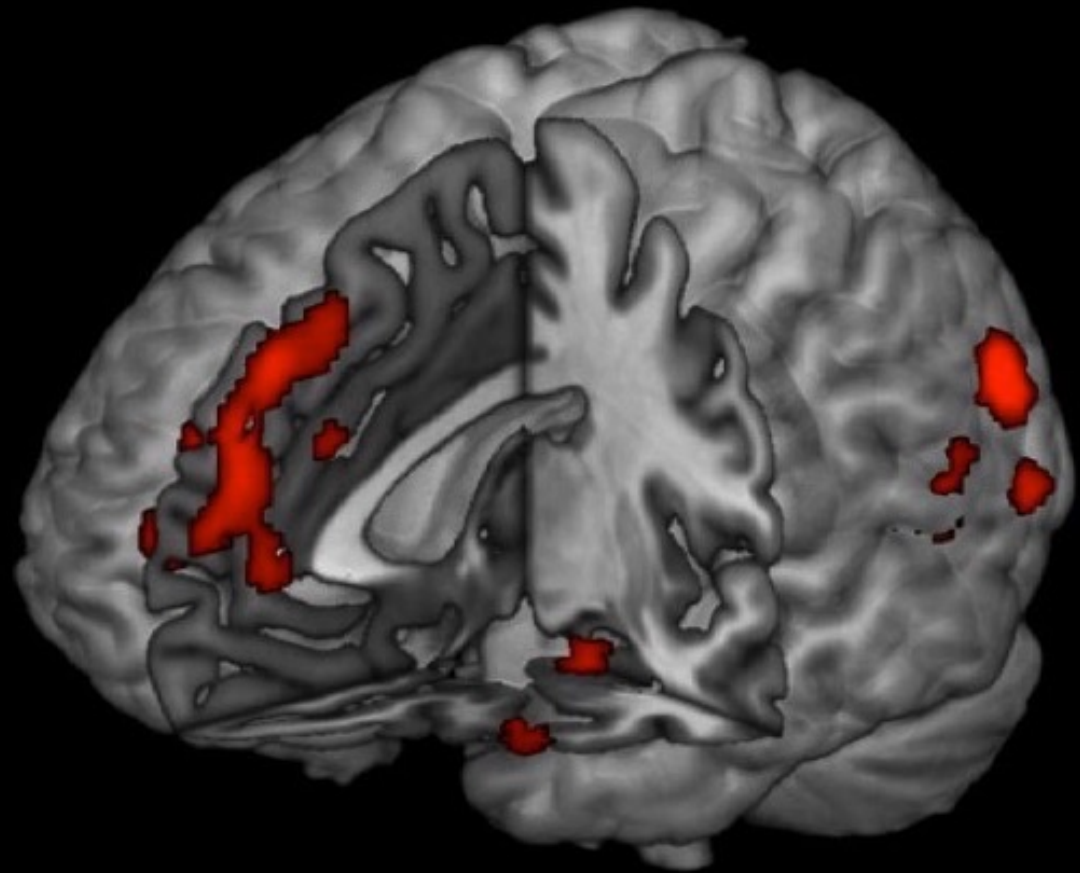


An Introduction to Neuroscience

Dr. Olav E Krigolson
krigolson@gmail.com



Administrative Stuff

Attendance

Exams

Expectations

Course Outline

Week I: Neurons

Condition: Amyotrophic Lateral Sclerosis (ALS)

Topic: Neurons

Technique: Single Unit Recordings

October 4th, 2023

Week II: Sensory Perception

Condition: Phantom Limbs

Topic: Proprioception and Audition

Technique: Functional Imaging

October 11th, 2023

Week III: Vision

Condition: Aging

Topic: Vision

Technique: Electroencephalography

October 18th, 2023

Week IV: Motor Control

Condition: Parkinson's Disease

Topic: Movement Planning and Control

Technique: Transcranial Magnetic Stimulation

October 25th, 2023

Week V: Attention

Condition: Neglect

Topic: Attention

Technique: Patient Studies

November 1st, 2023

Week VI: Memory

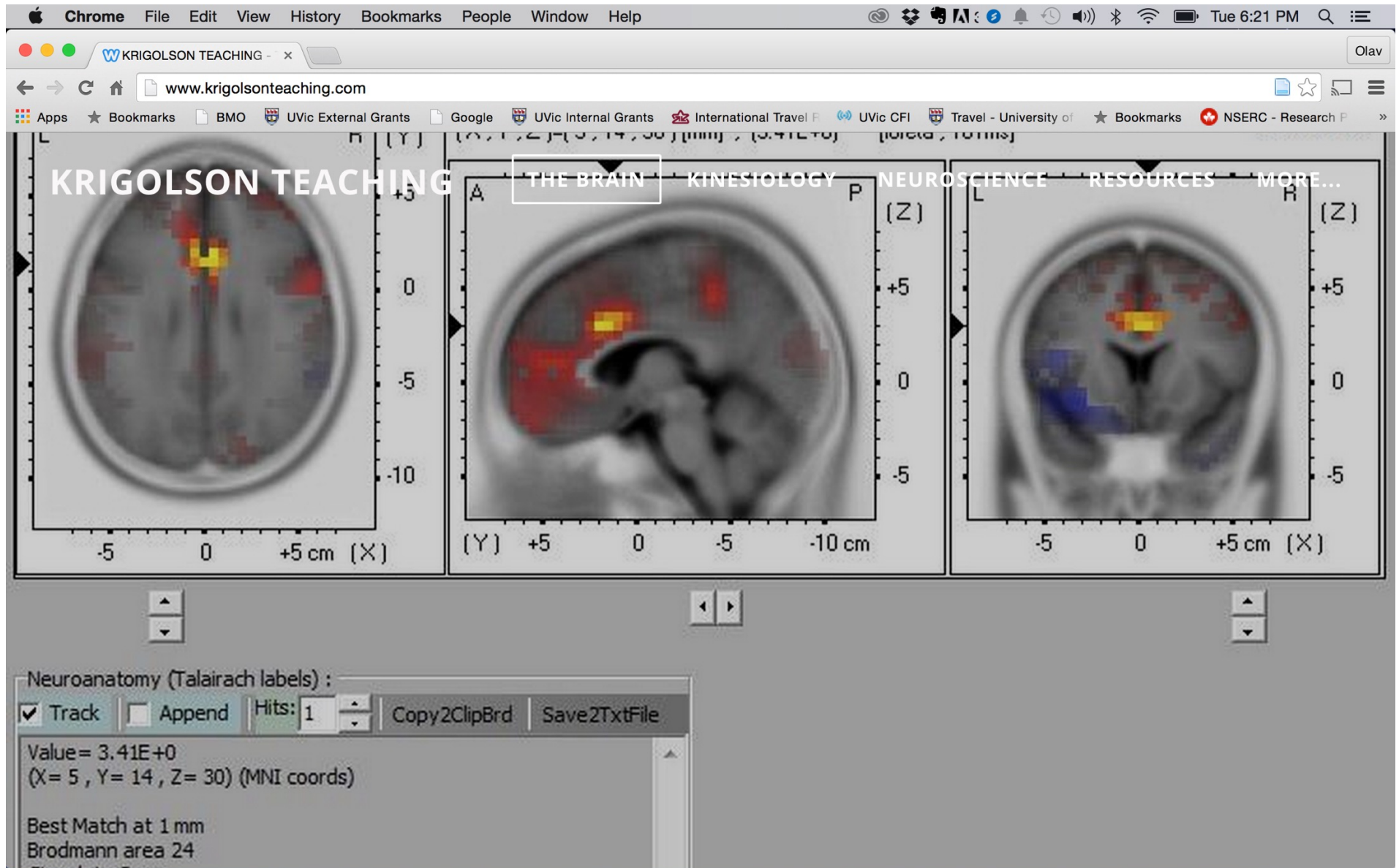
Condition: Alzheimer's Disease

Topic: Memory

Technique: Optical Imaging

November 8th, 2023

www.krigolsonteaching.com



Me

Formal Training

Expertise

Style

Disclaimers

Knowing Everything

Scope of Course

BS in the Media

Dr. Phil

Pace of Research

Lecture Format

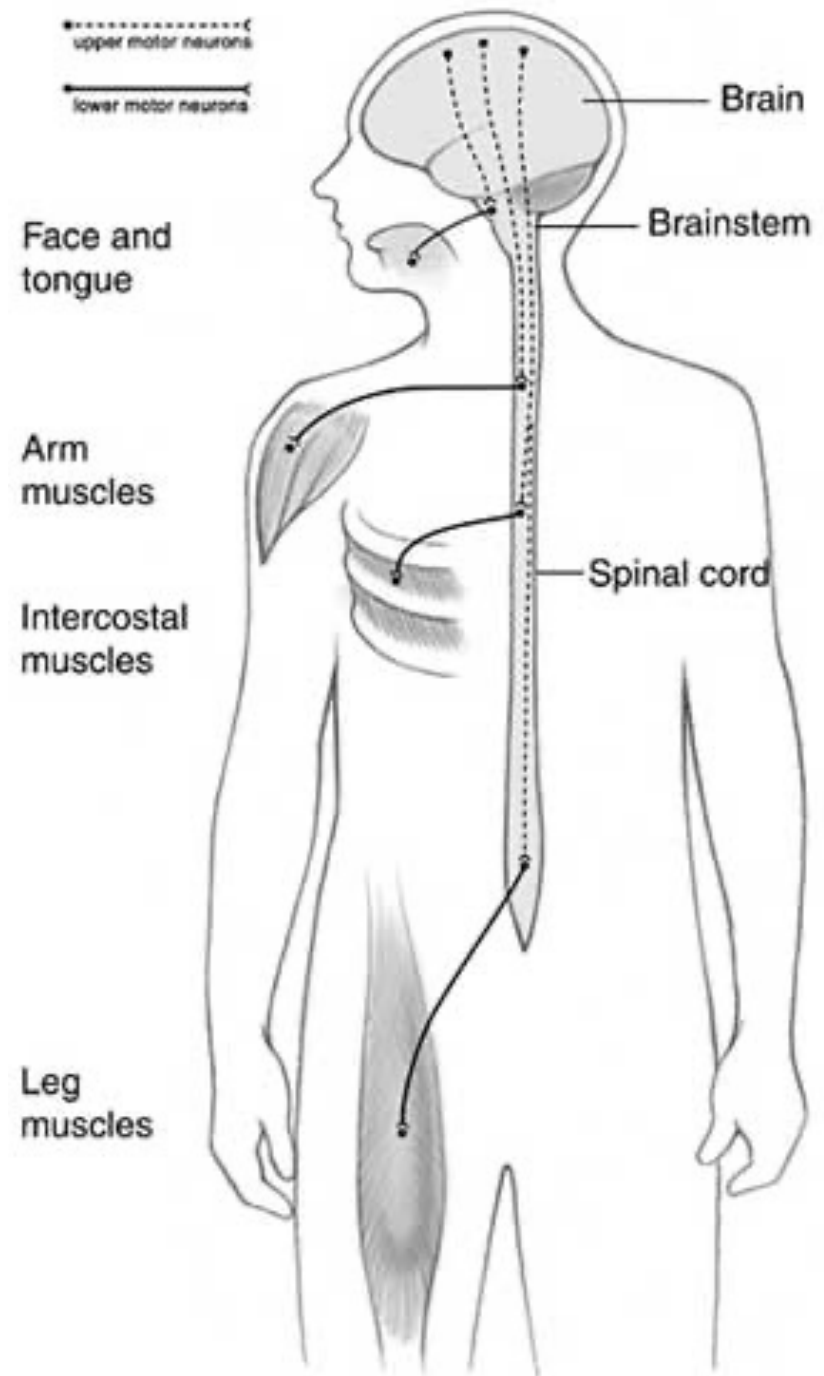
1. Clinical Issue (relevant to topic)
2. Topic
3. Technique

Topic One: Neurons

Lecture 1A:
Amyotrophic Lateral Sclerosis

What is amyotrophic lateral sclerosis?

It is a progressive neurological disease that affects the control of muscle movement due to its damaging effects on motor neurons in the spinal cord and the brain



Significance of the Name of this Disease

- A-myotrophic comes from Greek
- “A” = no/negative
- “myo” = muscle
- “trophic” = nourishment
- “No Muscle Nourishment”

- Lateral = defines location of the nerve cells that signal and control the muscles

- Sclerosis = scarring and hardening in the degenerating region

Other common names for this disease:

- Motor neuron disease
- Charcot's disease
- Lou Gehrig's disease

Nature and Characteristics of ALS

- Forms:
 - Two types of ALS:
 - Sporadic – no family history
 - Familial – family history/background
 - 90% of the known cases are sporadic

Who Gets ALS?

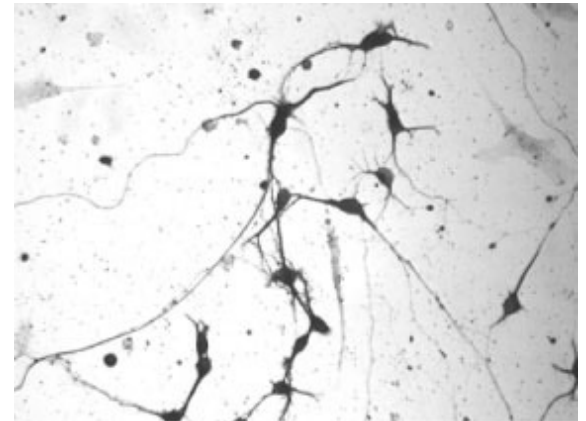
- According to the ALS CARE Database, 60% of the people with ALS in the database are men and 93% of patients in the database are caucasian
- Normally occurs in people between 40 to 70 years of age
 - Also can occur in people in their 20' s and 30' s

Mechanism of ALS

- starting point is a mutation of Chromosome 21 (also suspect in many other conditions, probably the most common being Down's)
- Most Common: the mutation changes the SOD1 gene/protein (currently over 21 "types of ALS" though)
- SOD1 change results in superoxide radicals not being neutralized
- The radicals "attack" the motor neurons and degrade them (many effects here – inflammation? over-excitation?)
- Muscles are not able to be stimulated

Symptoms of ALS

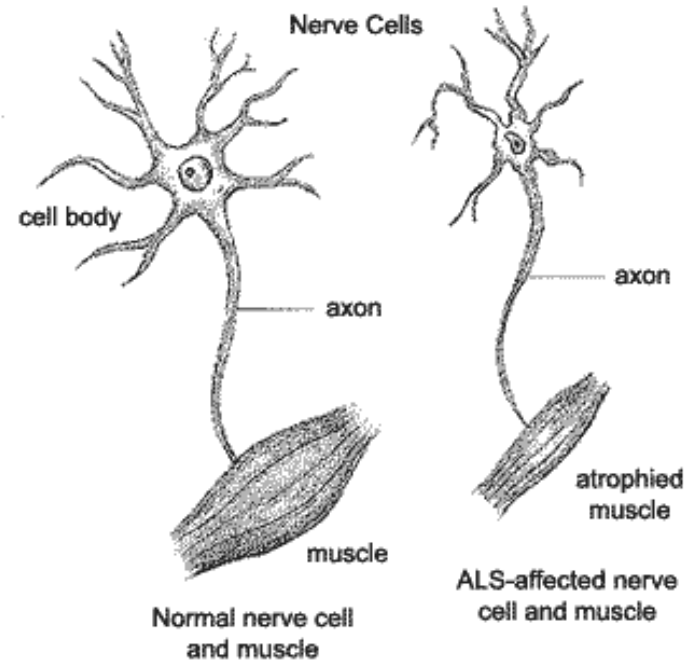
- First signs and symptoms (frequently overlooked)
 - Twitching and cramping of muscles (especially in hands and feet)
 - Stiffness
 - Weakness (especially in hands, arms and legs)
 - Slurred speech



Picture taken from the National Institute of Aging

Symptoms continued . . .

- Later signs and symptoms:
 - Difficulty chewing and swallowing
 - Shortness of breath
 - Muscle weakness due to wasting away of muscles
 - Causes muscles to become smaller
 - Respiratory failure
 - Paralysis



Picture from the ALS Association

This picture from the Neuromuscular website shows the wasting away of a person's hands and arms

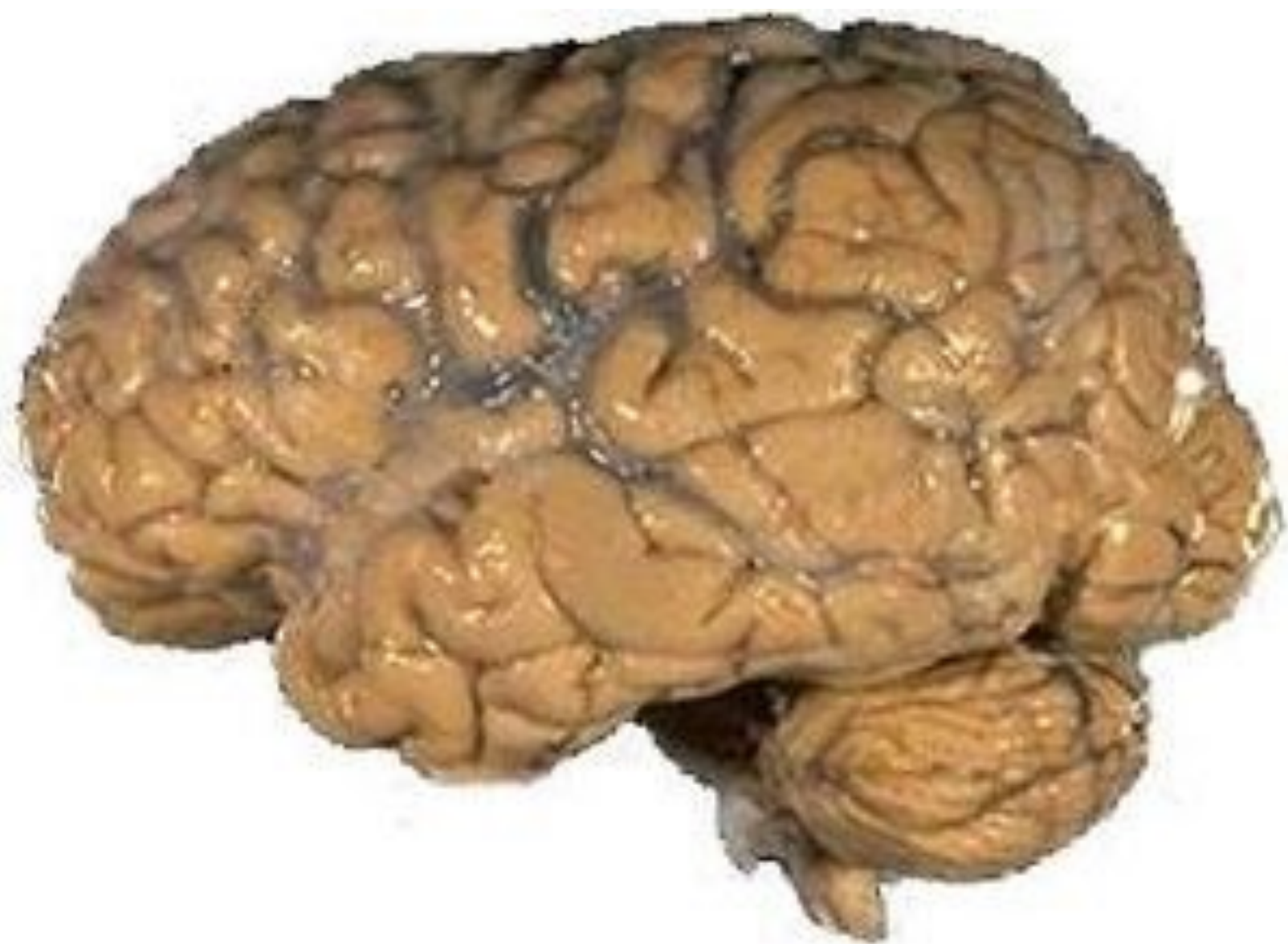


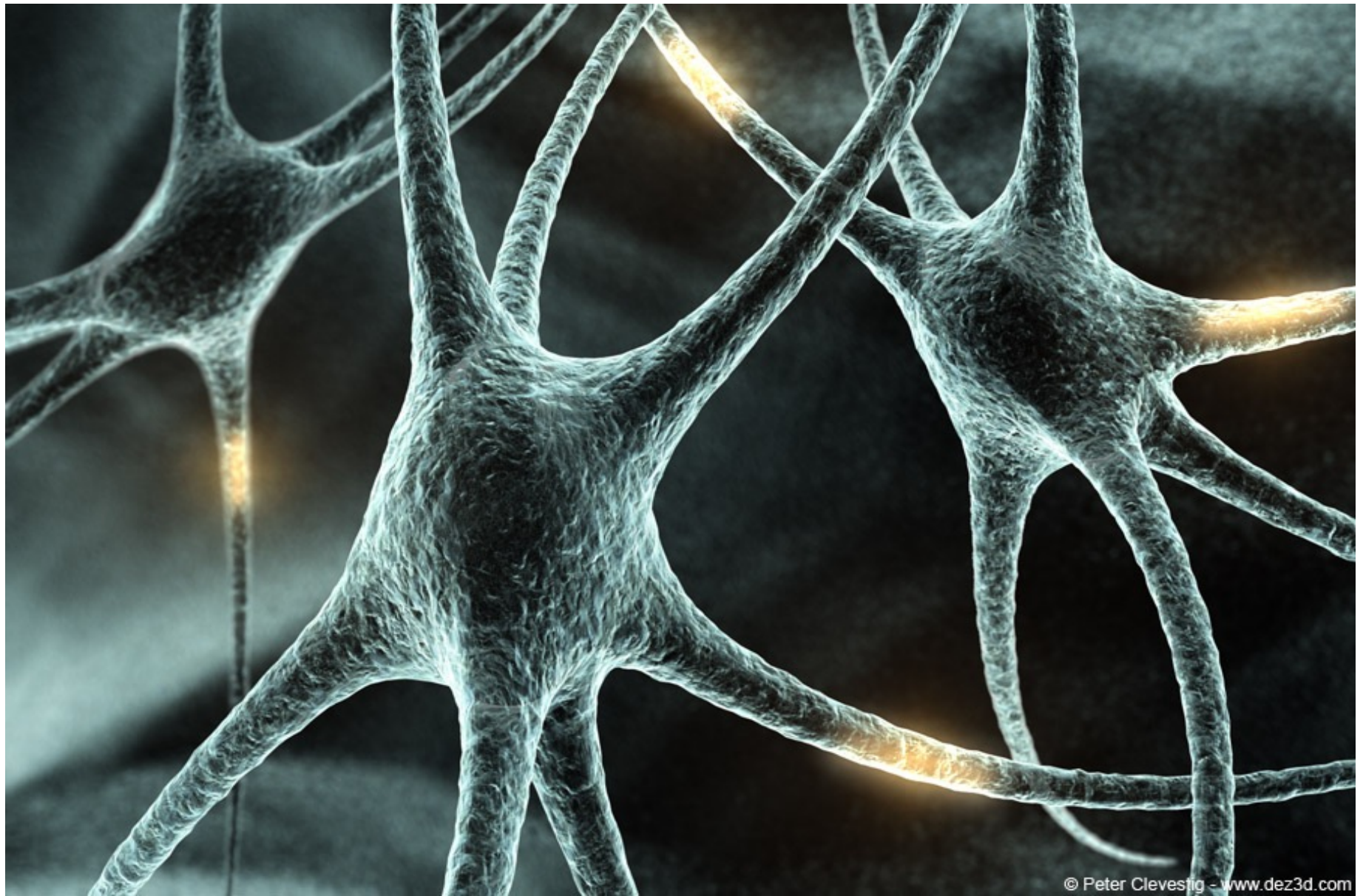
- Symptoms, or patterns of symptoms, are not the same for each ALS individual
- However, progressive muscle weakness and paralysis are universally experienced
- Since ALS attacks only motor neurons, the sense of sight, touch, hearing, taste, and smell are not affected
- Patients usually only live 3 to 5 years after they are diagnosed
- There are some cases; however, where individuals have lived 10 or more years

Diagnosing ALS

- 5,600 people in the US are diagnosed with ALS each year (about 700 in Canada)
- ALS is a very difficult disease to diagnose

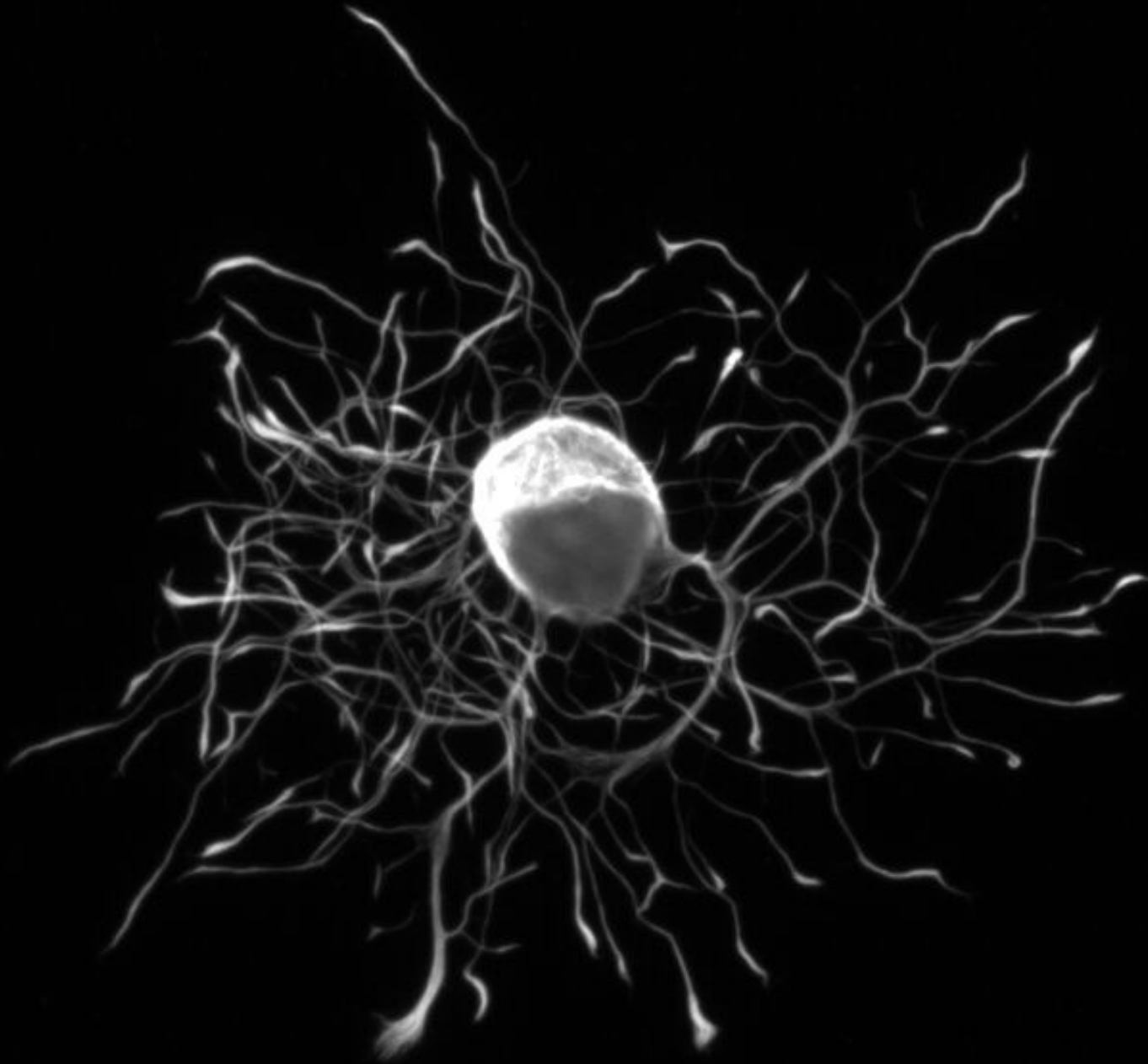
Lecture 1B: Neurons

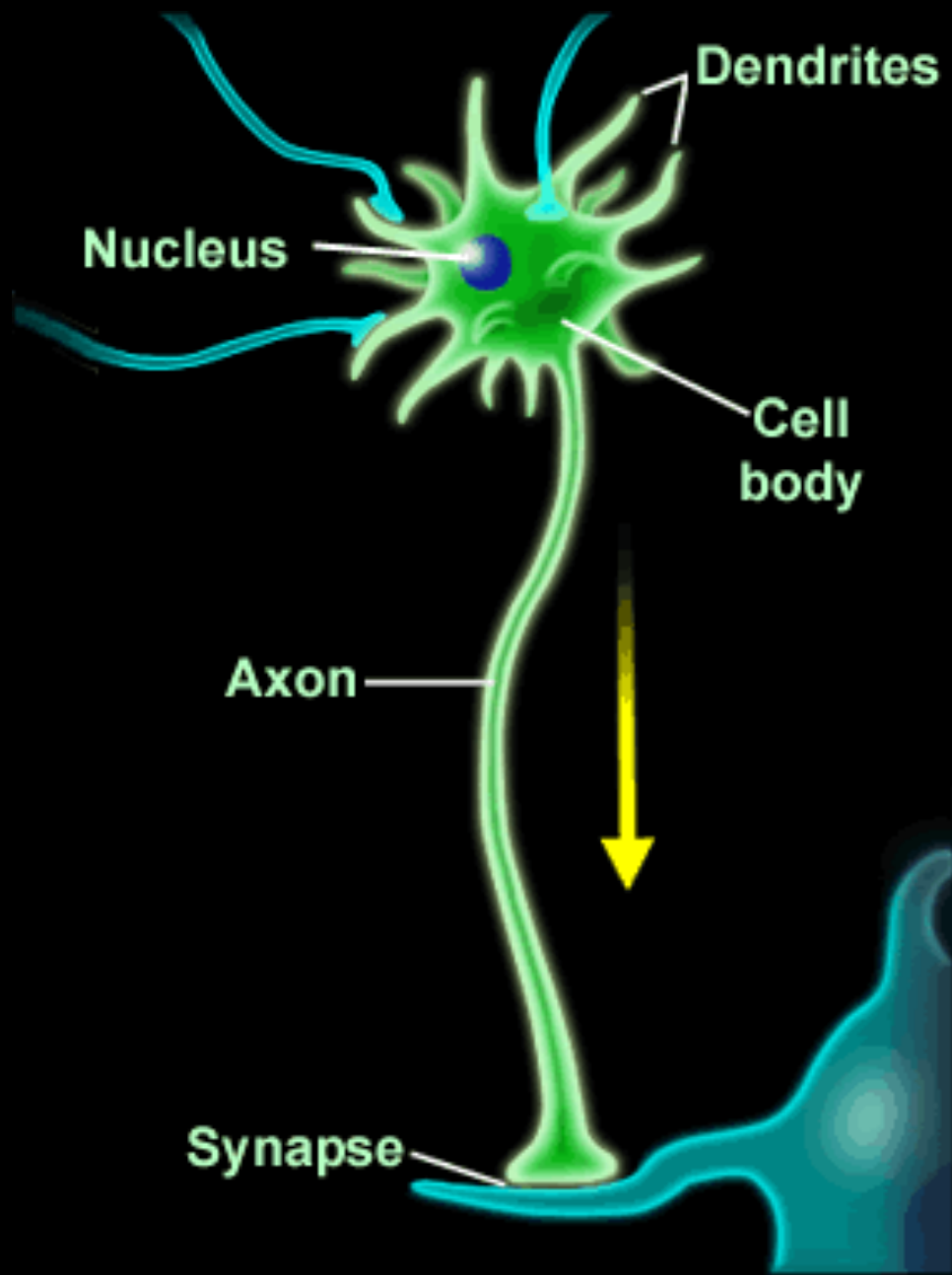


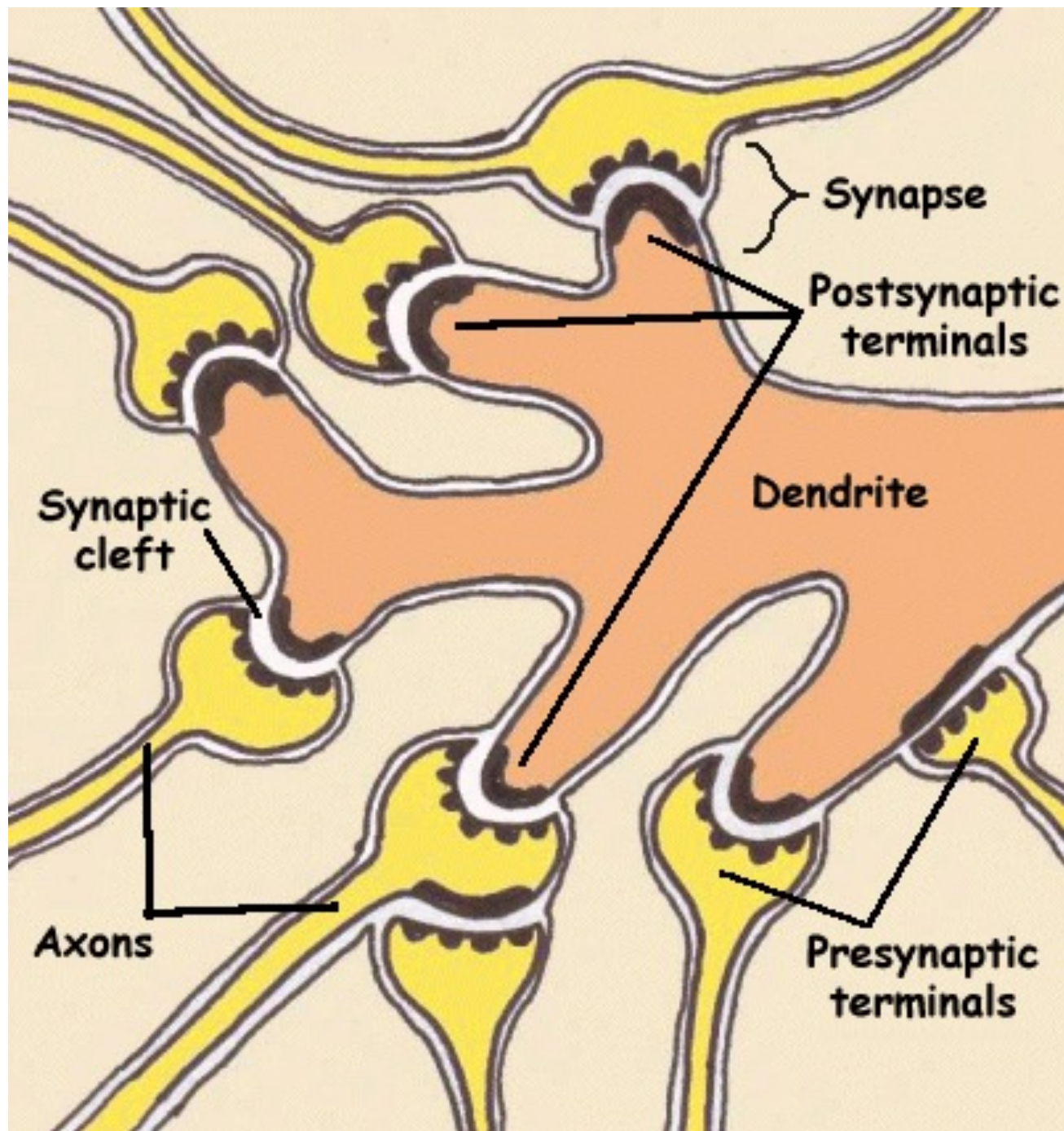


Human Brain: 86 billion neurons
0.15 quadrillion nerve synapses (million x billion)

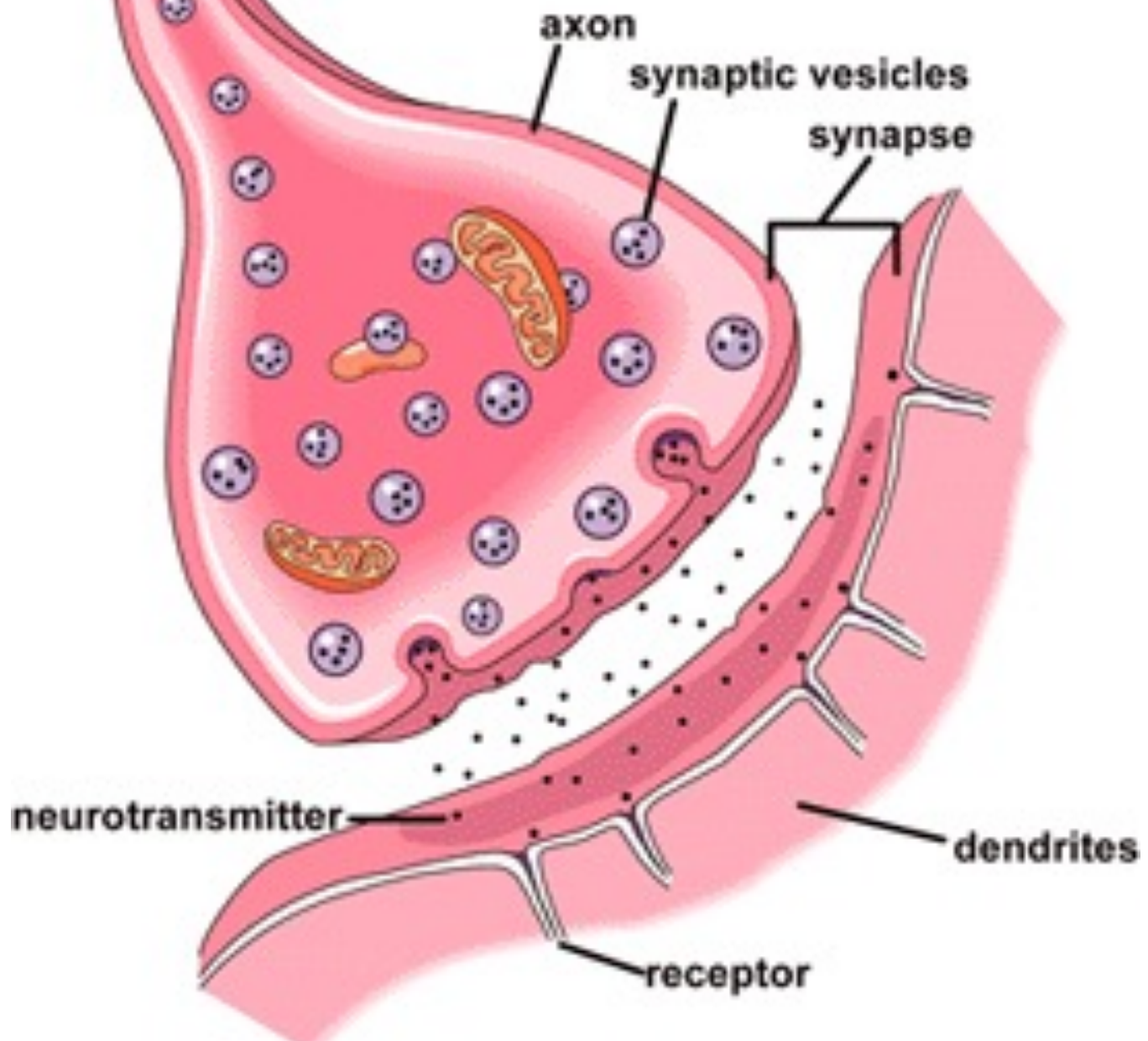
Human Brain: 150,000 km of nerve cells

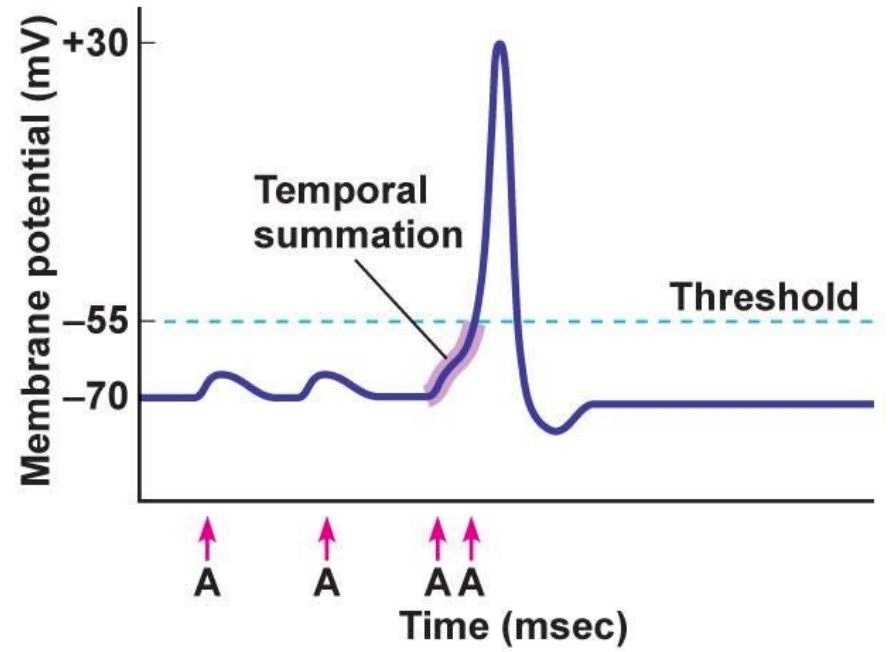
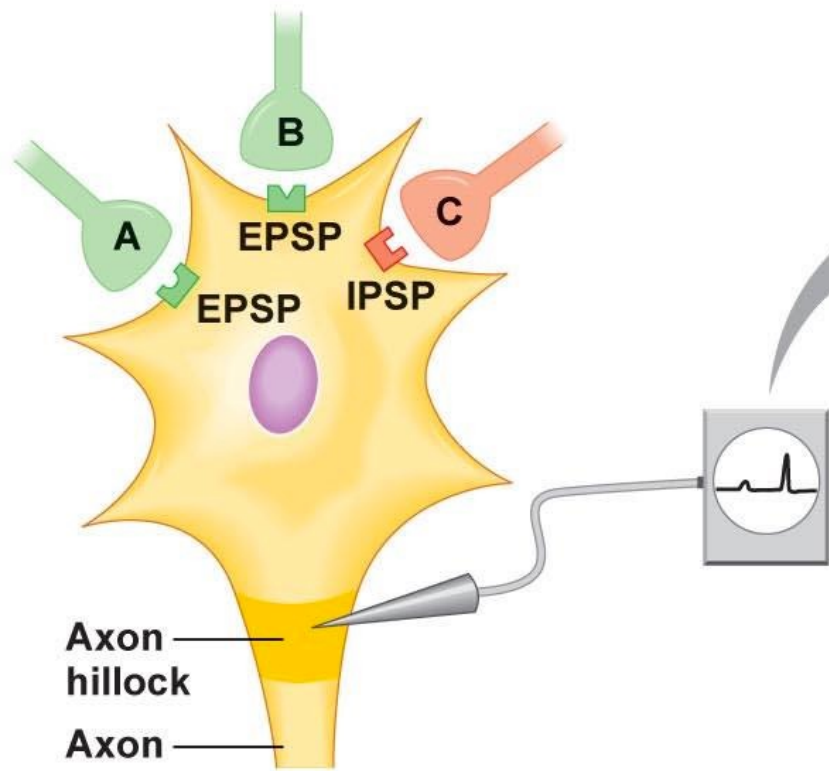




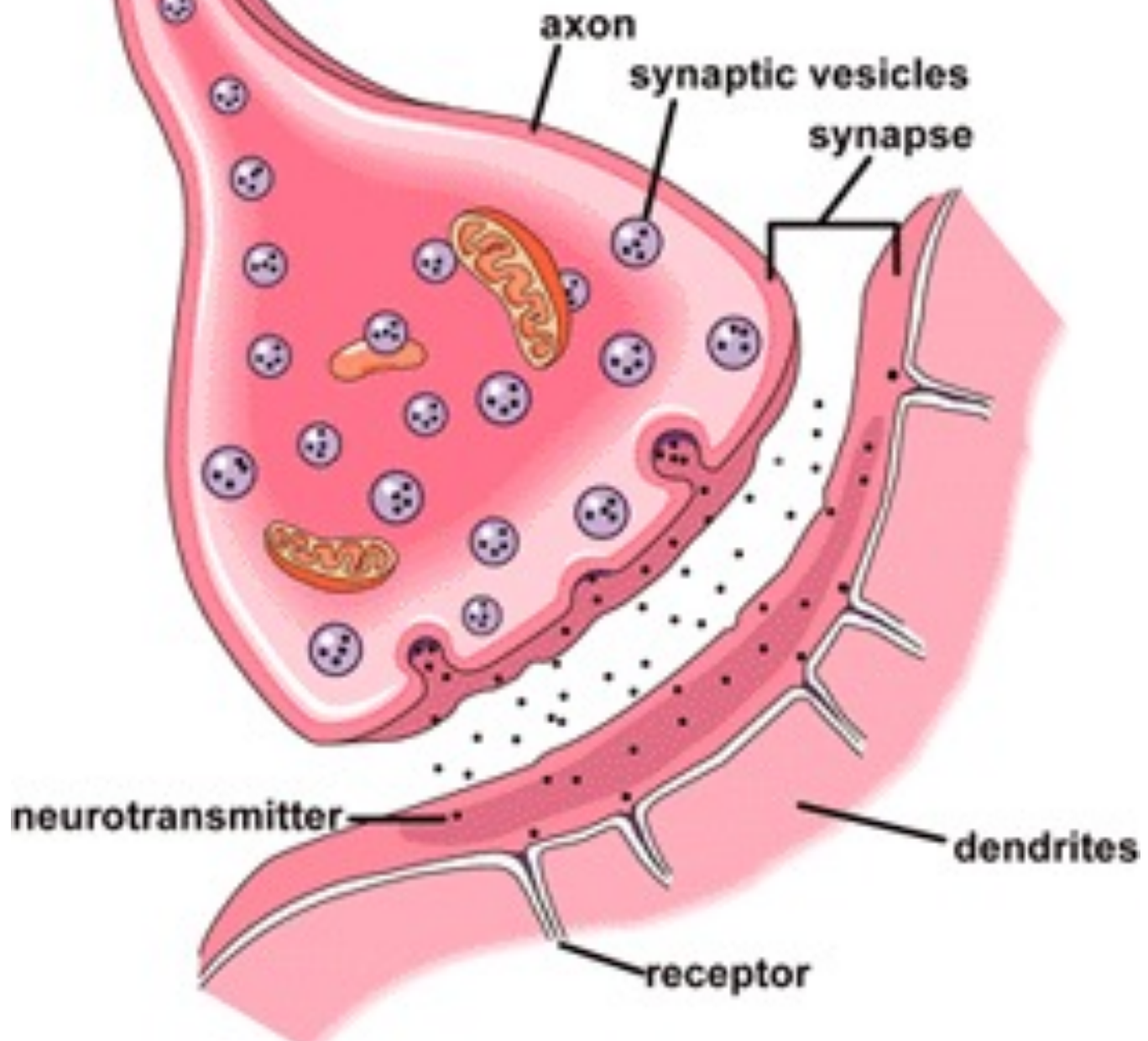


Synapse



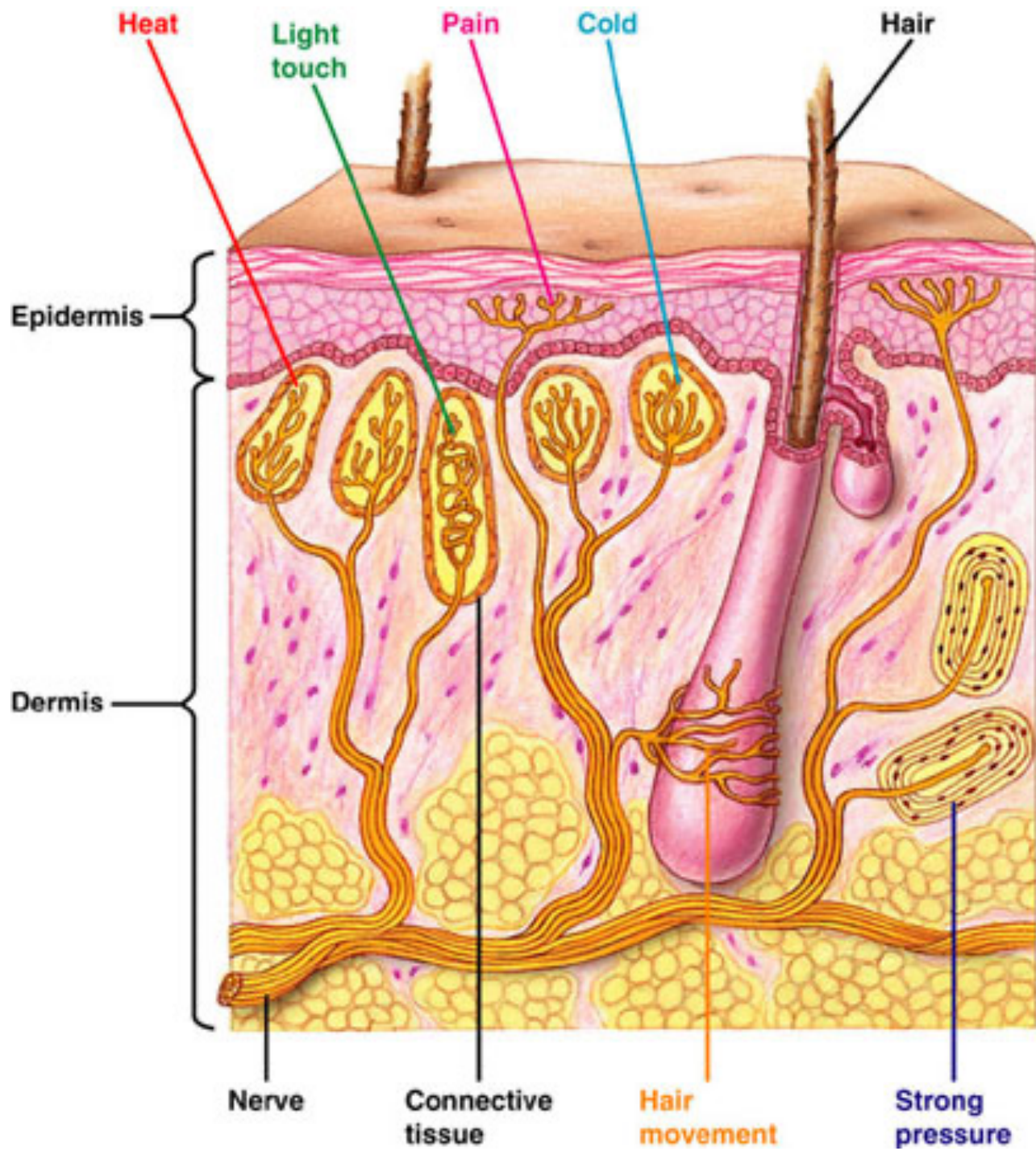


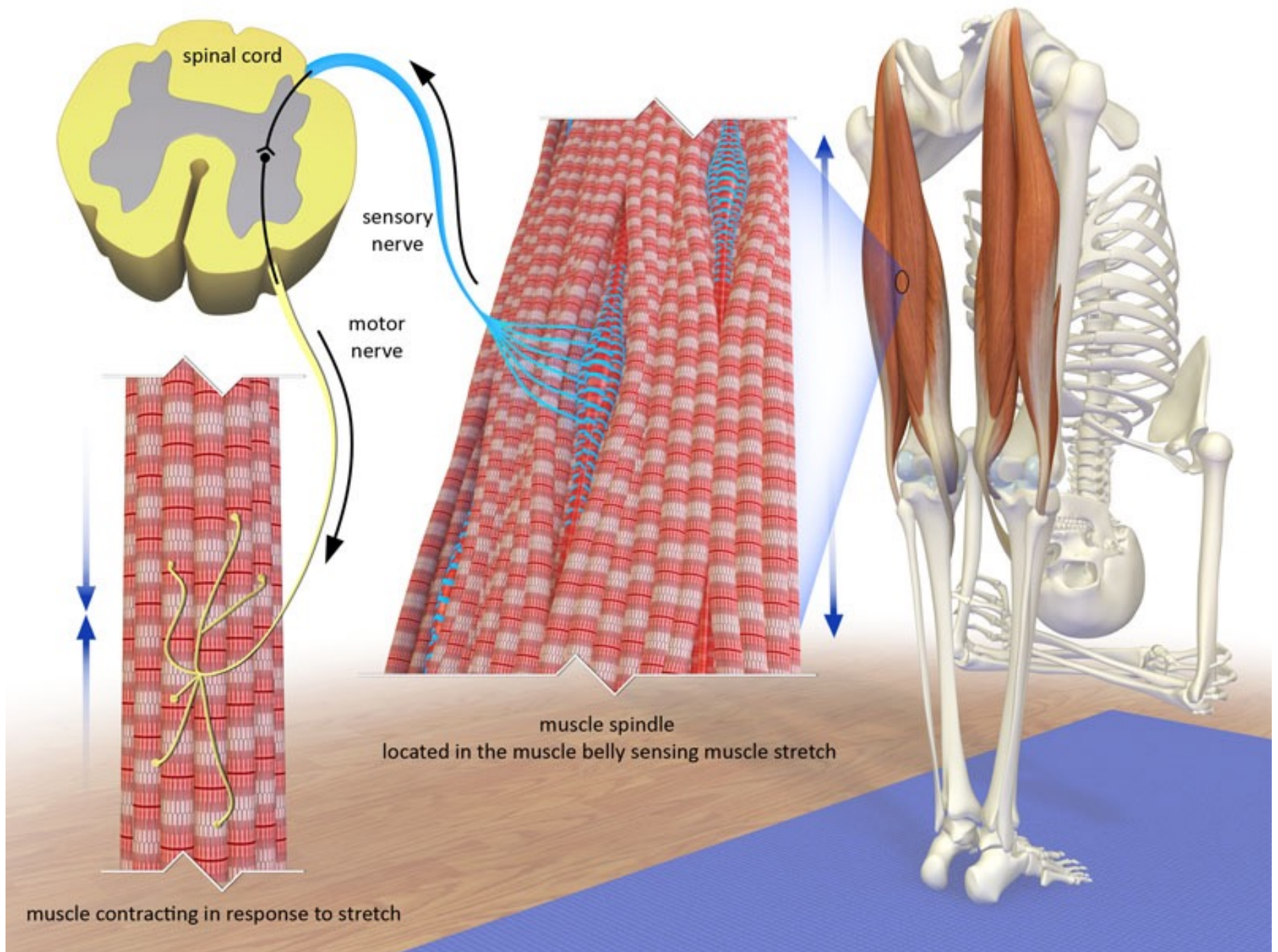
Synapse

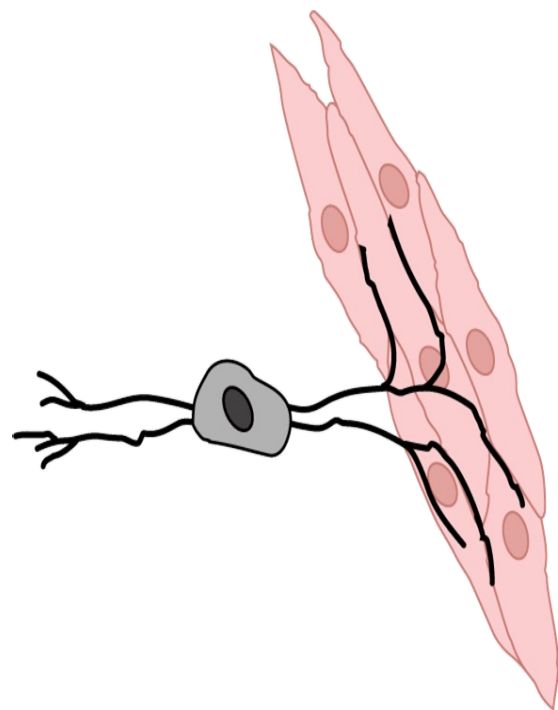
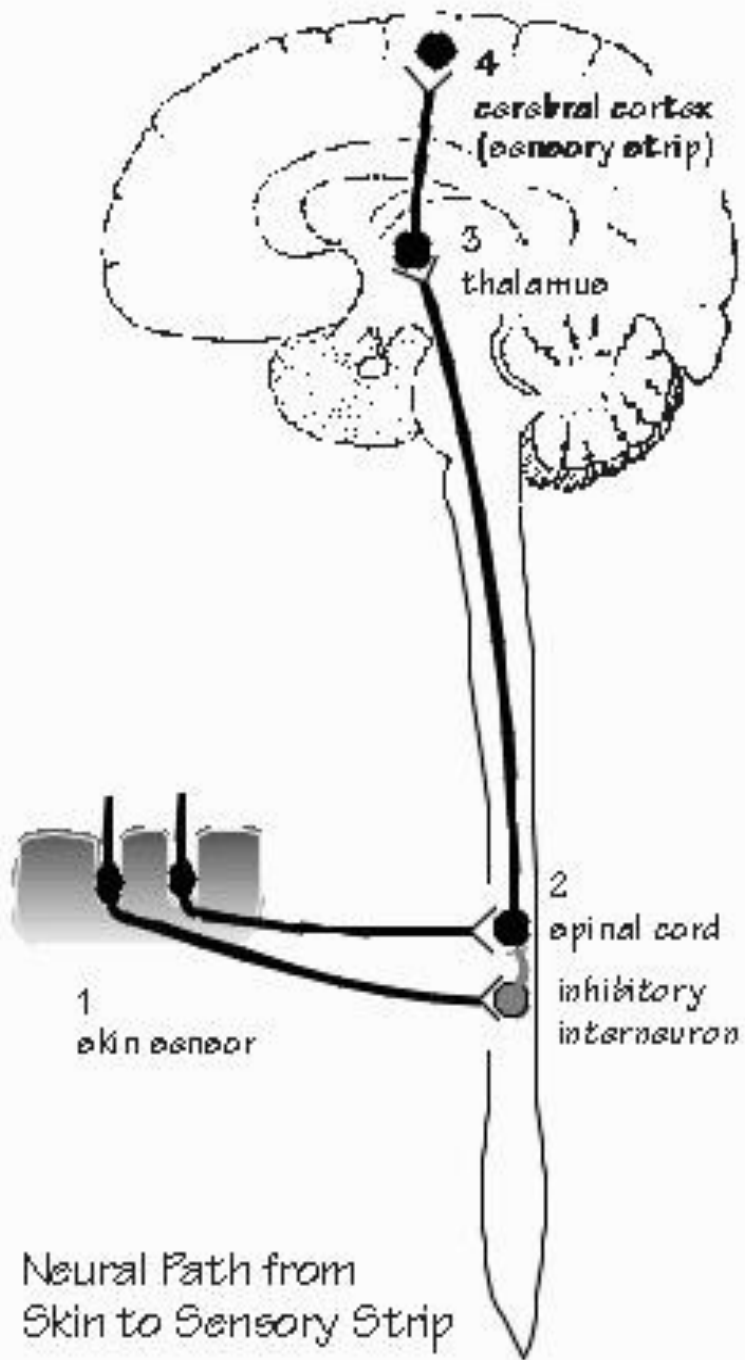


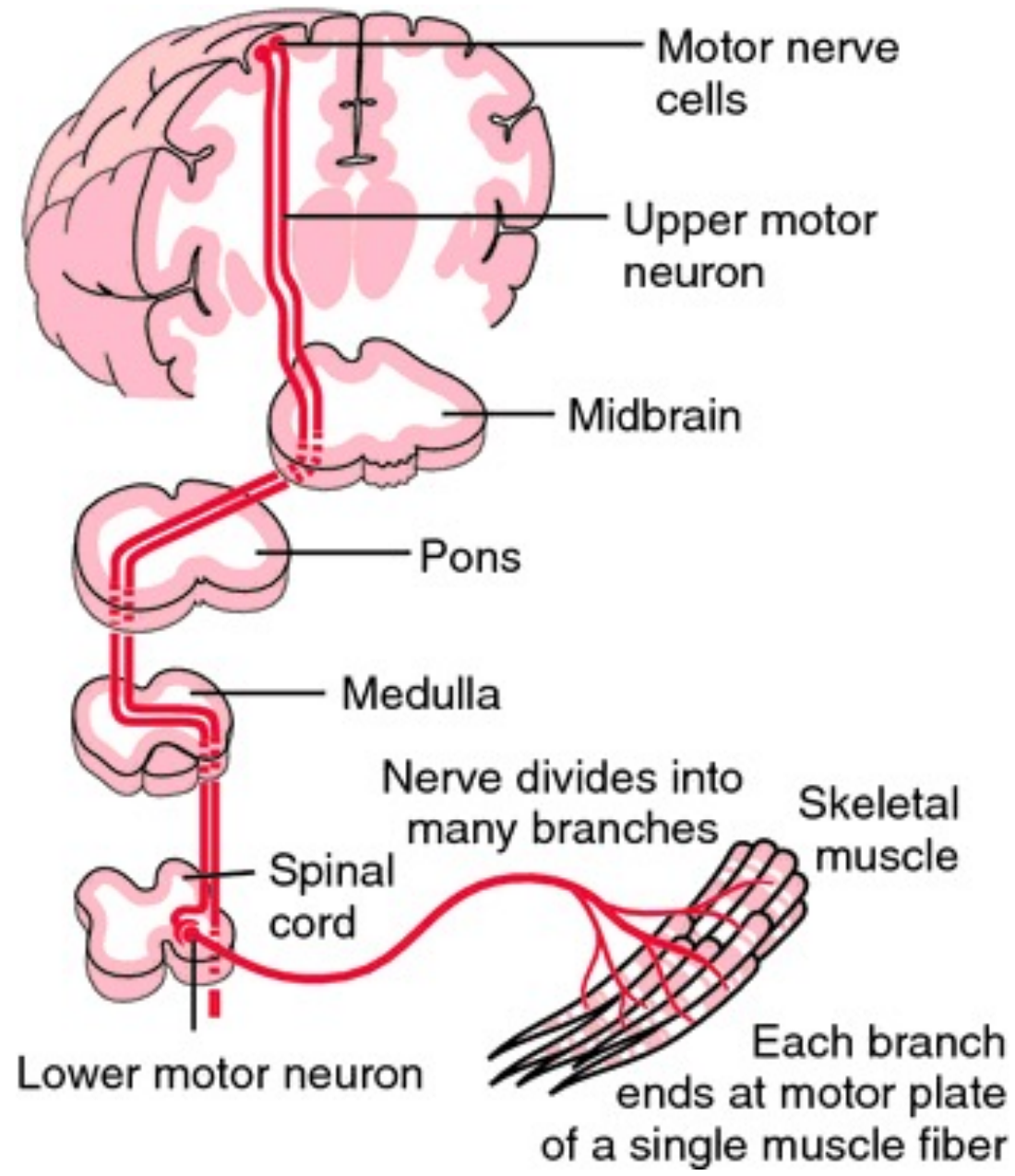
0 or 1

The Neuron as a Detector

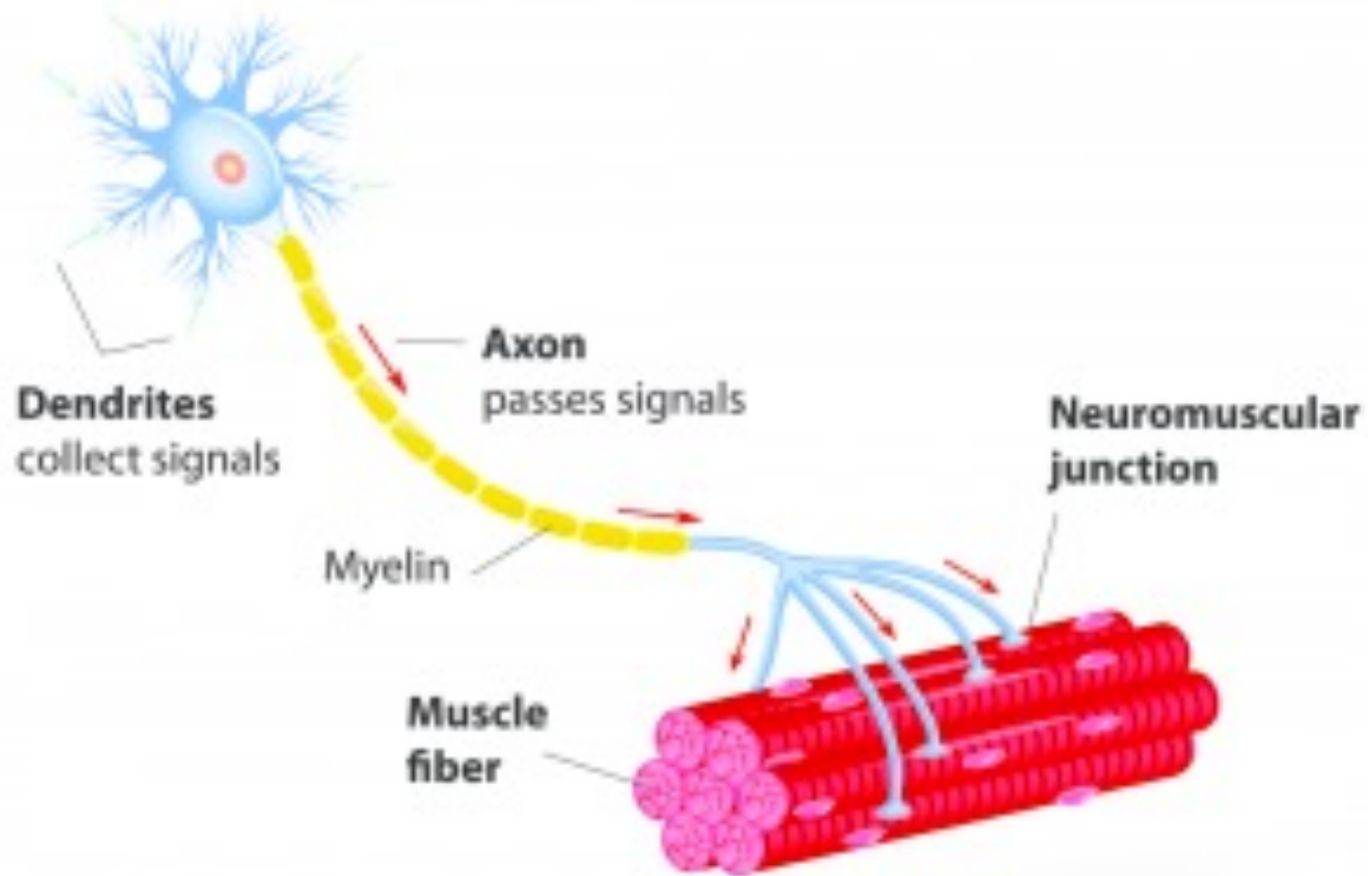






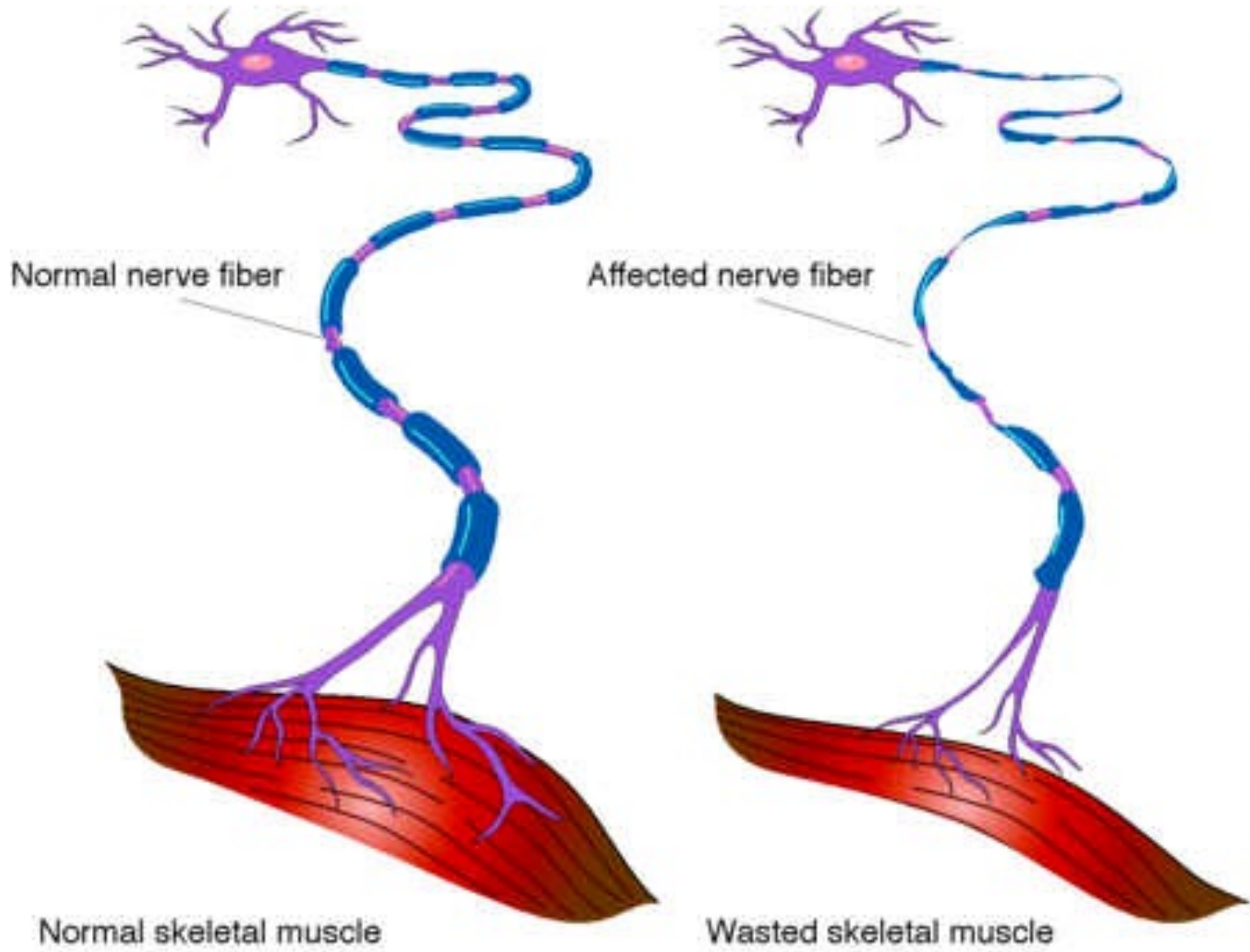


MOTOR NEURON



NORMAL SPINAL NEURON

DISEASED SPINAL NEURON



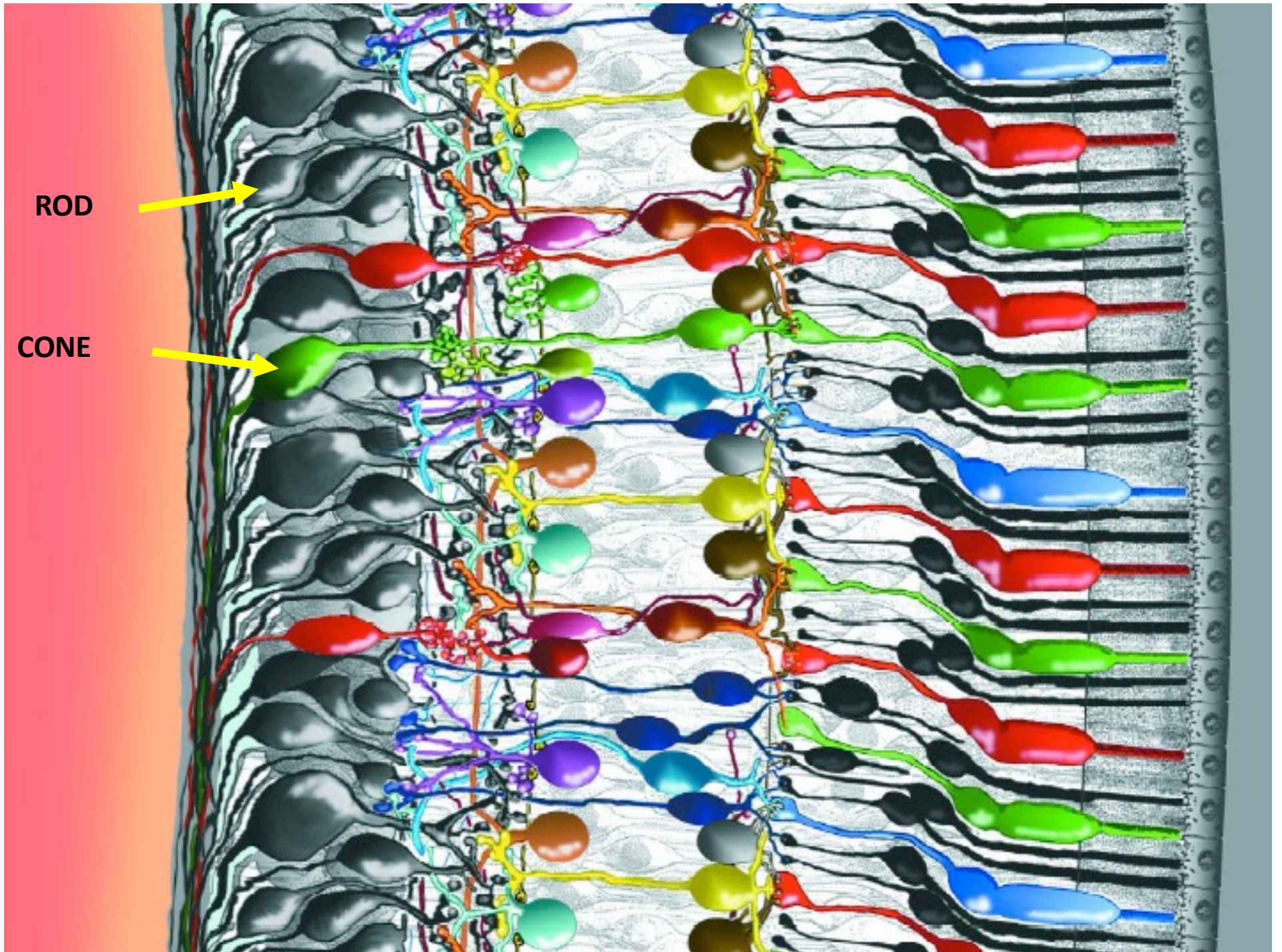
Normal nerve fiber

Affected nerve fiber

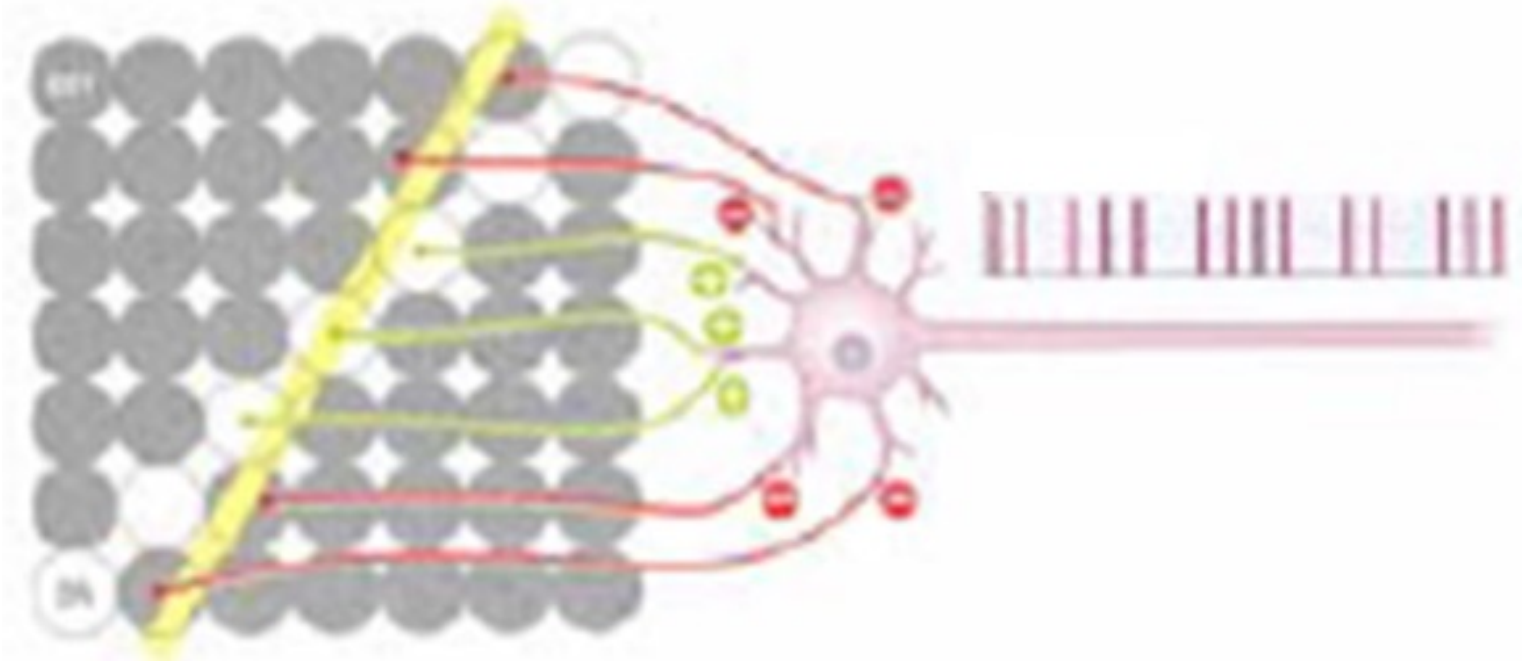
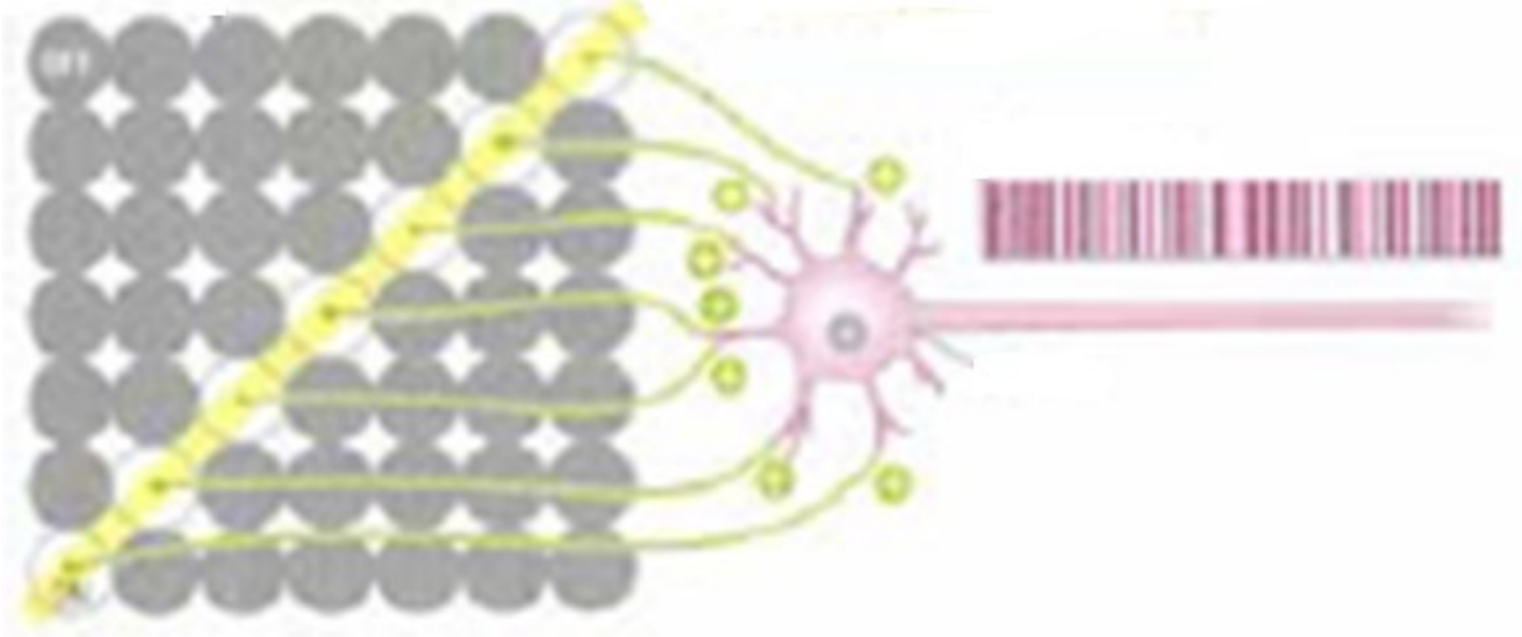
Normal skeletal muscle

Wasted skeletal muscle

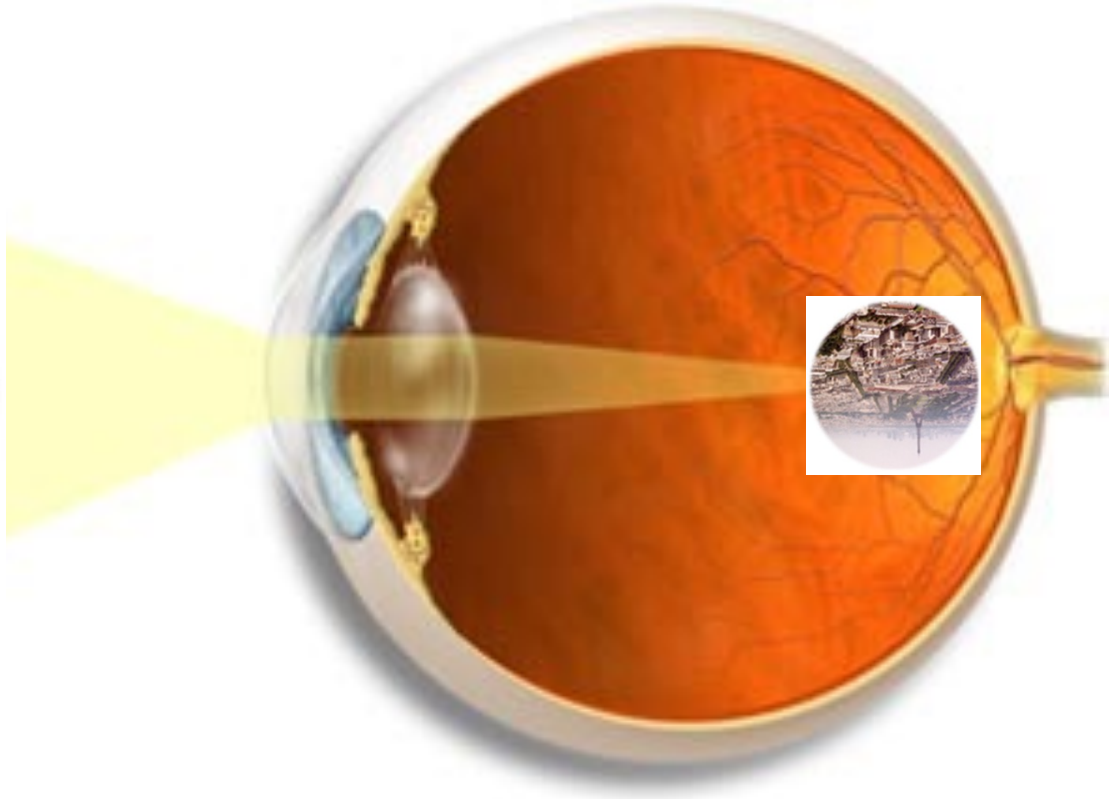
Other Types of Neurons

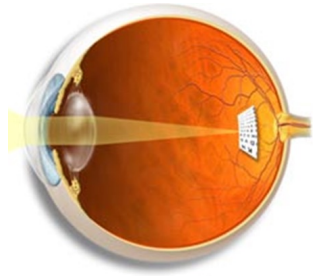


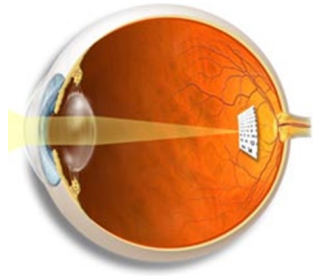


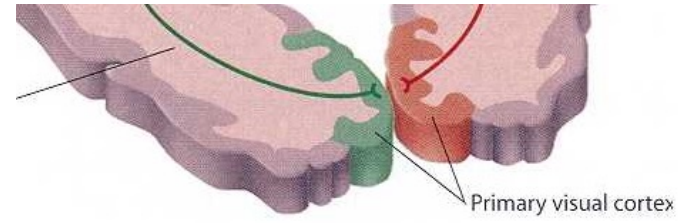
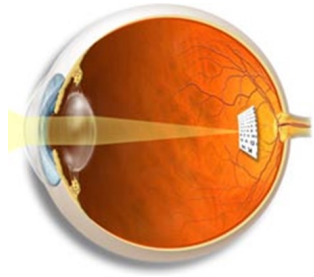


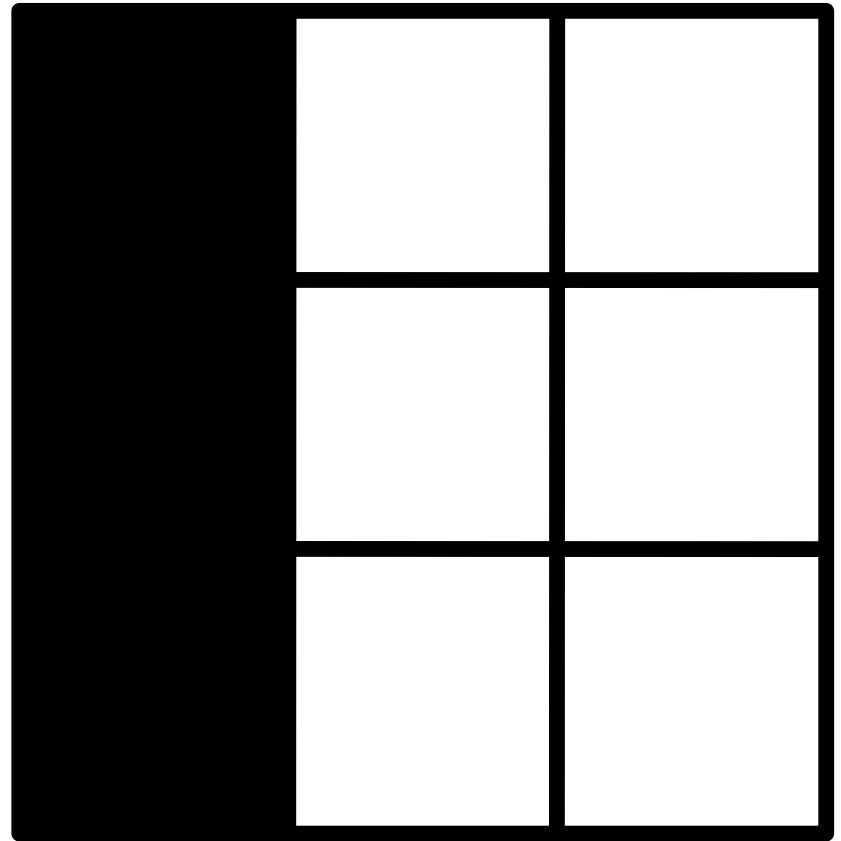
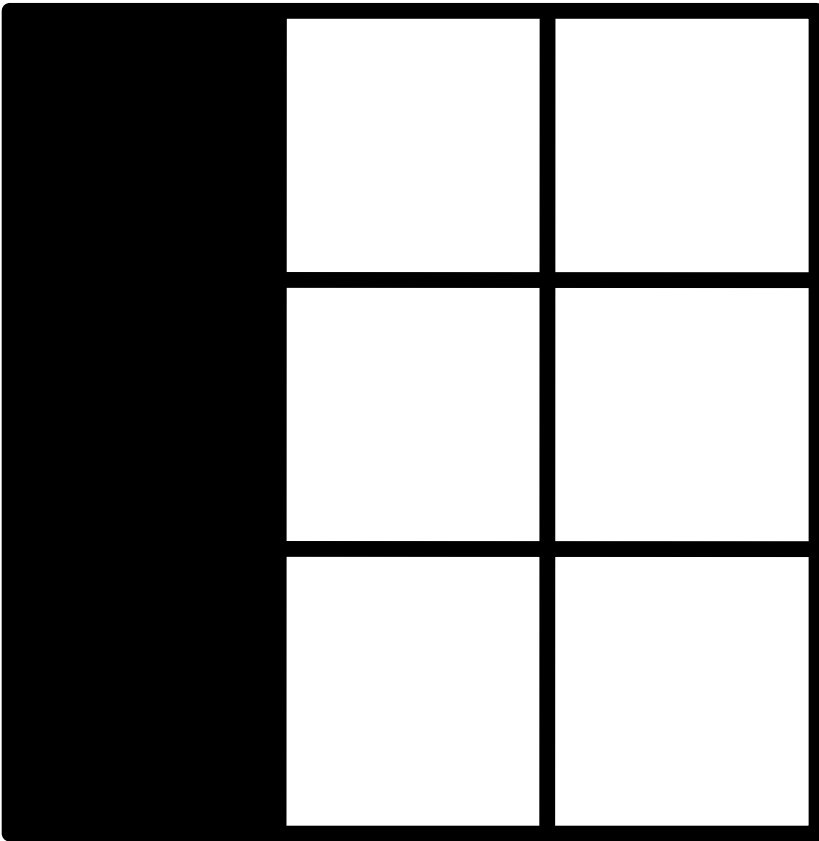
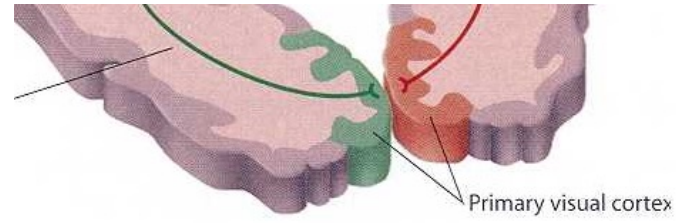
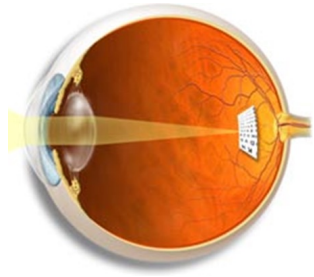
Layers of Detectors

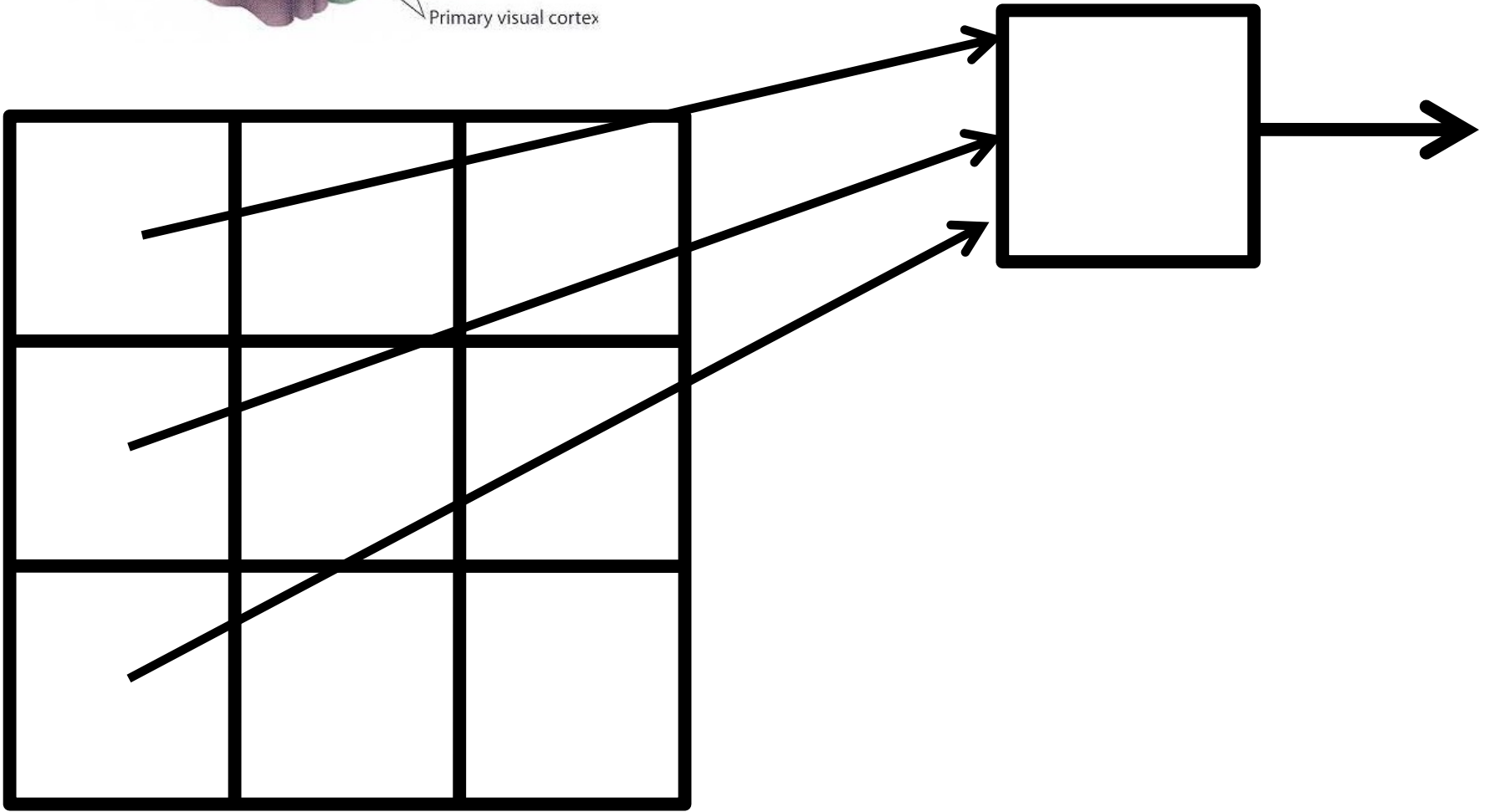
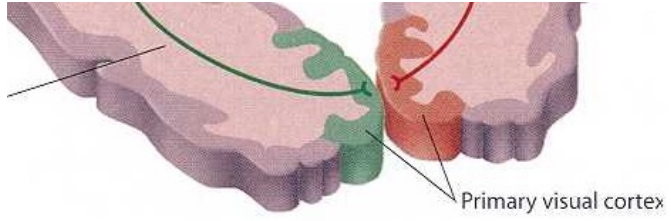


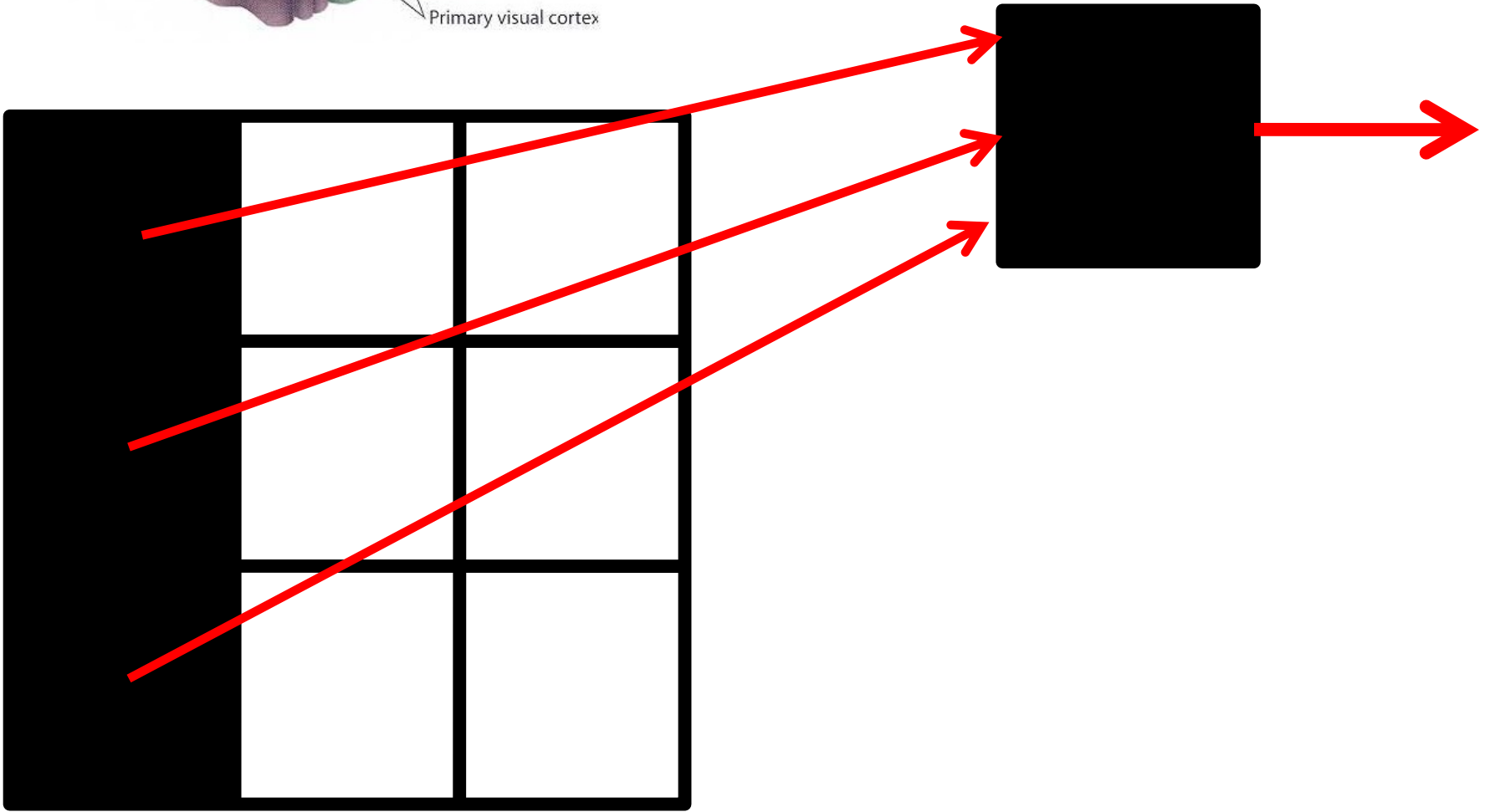
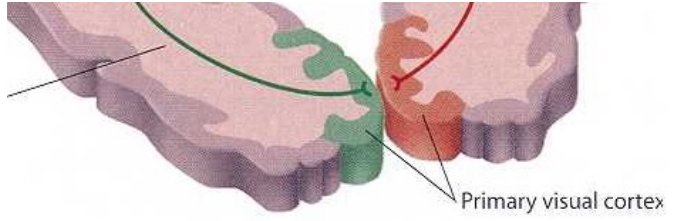


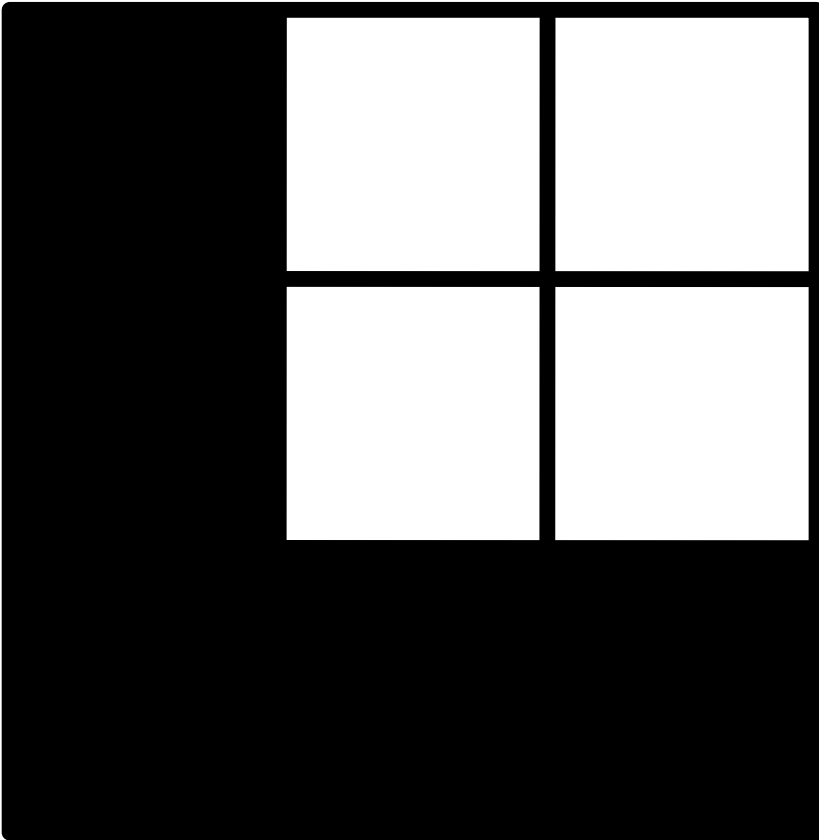
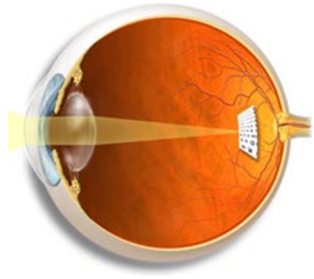


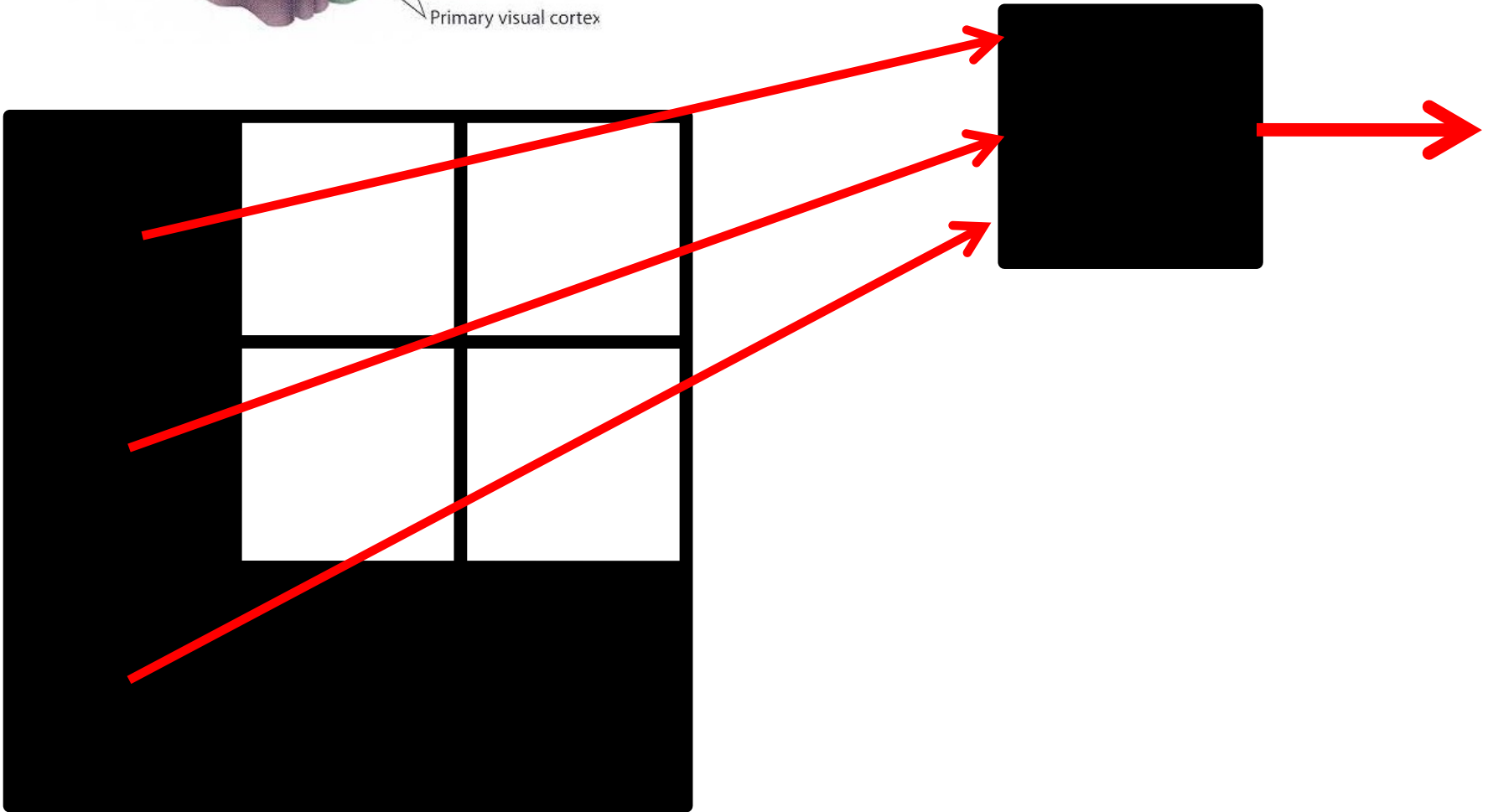
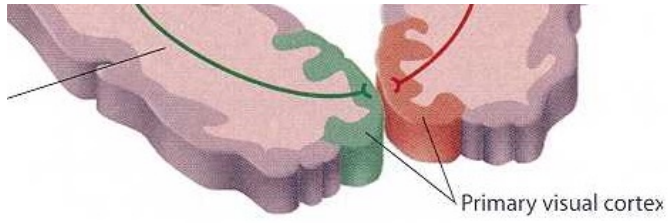


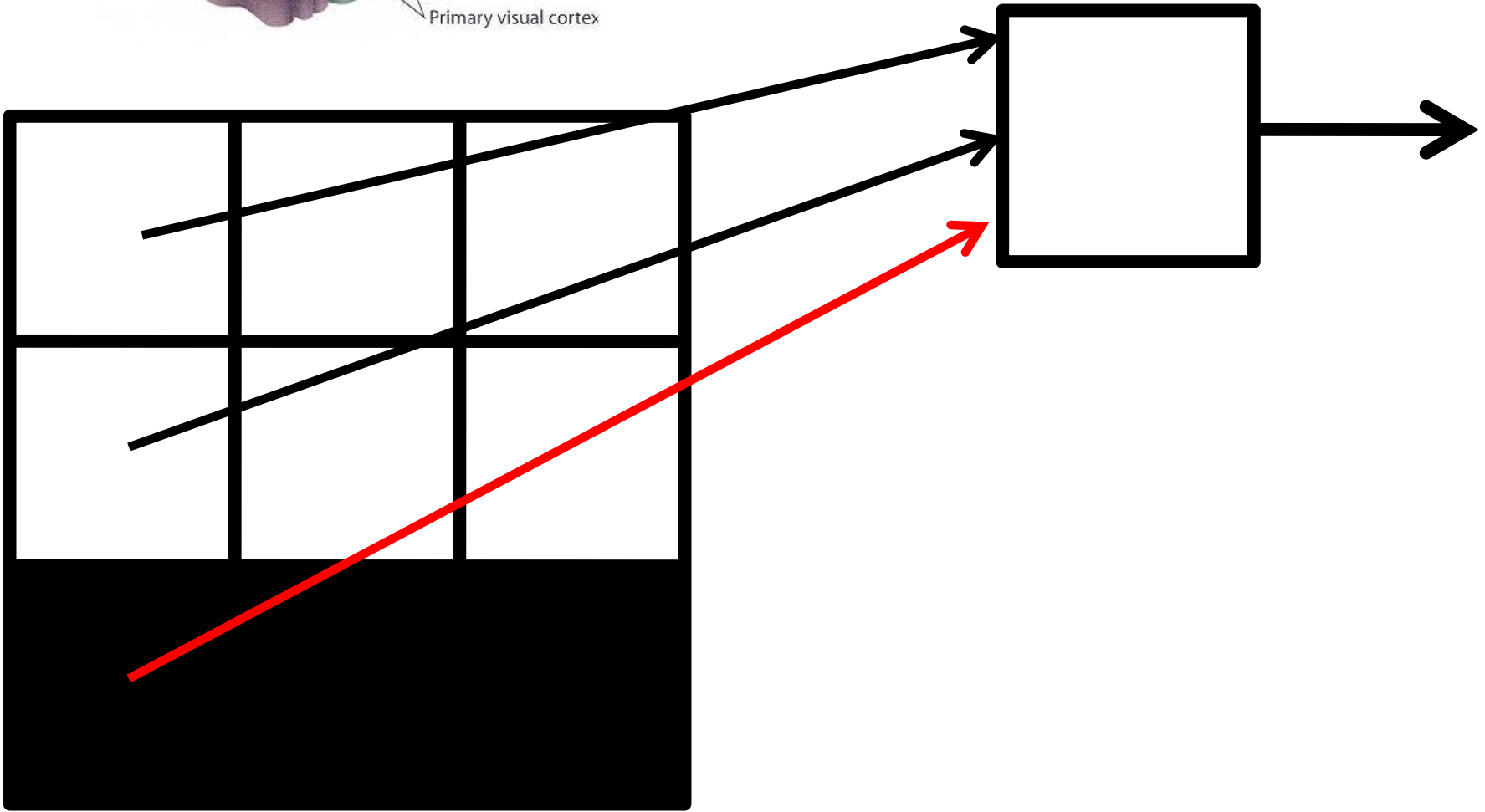
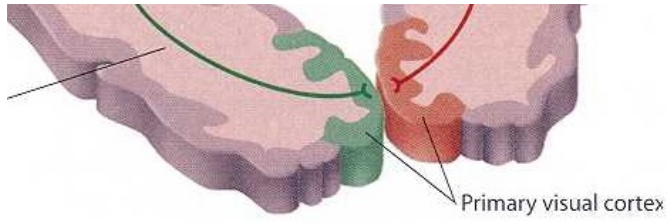


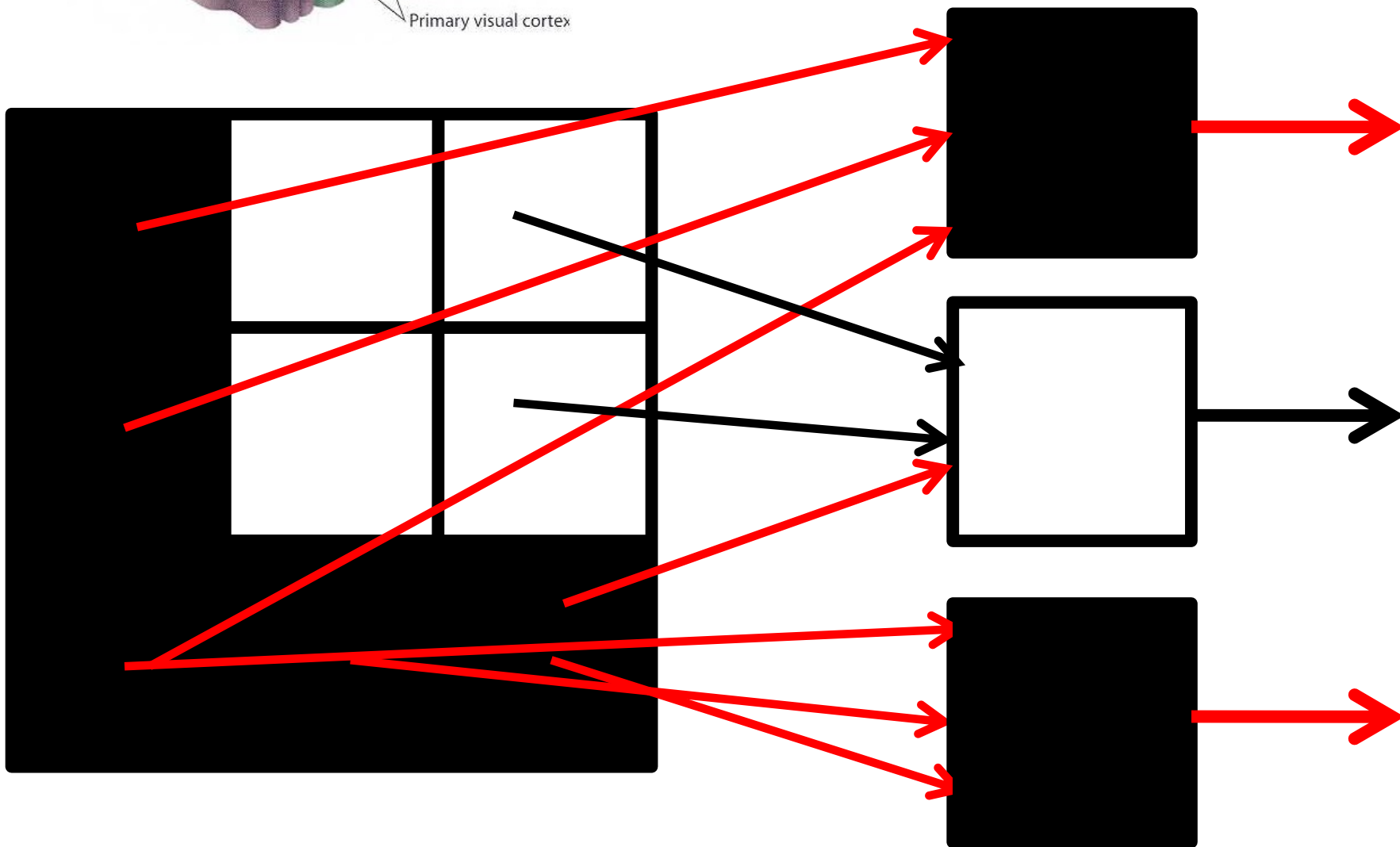
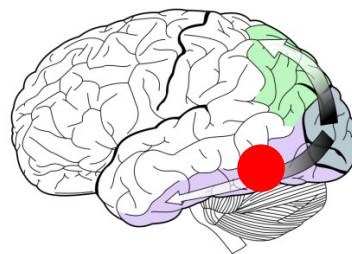
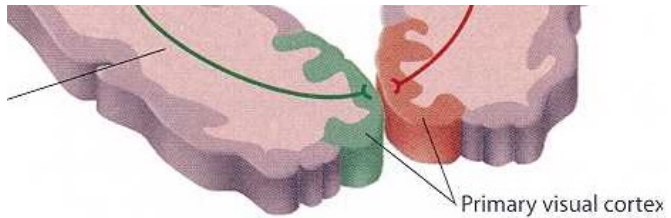


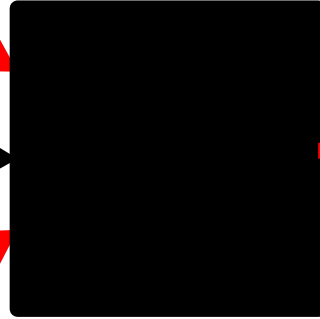
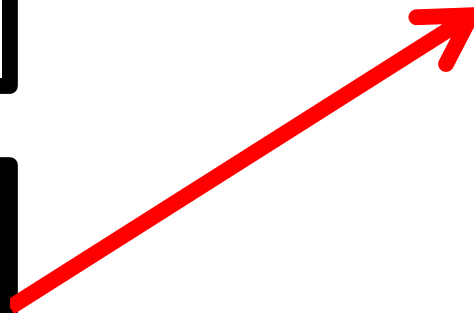
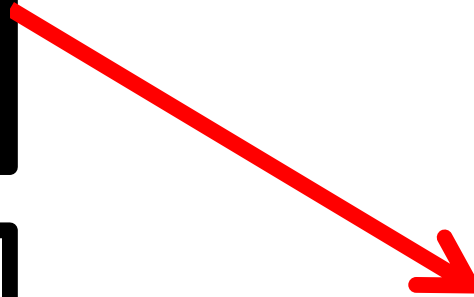
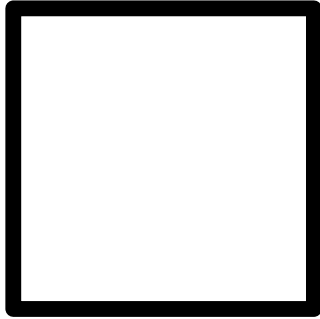
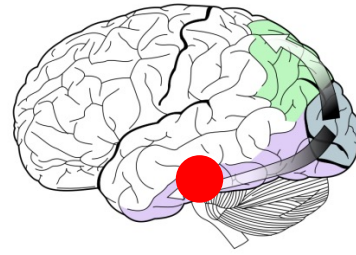
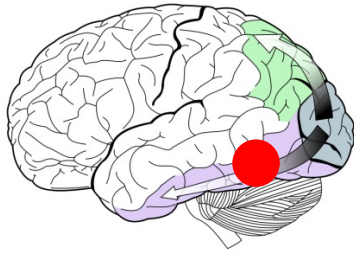




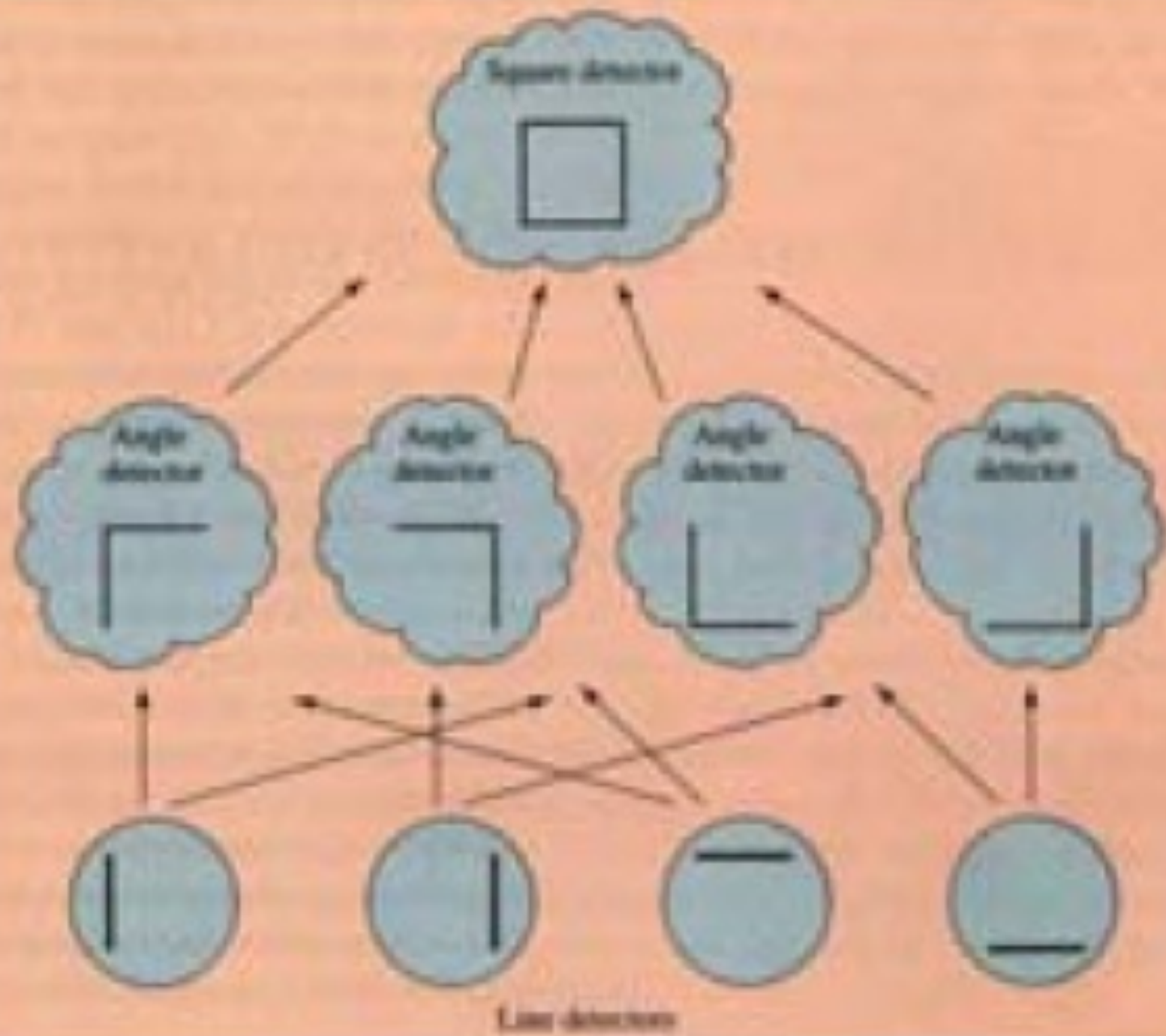


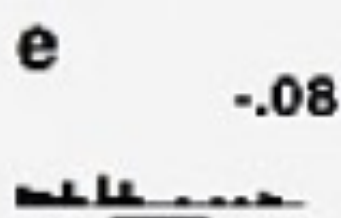
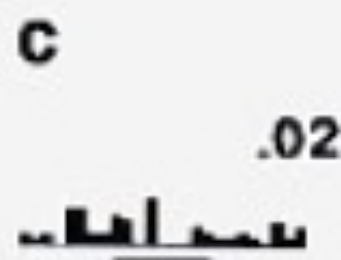
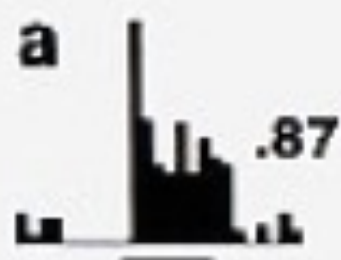




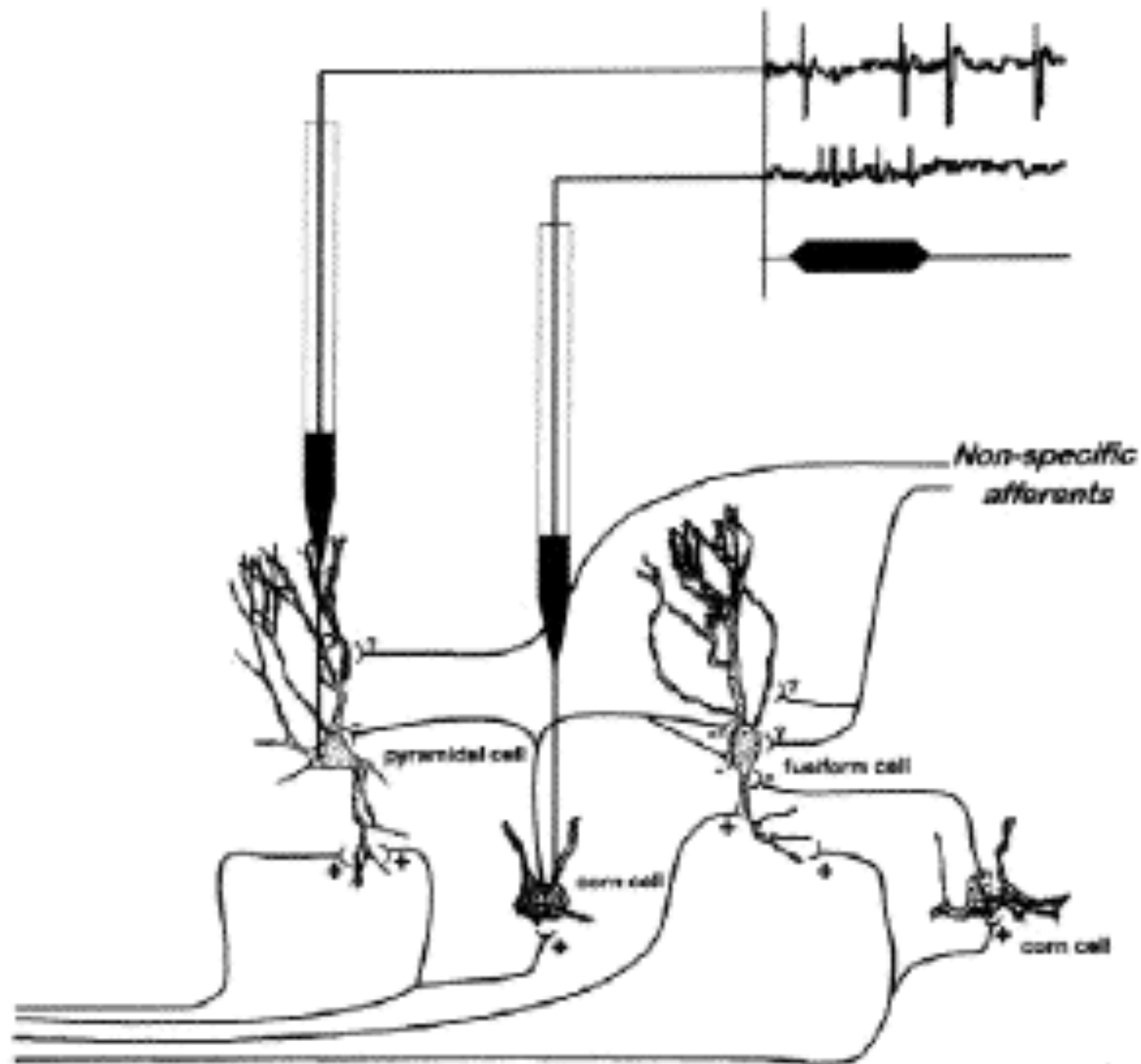


“L”



B

Lecture 1C: Techniques



Auditory Nerve (same CF)

