

Welcome to CSE/NEUBEH 528: Computational Neuroscience

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Today's Agenda

- ◆ Course Info and Logistics
- ◆ Motivation
 - ⇒ What is Computational Neuroscience?
 - ⇒ Illustrative Examples
- ◆ Neurobiology 101: Neurons and Networks

Course Information

- ◆ Browse class web page for syllabus and course information:
 - ⇒ <http://www.cs.washington.edu/education/courses/528/>

- ◆ Lecture slides will be made available on the website

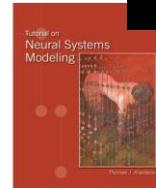
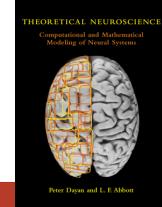
- ◆ Textbooks

- ⇒ Required:

*Theoretical Neuroscience:
Computational and Mathematical Modeling
of Neural Systems* by P. Dayan & L. Abbott

- ⇒ Recommended:

Tutorial on Neural Systems Modelling
by T. Anastasio



Course Topics

- ◆ *Descriptive Models of the Brain*

- ⇒ How is information about the external world *encoded* in neurons and networks? (Chapters 1 and 2)
- ⇒ How can we *decode* neural information? (Chapters 3 and 4)

- ◆ *Mechanistic Models of Brain Cells and Circuits*

- ⇒ How can we reproduce the behavior of a *single neuron* in a computer simulation? (Chapters 5 and 6)
- ⇒ How do we model a *network* of neurons? (Chapter 7)

- ◆ *Interpretive Models of the Brain*

- ⇒ Why do brain circuits operate the way they do?
- ⇒ What are the *computational principles* underlying their operation? (Chapters 7-10)

Course Goals

- ◆ **General Goals: Be able to**
 1. **Quantitatively describe** what a given component of a neural system is doing based on experimental data
 2. **Simulate on a computer** the behavior of neurons and networks in a neural system
 3. **Formulate computational principles** underlying the operation of neural systems
- ◆ We would like to enhance *interdisciplinary cross-talk*
Neuroscience  **Computing and Engineering**
(Experiments, methods, protocols, data, ...) (Computational principles, algorithms, simulation software/hardware, ...)

Workload and Grading

- ◆ Course grade (out of 4.0) will be based on homeworks and a final group project according to:
 - ⇒ Homeworks: 70%
 - ⇒ Final Project: 30%
- ◆ No midterm or final
- ◆ **Homework exercises**: Either written or Matlab-based
 - ⇒ Go over Matlab tutorials and homework on class website
- ◆ **Group Project**: As part of a group of 1-3 persons, investigate a "mini-research" question using methods from this course
 - ⇒ Each group will submit a report and give a presentation

Let's begin...

What is Computational Neuroscience?

Computational Neuroscience

- ◆ “The goal of computational neuroscience is to explain in computational terms how brains generate behaviors” (Sejnowski)
- ◆ Computational neuroscience provides tools and methods for “characterizing *what* nervous systems do, determining *how* they function, and understanding *why* they operate in particular ways” (Dayan and Abbott)
 - ⇒ Descriptive Models (*What*)
 - ⇒ Mechanistic Models (*How*)
 - ⇒ Interpretive Models (*Why*)

An Example: “Receptive Fields”

- ◆ What is the *receptive field* of a brain cell (neuron)?
 - ⇒ Any ideas?

An Example: “Receptive Fields”

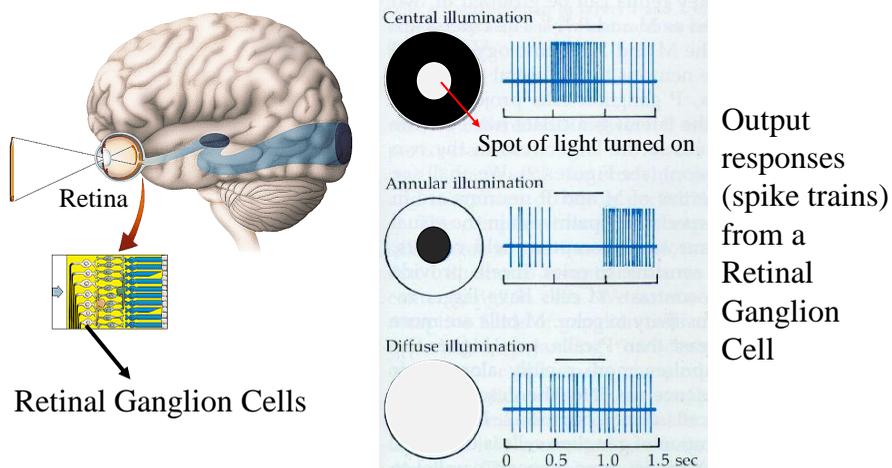
- ◆ What is the *receptive field* of a brain cell (neuron)?
- ◆ Classical Definition: The region of sensory space that activates a neuron (Hartline, 1938)
 - ⇒ Example: Region of the retina where a spot of light activates a retinal cell
- ◆ Current Definition: *Specific properties* of a sensory stimulus that generate a strong response from the cell
 - ⇒ Example: A circular spot of light that turns on at a particular location on the retina

An Example: Cortical Receptive Fields

Let's look at:

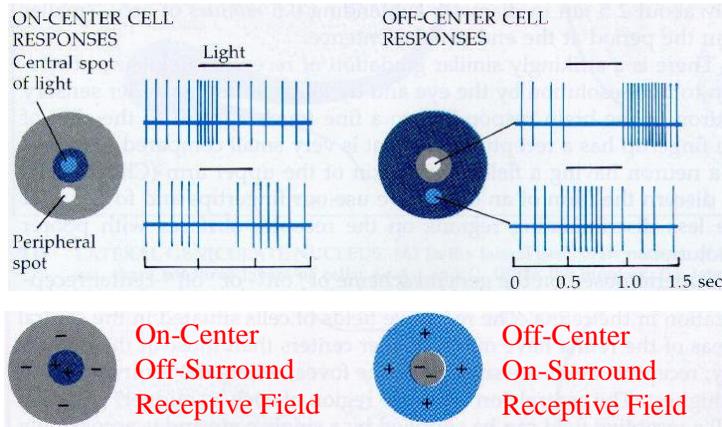
- I. A *Descriptive Model* of Receptive Fields
- II. A *Mechanistic Model* of Receptive Fields
- III. An *Interpretive Model* of Receptive Fields

I. Descriptive Model of Receptive Fields



I. Descriptive Model of Receptive Fields

Mapping a retinal receptive field with spots of light

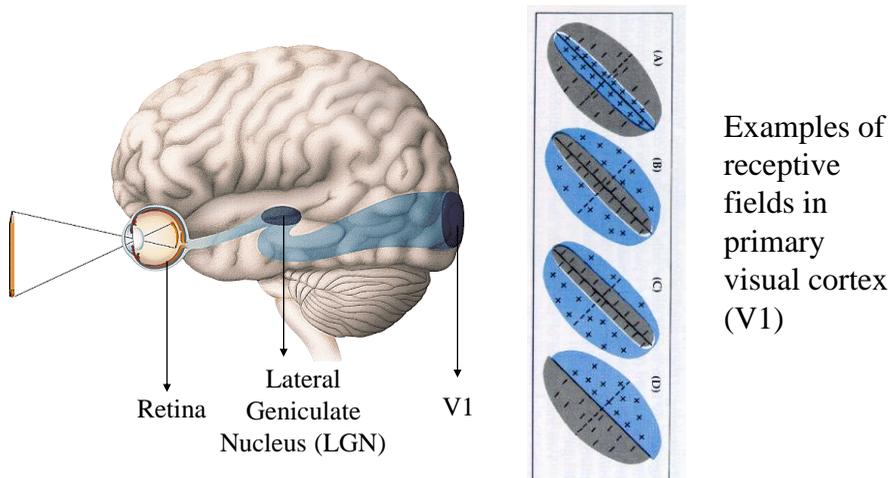


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(From Nicholls et al., 1992)

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Descriptive Models: Cortical Receptive Fields



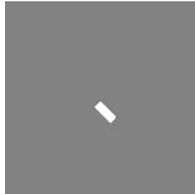
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(From Nicholls et al., 1992)

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Extracting a *Quantitative* Descriptive Model

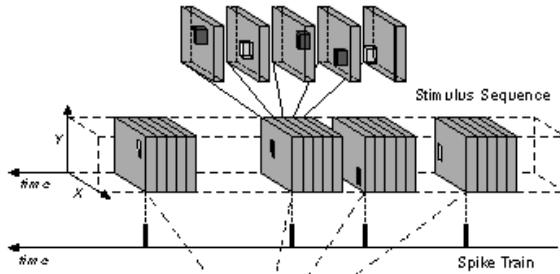
- ◆ The Reverse Correlation Method (Brief intro for now)



Random Bars Sequence
(white noise stimulus)

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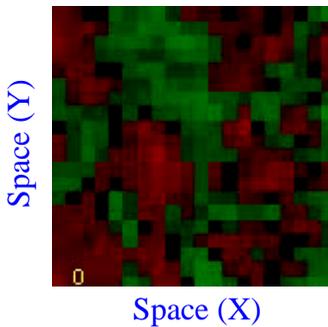


For each output spike, look back and record stimulus sequence occurring before the spike;
Compute the average sequence

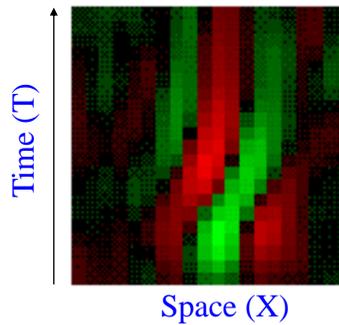
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A Quantitative Model of a V1 Receptive Field

Spatial Receptive Field for $T = 0-300$ ms



Space-Time Receptive Field



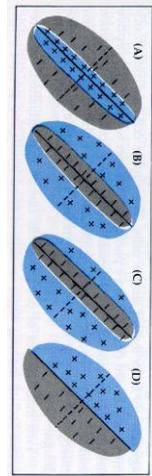
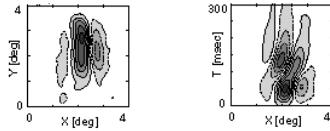
(Copyright 1995, Izumi Ohzawa)

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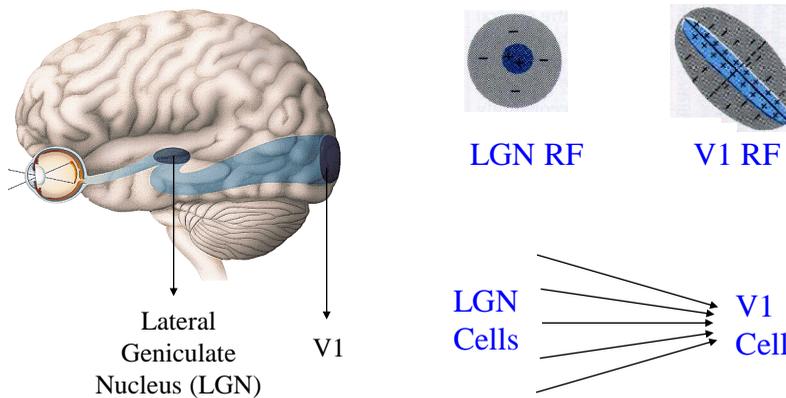
II. Mechanistic Model of Receptive Fields

- ◆ The Question: *How* are receptive fields constructed using the neural circuitry of the visual cortex?

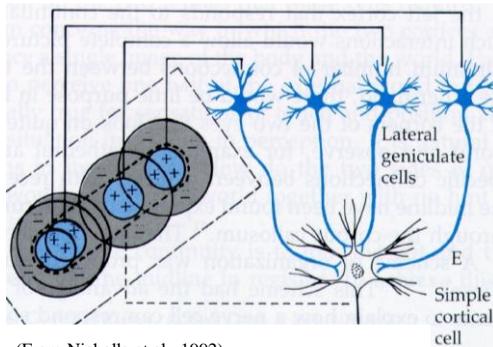


How are these *oriented* receptive fields obtained?

II. Mechanistic Model of Receptive Fields: V1



II. Mechanistic Model of Receptive Fields: V1



(From Nicholls et al., 1992)

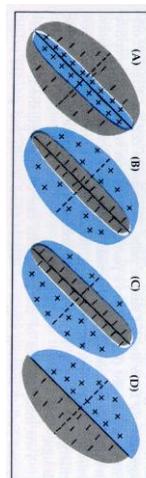
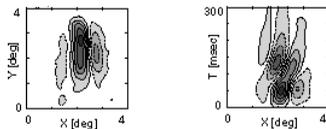
Model suggested by Hubel & Wiesel in the 1960s: V1 RFs are created from converging LGN inputs

Center-surround LGN RFs are *displaced along preferred orientation* of V1 cell

This simple model is still controversial!

III. Interpretive Model of Receptive Fields

◆ The Question: *Why* are receptive fields in V1 shaped in this way?



What are the computational advantages of such receptive fields?

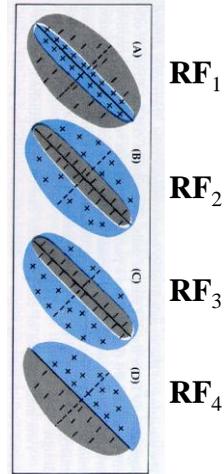
III. Interpretive Model of Receptive Fields

- ◆ **Computational Hypothesis:** Suppose the goal is to *represent images as faithfully and efficiently as possible* using neurons with receptive fields $\mathbf{RF}_1, \mathbf{RF}_2$, etc.

- ◆ Given image \mathbf{I} , want to **reconstruct** \mathbf{I} using neural responses r_1, r_2 etc.:

$$\hat{\mathbf{I}} = \sum_i \mathbf{RF}_i r_i$$

- ◆ **Idea:** Find the \mathbf{RF}_i that *minimize* the squared pixelwise **errors**: $\|\mathbf{I} - \hat{\mathbf{I}}\|^2$ and are as *independent* from each other as possible



III. Interpretive Model of Receptive Fields

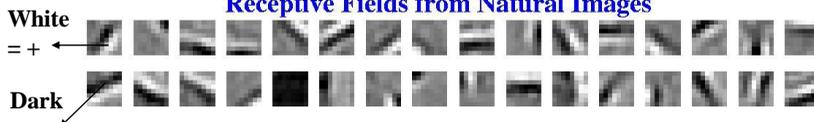
- ◆ Start out with **random** \mathbf{RF}_i and run your algorithm on natural images

Natural Images



□ Receptive Field Size

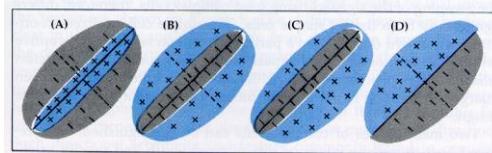
Receptive Fields from Natural Images



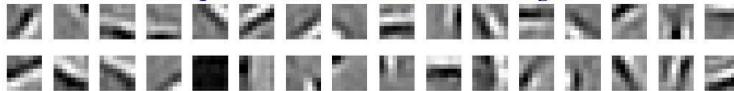
III. Interpretive Model of Receptive Fields

- ◆ **Conclusion:** The brain may be trying to find *faithful and efficient* representations of an animal's natural environment

Receptive Fields in V1



Receptive Fields from Natural Images



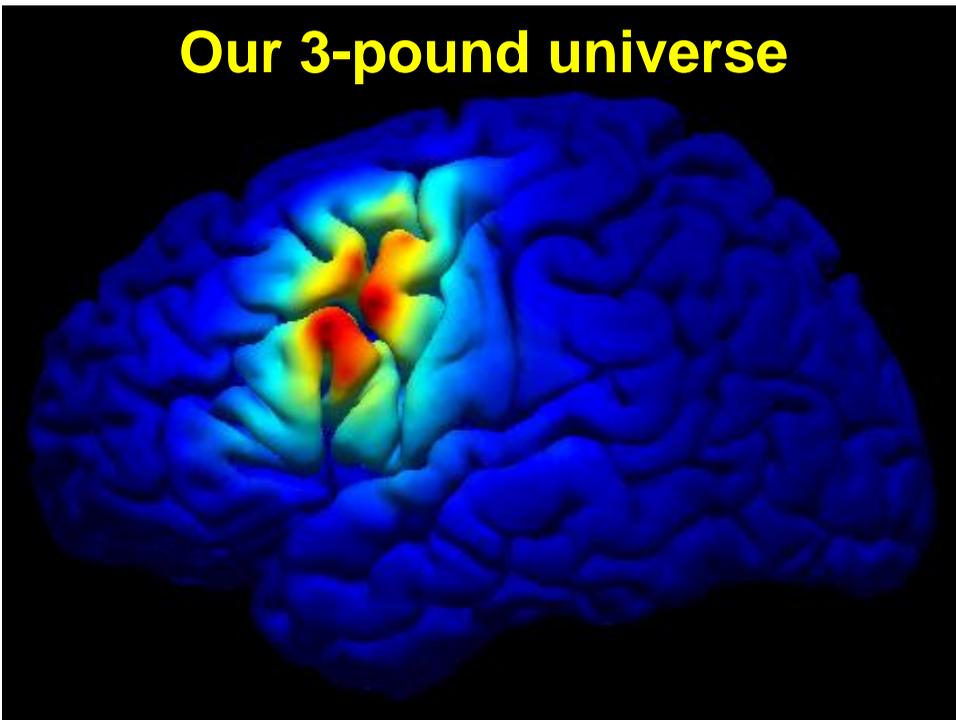
We will explore a variety of *Descriptive*,
Mechanistic, and *Interpretive* models
throughout this course

Neurobiology 101: Brain regions, neurons, and synapses

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Our 3-pound universe



Major Brain Regions: Brain Stem & Cerebellum

Medulla

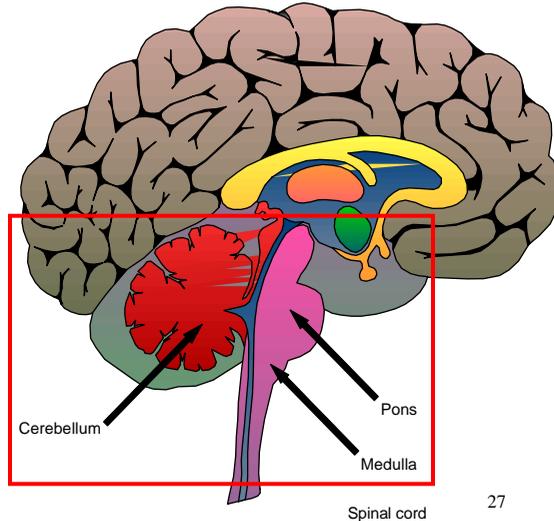
Breathing, muscle tone and blood pressure

Pons

Connects brainstem with cerebellum & involved in sleep and arousal

Cerebellum

Coordination of voluntary movements and sense of equilibrium



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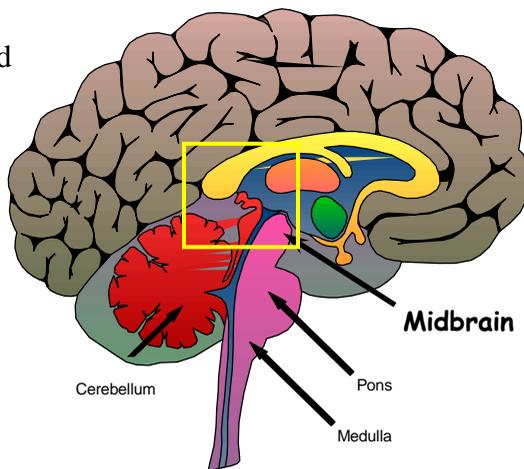
Major Brain Regions: Midbrain & Retic. Formation

Midbrain

Eye movements, visual and auditory reflexes

Reticular Formation

Modulates muscle reflexes, breathing & pain perception. Also regulates sleep, wakefulness & arousal



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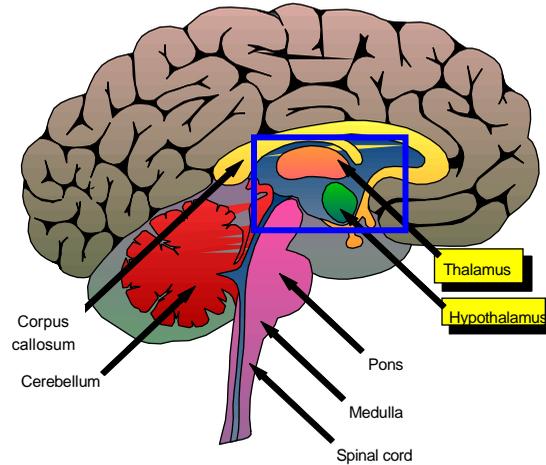
Major Brain Regions: Thalamus & Hypothalamus

Thalamus

“Relay station” for all sensory info (except smell) to the cortex

Hypothalamus

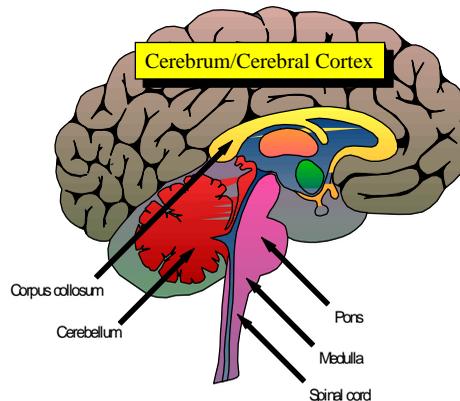
Regulates basic needs
fighting, fleeing,
feeding, and
mating



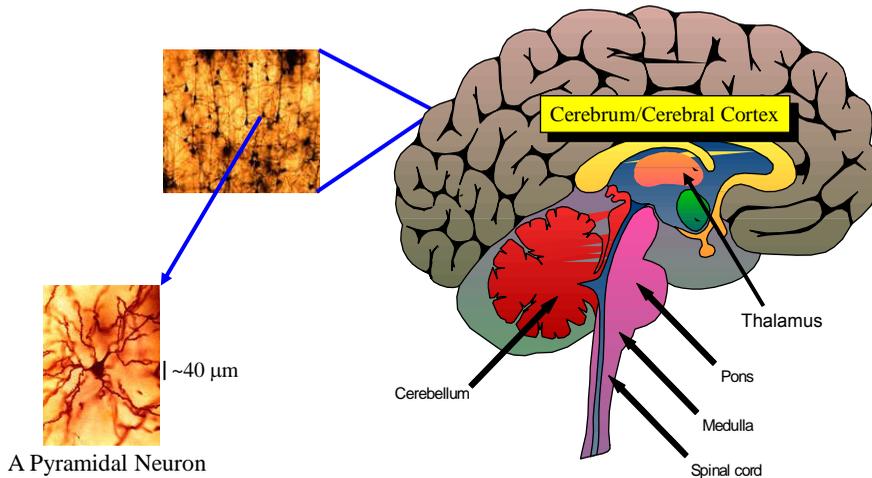
Major Brain Regions: Cerebral Hemispheres

◆ Consists of: Cerebral cortex, basal ganglia, hippocampus, and amygdala

◆ Involved in perception and motor control, cognitive functions, emotion, memory, and learning

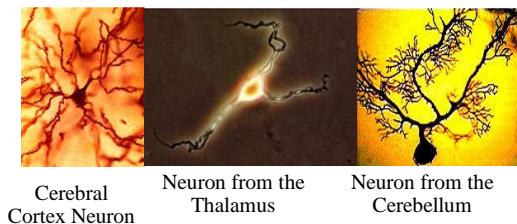


Enter...the neuron (“brain cell”)



A Pyramidal Neuron

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Cerebral Cortex Neuron

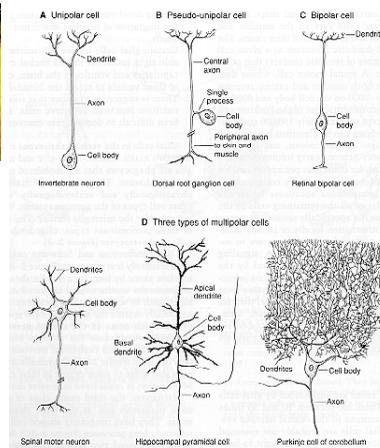
Neuron from the Thalamus

Neuron from the Cerebellum

Neuron Doctrine/ Dogma:

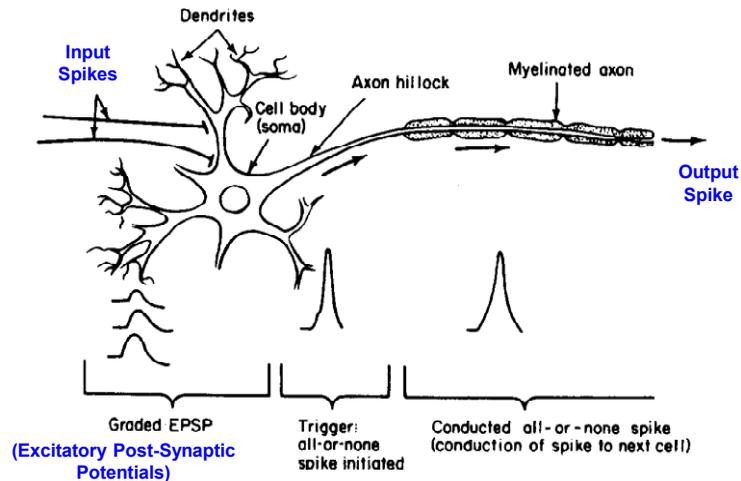
“The neuron is the appropriate basis for understanding the computational and functional properties of the brain”

First suggested in 1891 by Waldeyer



From Kandel, Schwartz, Jessel, Principles of Neural Science, 3rd edn., 1991, pg. 21

The Idealized Neuron

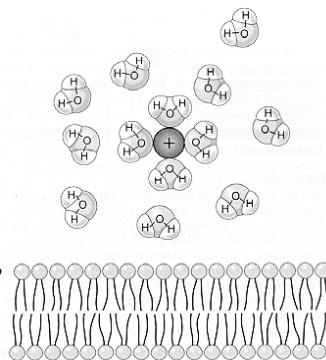


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What is a Neuron?

- ◆ A “leaky bag of charged liquid”
- ◆ Contents of the neuron enclosed within a *cell membrane*
- ◆ Cell membrane is a *lipid bilayer*
 - ⇒ Bilayer is impermeable to charged ions such as Na^+ , Cl^- , K^+ , and Ca^{2+}



From Kandel, Schwartz, Jessel, Principles of Neural Science, 3rd edn., 1991, pg. 67

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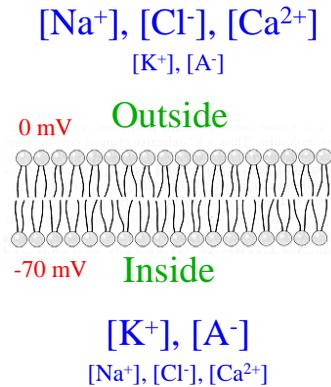
The Electrical Personality of a Neuron

- ◆ Each neuron maintains a *potential difference* across its membrane

- ↳ Inside is -70 to -80 mV relative to outside

- ↳ Ion concentration $[\text{Na}^+]$, $[\text{Cl}^-]$ and $[\text{Ca}^{2+}]$ higher outside; $[\text{K}^+]$ and organic anions $[\text{A}^-]$ higher inside

- ↳ *Ionic pump* maintains -70 mV difference by expelling Na^+ out and allowing K^+ ions in

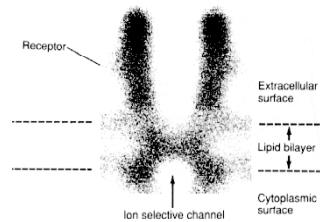
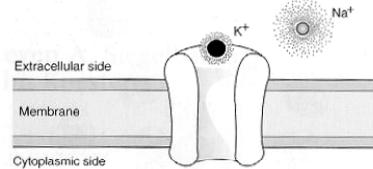


Influencing a Neuron's Electrical Personality

How can the electrical potential difference be changed in local regions of a neuron?

Membrane Proteins: The Gatekeepers

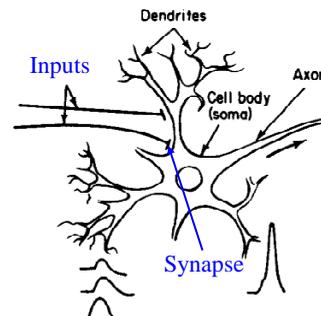
- ◆ Proteins in membranes act as **pores** or **channels** that allow specific ions to pass through.
 - ⇒ E.g. Pass K^+ but not Cl^- or Na^+
- ◆ These “ionic channels” are *gated*
 - ⇒ **Voltage-gated**: Probability of opening depends on membrane voltage
 - ⇒ **Chemically-gated**: Binding to a chemical causes channel to open
 - ⇒ **Mechanically-gated**: Sensitive to pressure or stretch



From Kandel, Schwartz, Jessel, Principles of Neural Science, 3rd edn., 1991, pgs. 68 & 137

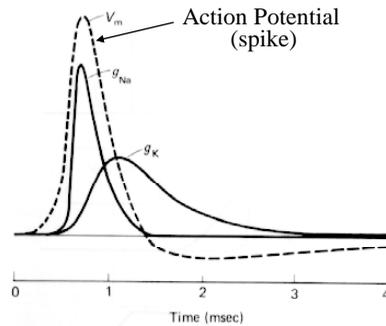
Gated Channels allow Neuronal Signaling

- ◆ Inputs from other neurons → **chemically-gated channels** (at “**synapses**”) → Changes in local membrane potential
- ◆ This causes opening/closing of **voltage-gated channels** in dendrites, body, and axon, resulting in **depolarization** (positive change in voltage) or **hyperpolarization** (negative change)



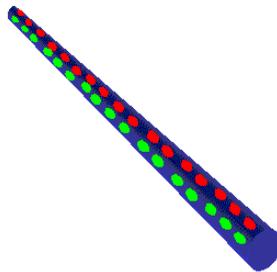
The Output of a Neuron: Action Potentials

- ◆ Voltage-gated channels cause action potentials (spikes)
 1. Rapid Na^+ influx causes rising edge
 2. Na^+ channels deactivate
 3. K^+ outflux restores membrane potential
- ◆ Positive feedback causes spike
 - ⇒ Na^+ influx increases membrane potential, causing *more* Na^+ influx



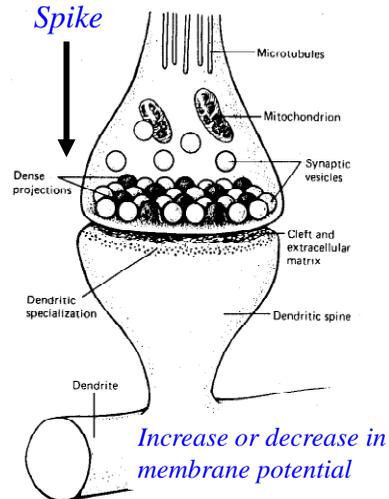
From Kandel, Schwartz, Jessel, Principles of Neural Science, 3rd edn., 1991, pg. 110

Propagation of a Spike along an Axon



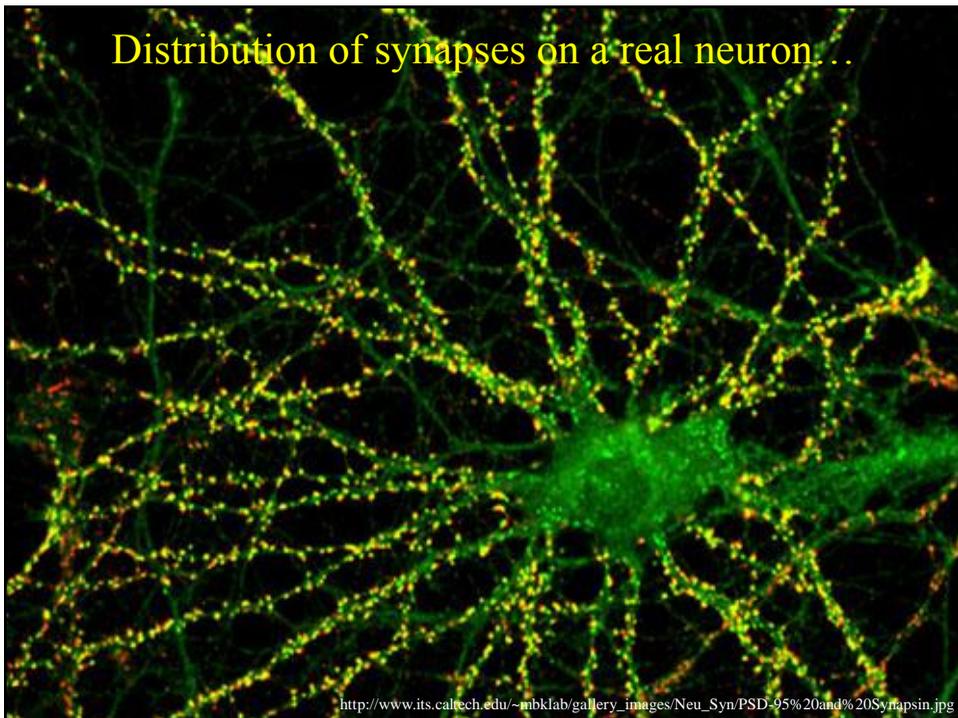
Communication between Neurons: Synapses

- ◆ Synapses are the “connections” between neurons
 - ⇒ **Electrical** synapses (gap junctions)
 - ⇒ **Chemical** synapses (use neurotransmitters)
- ◆ Synapses can be excitatory or inhibitory
- ◆ Synapse Doctrine: Synapses are the basis for **memory** and **learning**



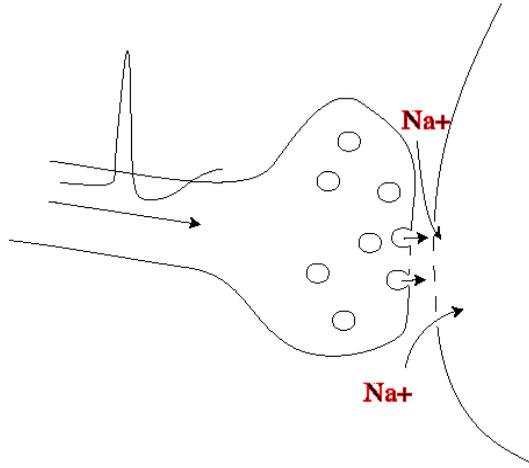
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Distribution of synapses on a real neuron...



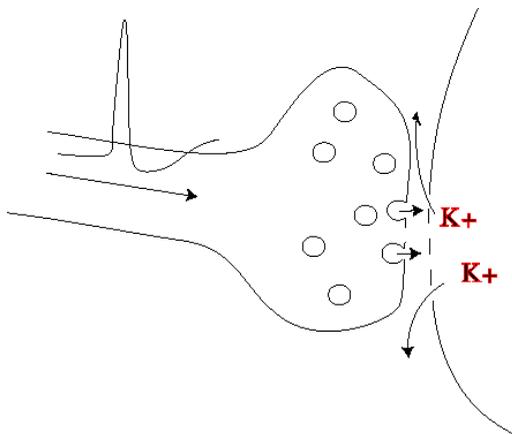
http://www.its.caltech.edu/~mbklab/gallery_images/Neu_Syn/PSD-95%20and%20Synapsin.jpg

An **Excitatory** Synapse



Input spike →
Neurotransmitter
release →
Binds to Na
channels (which
open) →
Na⁺ influx →
Depolarization due
to EPSP (excitatory
postsynaptic
potential)

An **Inhibitory** Synapse



Input spike →
Neurotransmitter
release →
Binds to K
channels →
K⁺ leaves cell →
Hyperpolarization due
to IPSP (inhibitory
postsynaptic potential)

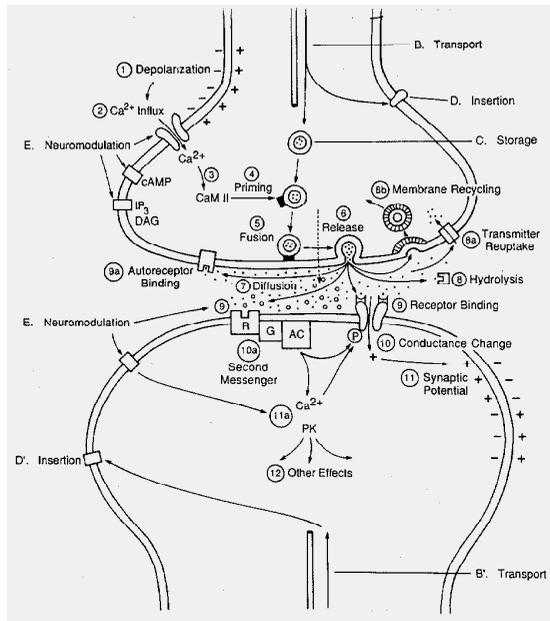
Down in the Synaptic Engine Room

A reductionist's dream! (or nightmare?)

Note: Even this is a simplification!

From Kandel, Schwartz, Jessel, Principles of Neural Science, 3rd edn., 1991

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Synaptic plasticity: Adapting the connections

- ◆ **Long Term Potentiation (LTP)**: Increase in synaptic strength that lasts for several hours or more
 - ⇒ Measured as an increase in the excitatory postsynaptic potential (EPSP) caused by presynaptic spikes

LTP observed as an increase in size or slope of EPSP for the same presynaptic input



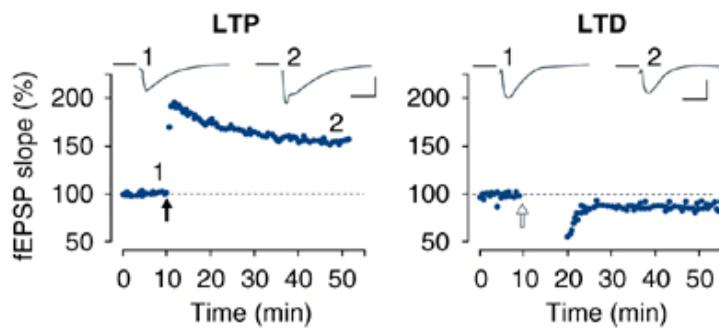
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Types of Synaptic Plasticity

- ◆ **LTP**: synaptic strength increases after prolonged pairing of presynaptic and postsynaptic spiking (*correlated firing of two connected neurons*).
- ◆ **Long Term Depression (LTD)**: Reduction in synaptic strength that lasts for several hours or more

Example of measured synaptic plasticity



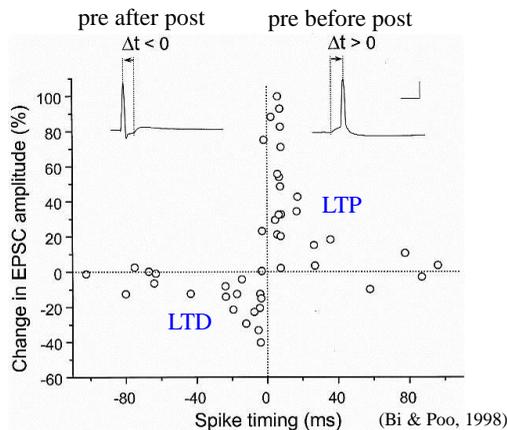
(From: http://www.nature.com/npp/journal/v33/n1/fig_tab/1301559f1.html)

Types of Synaptic Plasticity

- ◆ **LTP**: synaptic strength increases after prolonged pairing of presynaptic and postsynaptic spiking (*correlated firing of two connected neurons*).
- ◆ **Long Term Depression (LTD)**: Reduction in synaptic strength that lasts for several hours or more
- ◆ **Spike-Timing Dependent Plasticity**: LTP/LTD depends on relative timing of pre/postsynaptic spiking

Spike-Timing Dependent Plasticity

- ◆ Amount of increase or decrease in synaptic strength (LTP/LTD) depends on **relative timing of pre & postsynaptic spikes**



Comparing Neural versus Digital Computing

- ◆ **Device count:**
 - ⇒ Human Brain: 10^{11} neurons (each neuron $\sim 10^4$ connections)
 - ⇒ Silicon Chip: 10^{10} transistors with sparse connectivity
- ◆ **Device speed:**
 - ⇒ Biology has up to $100\mu\text{s}$ temporal resolution
 - ⇒ Digital circuits will soon have a 100ps clock (10 GHz)
- ◆ **Computing paradigm:**
 - ⇒ Brain: Massively parallel computation & adaptive connectivity
 - ⇒ Digital Computers: sequential information processing via CPUs with fixed connectivity
- ◆ **Capabilities:**
 - ⇒ Digital computers excel in math & symbol processing...
 - ⇒ Brains: Better at solving ill-posed problems (vision, speech)

Conclusions and Summary

- ◆ Structure and organization of the brain suggests **computational analogies**
 - ⇒ **Information storage:** Physical/chemical structure of neurons and synapses
 - ⇒ **Information transmission:** Electrical and chemical signaling
 - ⇒ **Primary computing elements:** Neurons
 - ⇒ **Computational basis:** **Currently unknown** (but inching closer)
- ◆ We can understand neuronal computation by understanding the underlying primitives through:
 - ⇒ **Descriptive models**
 - ⇒ **Mechanistic models**
 - ⇒ **Interpretive models**

Next Class

- ◆ Descriptive Models

- ⇒ **Neural Encoding**

- ◆ Things to do:

- ⇒ Visit course website

- <http://www.cs.washington.edu/education/courses/528/>

- ⇒ Matlab practice: Homework 0 and tutorials online

- ⇒ Read Chapter 1 in Dayan & Abbott textbook