Introduction to Computational Neuroscience

Biol698 Math635 Biol498 Math430



Bibliography:

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* "Biophysics of Computation - Information processing in single neurons", by Christof Koch. Oxford University Press, 1999. ISBN 0-19-510491-9

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Neuroscience is the study of the brain

Brain function underlies all behavior

•Motor activity (walking, smiling, etc.)

•Higher order cognitive behavior (thinking, learning, memory, decision making)

The brain consists of neurons and glial cells.

Neurons are the basic signaling units of the nervous system.

Glial cells are important in maintaining the health of neurons and how they behave and interact

Neurons communicate with each other at synapses

Human Brain:

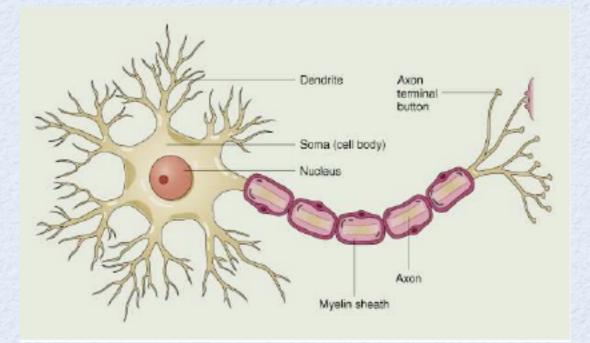
- 10¹¹ neurons
- 10¹⁵ synapses

Neurons typically receive inputs from a large number of other cells A typical 1 mm3 cortical tissue contains 10⁵ neurons Neuron size varies from the micrometer to the meter scales

The estimated number of different morphological classes of neurons in the vertebrate brain is 10,000

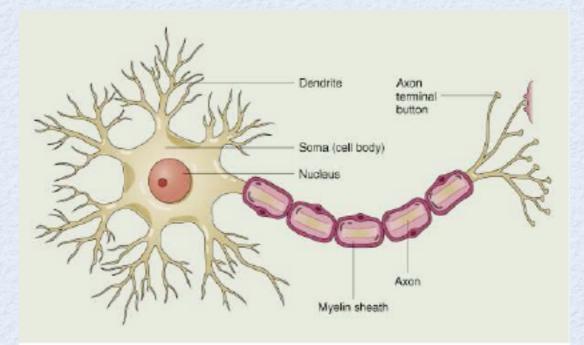
Neurons:

- Soma or cell body
- Dendrites
- Axon



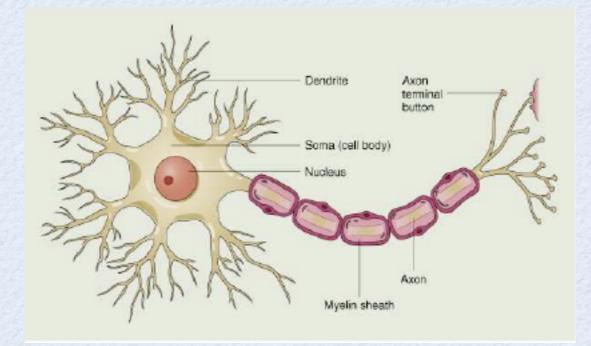
Soma:

- Site of the nucleus and all the normal cellular machinery
- Functionally the soma plays the role of integrating all of the inputs of the cell to produce some output



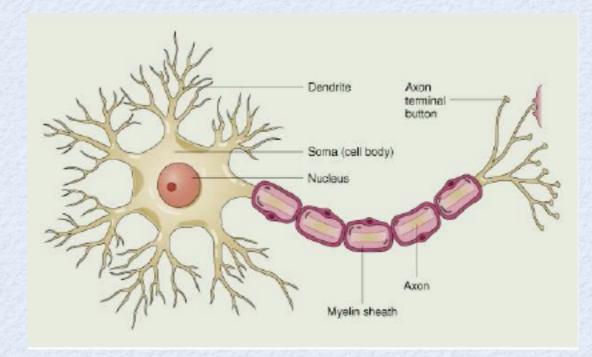
Dendrites:

- "Input line" to the neuron
- Dendrites that lie at the top of the neuron are called apical
- Dendrites that lie at the base of the neuron are called basal
- In many cases, the majority of the surface area of the cell is taken up by the dendrites



Axon:

- Major conducting unit of the neuron ("output line")
- The diameter of an axon may range from .2 to 20 µm and may extend for up to one meter
- It can convey information great distances by propagating a transient electrical signal called the action potential
- Large axons are surrounded by a fatty insulating sheath called myelin
- The sheath is interrupted at regular intervals by nodes of Ranvier.



Neurons, like all other cells, maintain a potential difference of about 65 mV across their external membrane

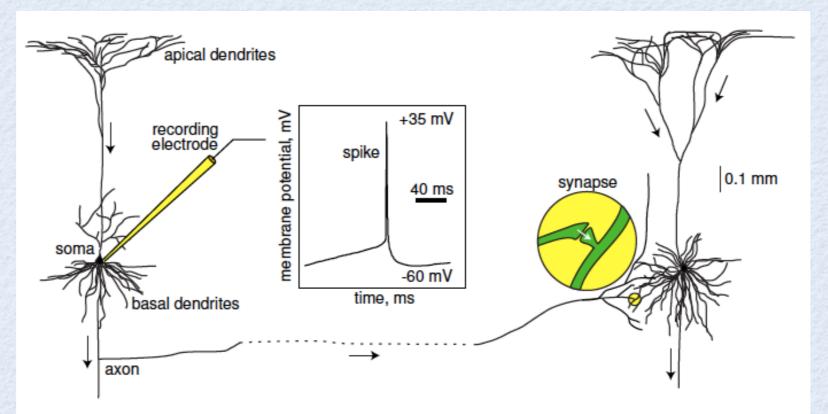


Figure 1.1: Two interconnected cortical pyramidal neurons (hand drawing) and *in vitro* recorded spike.

What distinguishes neurons and other excitable cells from most cells in the body is that this resting membrane potential can be altered and can, therefore, serve as a signaling mechanism

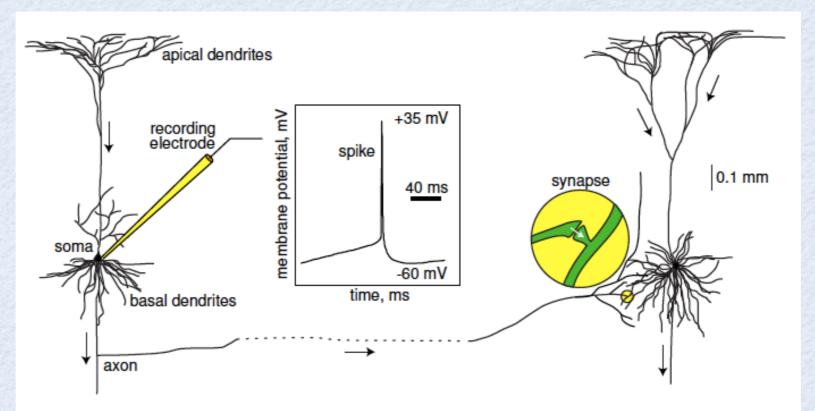


Figure 1.1: Two interconnected cortical pyramidal neurons (hand drawing) and *in vitro* recorded spike.

Action potentials:

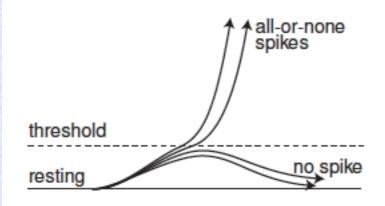
- Fast change in the neuron's membrane potential
- It is initiated at a specialized region of the soma called the axon hillock
- It is initiated when the membrane potential reaches some threshold value
- It propagates along the axon
- May last as short as 1 msec
- May travel at rates that vary between about 1 and 100 meters per second
- The shape and speed of each action potential does not depend on the stimulus

Refractory period:

 After each action potential, there is a period during which a second impulse cannot be initiated

Firing frequency:

- Stronger stimuli often produce higher frequencies of impulse firing
- Since the shape and speed of each action potential does not depend on the stimulus, information about the stimulus is often carried in the frequency of firing
- Limited by the refractory period
- Some neurons are capable of continuously generating action potentials, even without inputs
- The temporal pattern of firing may be quite complicated.





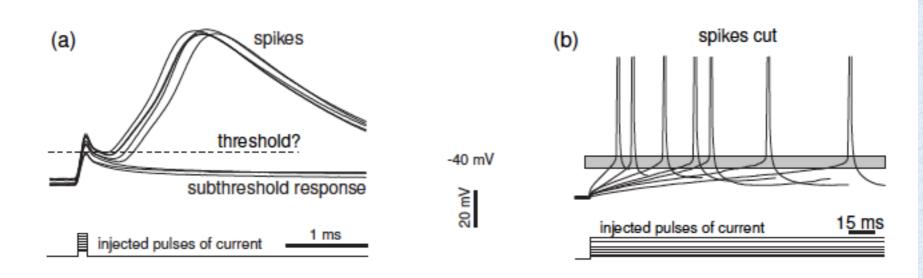


Figure 1.4: Where is the firing threshold? Shown are *in vitro* recordings of two layer 5 pyramidal neurons of rat. Notice the difference of voltage and time scales.

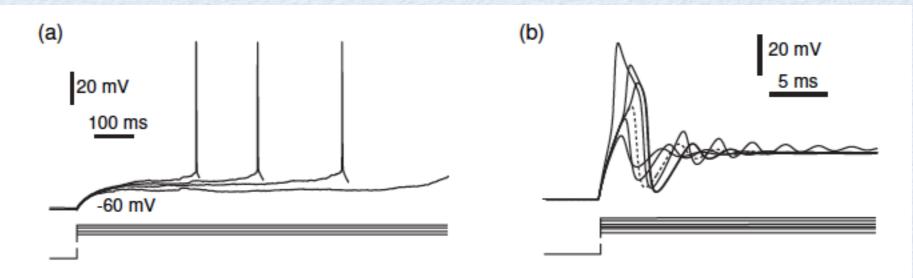


Figure 1.5: Where is the rheobase, i.e., the minimal current that fires the cell? (a) in vitro recordings of pyramidal neuron of layer 2/3 of rat's visual cortex show increasing latencies as the amplitude of the injected current decreases. (b) Simulation of the $I_{\text{Na,p}}+I_{\text{K}}$ -model shows spikes of graded amplitude.

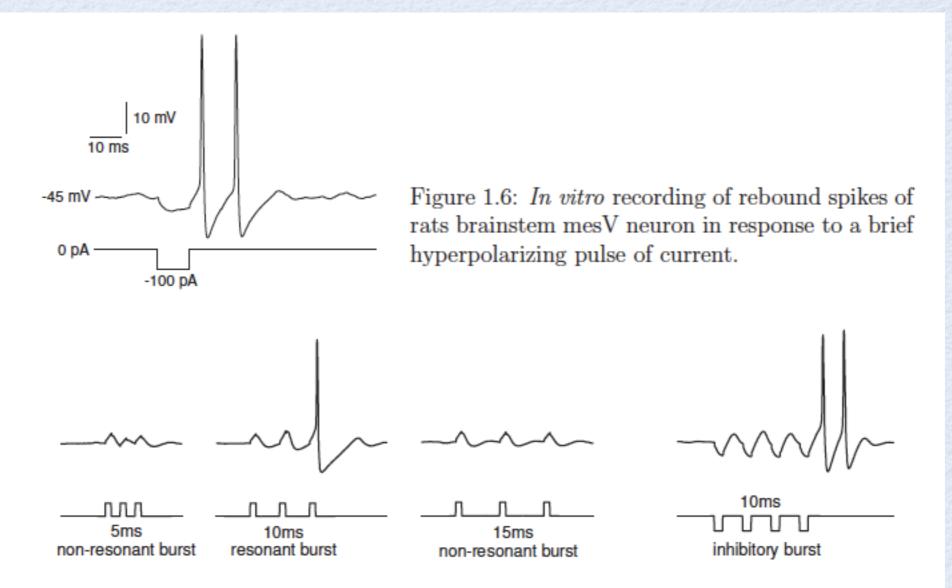


Figure 1.7: Resonant response of the mesencephalic V neuron of rat brainstem to pulses of injected current having 10 ms period (*in vitro*).

Dynamical systems tools:

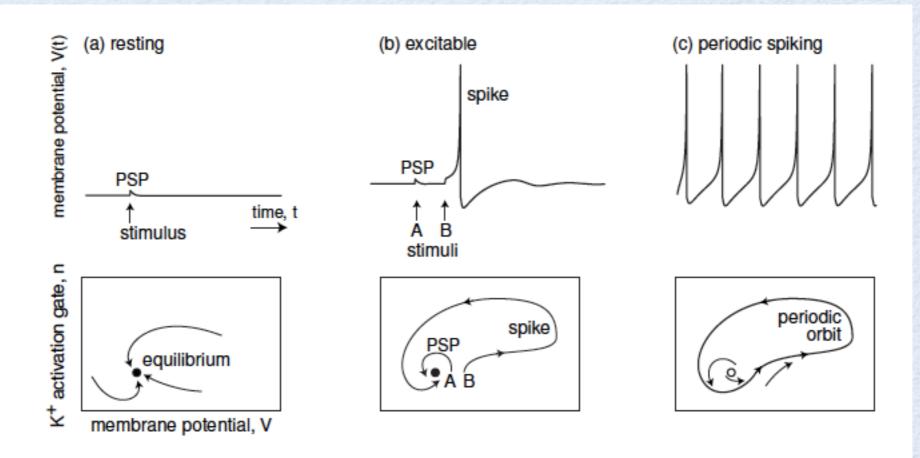


Figure 1.9: Resting, excitable, and periodic spiking activity correspond to a stable equilibrium (a and b) or limit cycle (c), respectively.

Bursting

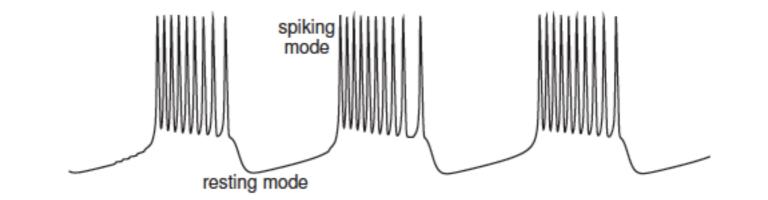


Figure 1.10: Rhythmic transitions between resting and spiking modes result in bursting behavior.

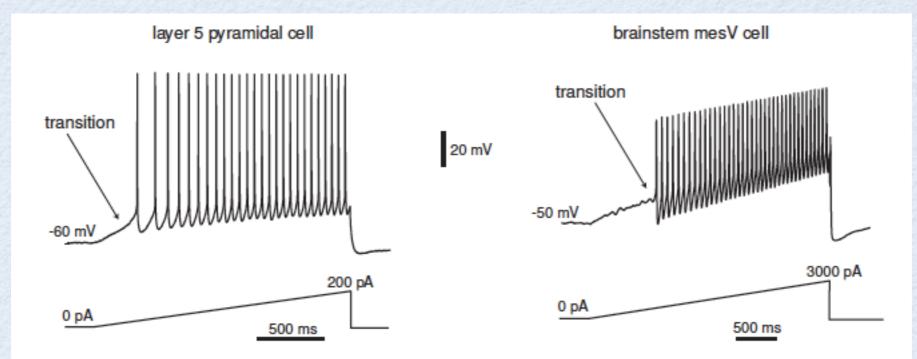
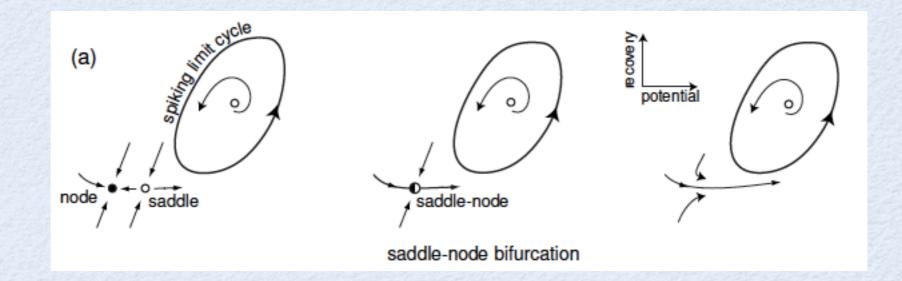
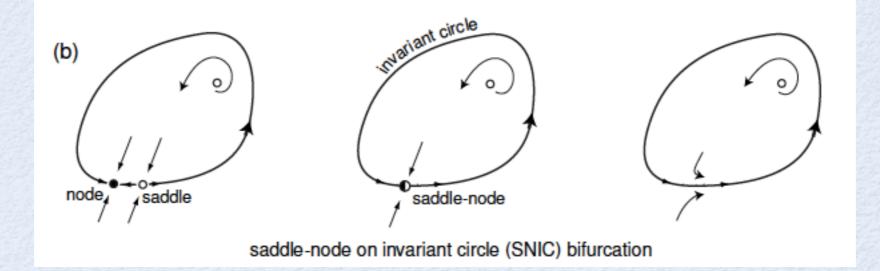
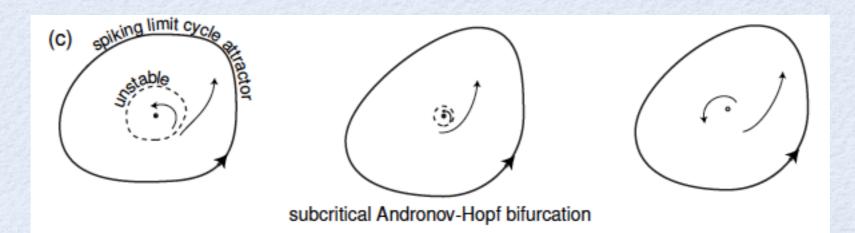


Figure 1.11: As the magnitude of the injected current slowly increases, the neurons bifurcate from resting (equilibrium) to tonic spiking (limit cycle) modes.







Bifurcations:

(d) (c) supercritical Andronov-Hopf bifurcation

Bifurcations:

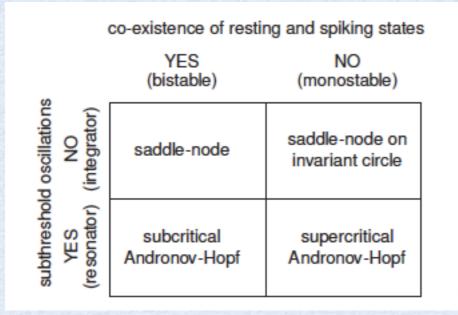


Figure 1.13: Classification of neurons into monostable/bistable integrators/resonators according to the bifurcation of the resting state in Fig. 1.12.

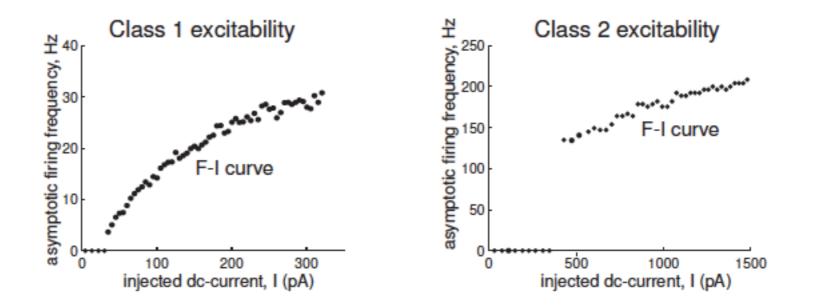


Figure 1.14: Frequency-current (F-I) curves of cortical pyramidal neuron and brainstem mesV neuron from Fig. 7.3. These are the same neurons used in the ramp experiment in Fig. 1.11.

Hodgkin-Huxley classification of neural excitability:

- Class 1: Action potentials can be generated with arbitrarily low frequency, depending on the strength of the applied current
- Class 2: Action potentials are generated in a certain frequency band that is relatively insensitive to changes in the strength of the applied current.
- Class 3: A single action potential is generated in response to a pulse of current. Repetitive (tonic) spiking can be generated only for extremely strong injected currents or not at all.

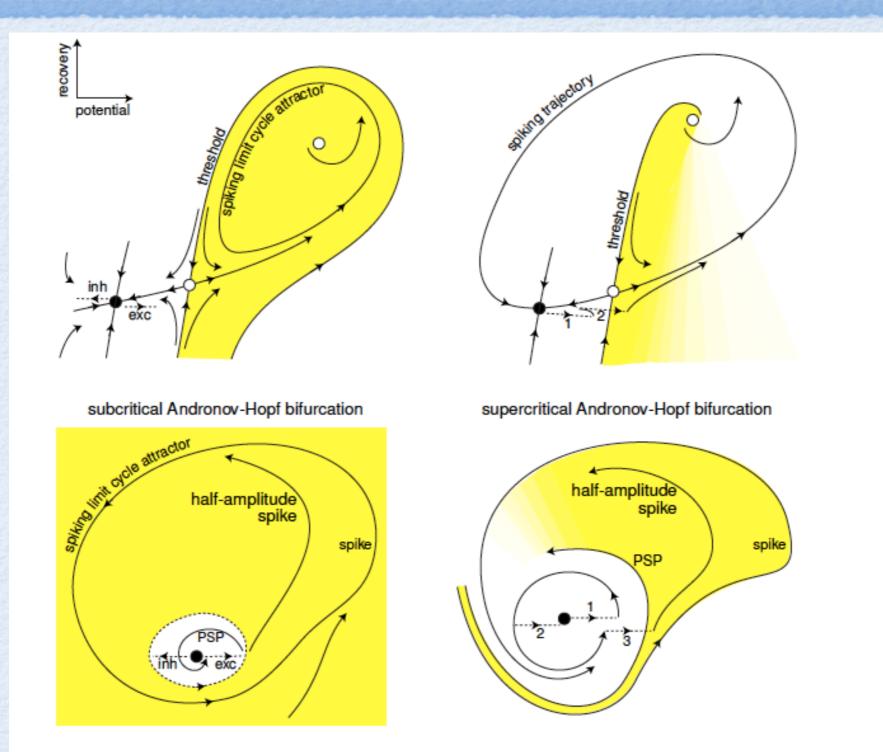


Figure 1.15: The geometry of phase portraits of excitable systems near 4 bifurcations can explain many neuro-computational properties (see Sect. 1.2.4 for detail).