monostable: the neuron does not exhibits the presence of resting and tonic spiking
- Resonator: there exist small amplitude oscillations of membrane potential

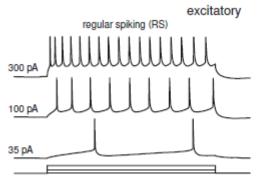
Integrators vs Resonators

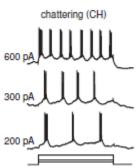


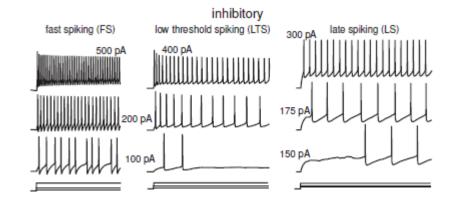


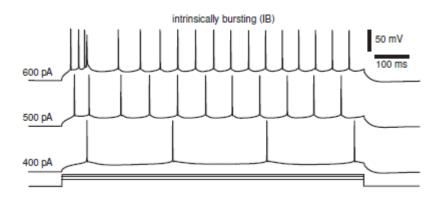
Firing Patterns

> The most fundamental classes of firing patterns are just 6



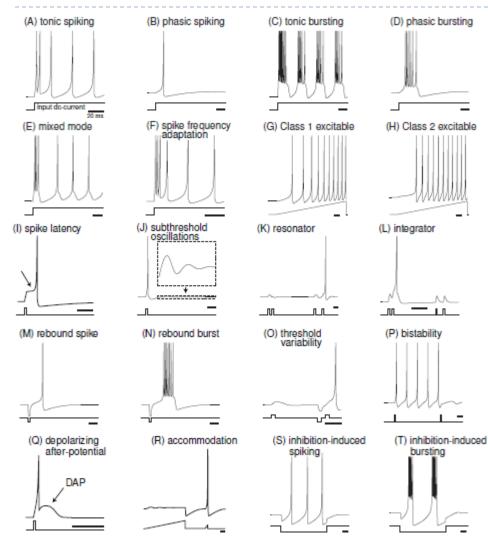








Neuro-Computational Features



- 20 Most prominent features of biological spiking neurons
- The Izhikevich model can simulate all of them
- Izhikevich's book Chapter 8

Papers:

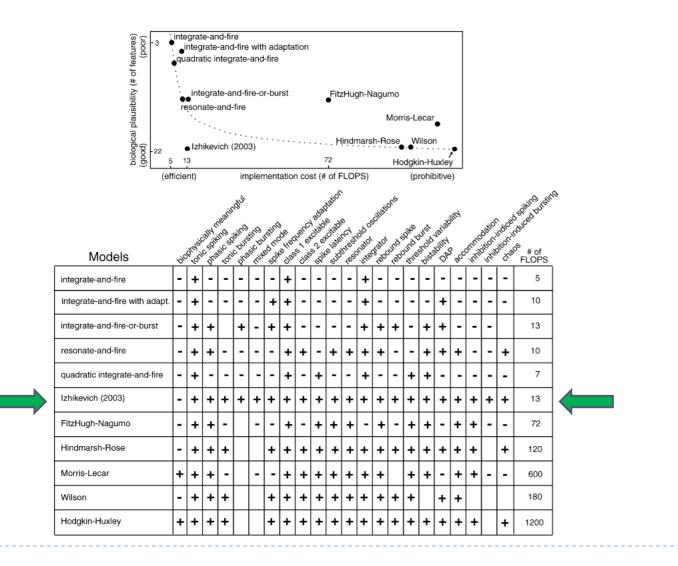
E.M. Izhikevich, "Which model to use for cortical spiking neurons?." *IEEE transactions on neural networks* 15.5 (2004): 1063-1070.

E.M. Izhikevich, "Simple model of spiking neurons." IEEE Transactions on neural networks 14.6 (2003): 1569-1572.

web:

http://izhikevich.org/publications/whichmod.htm

Which Model to Use for Cortical Spiking Neurons?



What happens if we connect spiking neurons to form a network?

TD BE CONTINUED

The Other Half of the Brain

Mounting evidence suggests that glial cells, overlooked for half a century, may be nearly as critical to thinking and learning as neurons are ...see you on Monday!!

