

Lecture 9, 06 Feb 2008

Vertebrate Physiology  
ECOL 437 (MCB/VetSci 437)  
Univ. of Arizona, spring 2008

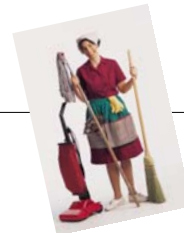
Kevin Bonine & Kevin Oh



1. [Synapses](#) (Ch12)
2. [Sensory Systems](#) (Ch13)

[http://eebweb.arizona.edu/eeb\\_course\\_websites.htm](http://eebweb.arizona.edu/eeb_course_websites.htm)<sup>1</sup>

Housekeeping, 06 February 2008



Upcoming Readings

today: [Ch13](#)

LAB Wed 06 Feb: [Catania 2002](#), [Barinaga 1999](#), [Malakoff 1999](#)

(see website for links to papers; "worksheet" via email)

Fri 08 Feb: [Ch13](#)

Mon 11 Feb: [Ch13](#)

Wed 13 Feb: [Ch13](#)

LAB Wed 13 Feb: none

Fri 15 Feb: [Exam 1](#), through [Ch13](#)

Lab discussion leaders: [20 Feb](#)

1pm – [Virsheena](#), [Mathew S. Arturo](#)

3pm – [Kat](#), [Clif](#), [Amber](#)

Lab discussion leaders: [06 Feb](#)

1pm – [Rittner](#), [Whitney](#)

3pm – [Roxanne](#), [Maria](#)

2

# PHYSIOLOGY & UA ADVANCE

**Christine Maric, Ph.D., FAHA, FASN**

Director, Diabetes Research  
Center for the study of Sex Differences  
Assistant Professor of Medicine  
Georgetown University Medical Center

Upcoming  
Physiology  
Seminar

**“Sex hormones in the  
pathophysiology of diabetic  
renal disease”**

**Friday February 8, 2008 11 a.m.**

Room 5403, Arizona Health Sciences Center

Also available on-line at:  
<http://www.physiology.arizona.edu/seminars>

(Refreshments served at 10:50 a.m.)

For additional information, please contact host: Heddwyn Brooks, 626-7702 [brooksh@emad.arizona.edu](mailto:brooksh@emad.arizona.edu)

\*This lecture is co-sponsored by the UA ADVANCE program,  
a program funded by the National Science Foundation under Grant  
No SBE-0548130, featuring young female scientists.\*

3

## The Edges of Life

Wednesday, February 6

Life's Final Edge? The Origin and Extinction of Species in a Human-Dominated Earth

Michael Rosenzweig, Professor, Ecology and Evolutionary Biology

Today, Earth's treasury of species, its biodiversity, faces an existential challenge and its outcome depends on man. Science now knows we've taken away enough land from nature to precipitate a mass extinction like the one that exterminated the dinosaurs 65 million years ago. Using reconciliation ecology, we can prevent this - and preserve life.

Wednesday, February 13

Life's Cognitive Edge: The Role of the Mind and What it Means to be Human

Anna Dornhaus, Assistant Professor, Ecology and Evolutionary Biology

Our human mind distinguishes us from other animal life-or does it? Recent research has revealed culture and social learning, tool use, complex communication, self-recognition, and planning for the future are not unique to the human experience. With these new findings, science is finally getting closer to understanding exactly what makes us human.

Wednesday, February 20

Life's Human Edge: Changing Perspectives on the End of Life

Michael Gill, Associate Professor, Philosophy

Nothing looms with more certainty than the final edge of one's own life. But in fact, the edge between life and death is anything but clear. This lecture will address the attempts that have been made to define the line between life and death and will explore the biological, legal, ethical, and spiritual debates that have raged around that line.

Wednesday, March 5

Life's Technological Edge: The Singularity is Near: When Humans Transcend Biology

Ray Kurzweil, via *Teleportec Teleporter*

Founder, Chairman and Chief Executive Officer, Kurzweil Technologies

Humanity is on the edge of a vast transformation, when what it means to be human will be both enriched and challenged. Inventor and futurist Ray Kurzweil will introduce this radically optimistic singularity, an era when we break our genetic shackles to create a nonbiological intelligence trillions of times more powerful than today. In this new world, humans will transcend biological limitations to achieve entirely new levels of progress and longevity.

*This lecture co-sponsored by: UA College of Engineering and UA College of Science*

**These do not count as physiology lectures.**

4

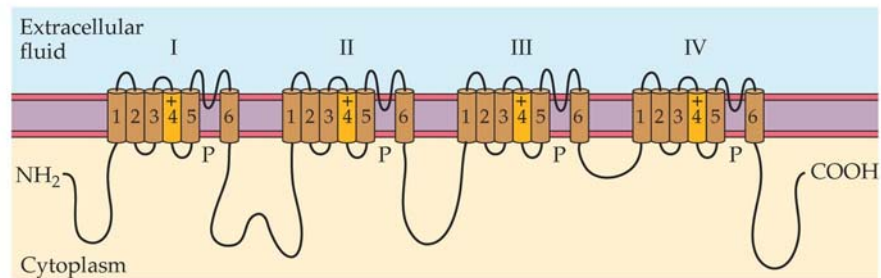
**TABLE 12.3 Ionotropic and metabotropic receptors: Structural, functional, and mechanistic differences**

Characteristic	Ionotropic receptors	Metabotropic receptors
Receptor molecule	Ligand-gated channel receptor	G protein-coupled receptor
Molecular structure	Five subunits around an ion channel	Protein with seven trans-membrane segments; no channel
Molecular action	Open ion channel	Activate G protein; metabolic cascade
Second messenger	No	Yes (usually)
Gating of ion channels	Direct	Indirect (or none)
Type of synaptic effect	Fast EPSP or IPSP	Slow PSPs; modulatory changes (in channel properties, cell metabolism, or gene expression)

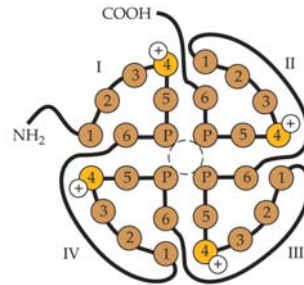
ANIMAL PHYSIOLOGY, Table 12.3 © Sinauer Associates, Inc.

(a) Topology of voltage-gated Na<sup>+</sup> channels

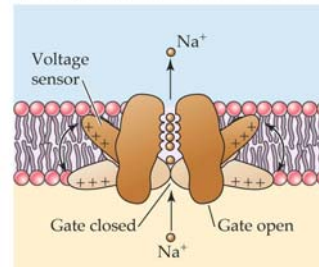
Hill et al. 2004, Fig. 11.17



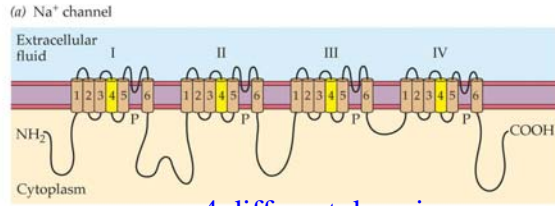
(b) Surface view of a Na<sup>+</sup> channel



(c) Voltage-dependent conformational change

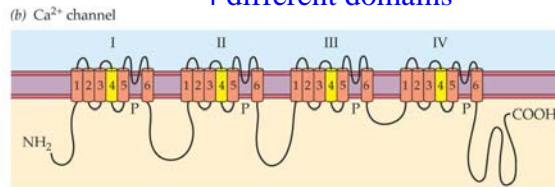


6

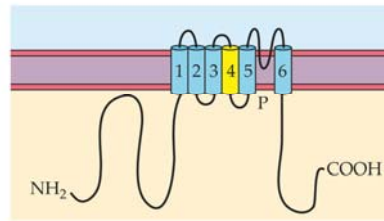


Hill et al. 2004, Fig. 11.18

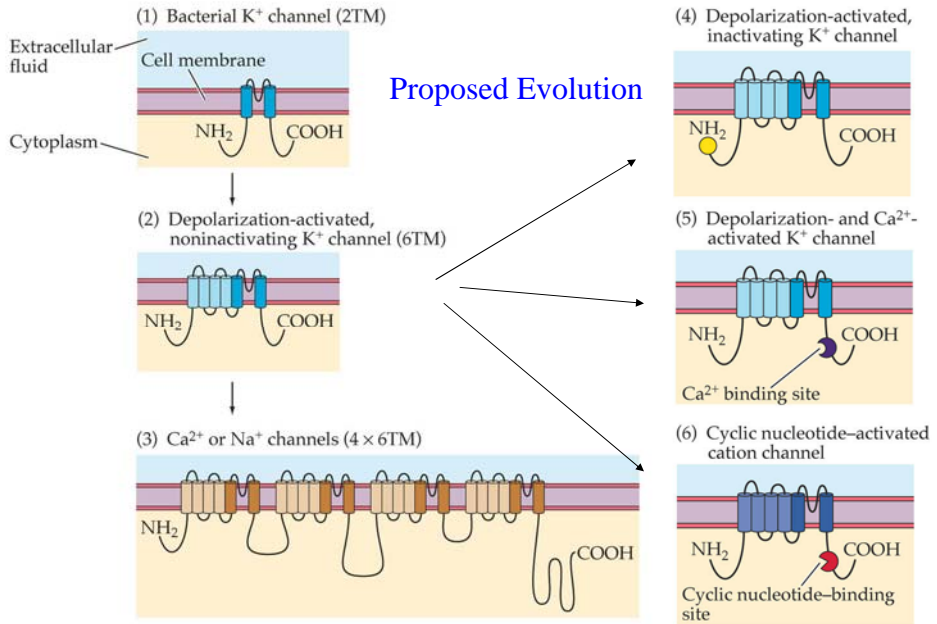
4 different domains



(c) K<sup>+</sup> channel 4 identical subunits



Voltage-gated channel superfamily



Voltage-gated Channels

ANIMAL PHYSIOLOGY, Box 11.1 (Part 1) © 201

Hill et al. 2004, pg. 301

# Neurotransmitters:

1. **small-molecule neurotransmitters**  
(often made in axon terminals; common)
2. **neuroactive peptides**  
(often made in soma and shipped down axon)

Nematodes use a lot of the same neurotransmitters.

9

**TABLE 12.2 Some neurotransmitters and receptors of vertebrate central nervous systems**  
These lists are not exhaustive; there are more transmitters, and more receptors for each transmitter.

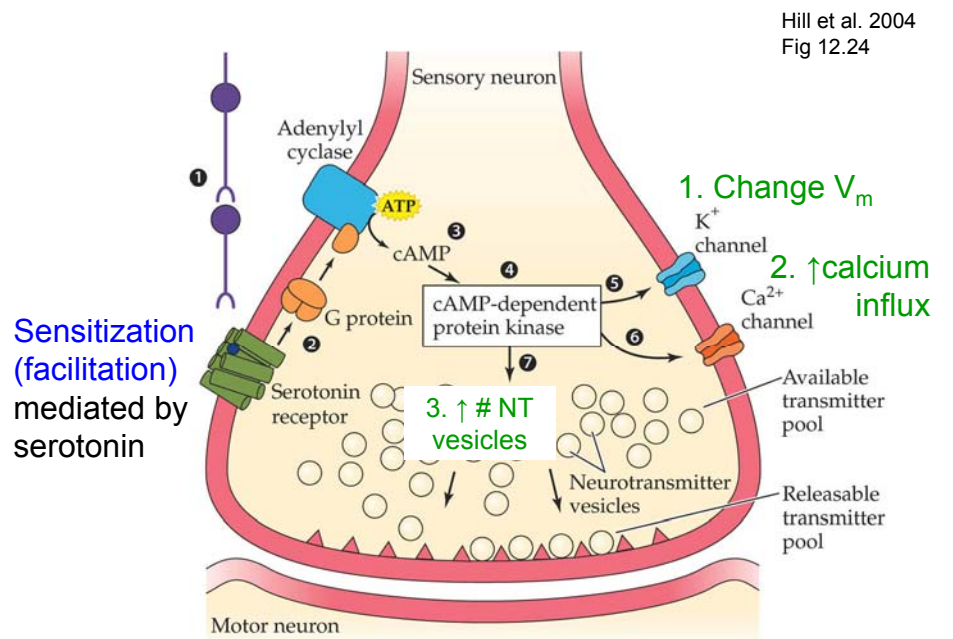
Neurotransmitter	Receptor class	Direct/ ionotropic	Indirect/ metabotropic	Common mode of action
<b>Amines</b>				
(10%) Acetylcholine	Nicotinic	X		EPSP
	Muscarinic M <sub>1</sub> -M <sub>5</sub>		X	G protein → IPSP
Dopamine	d <sub>1</sub>		X	
	d <sub>2</sub>		X	
	d <sub>3</sub>		X	
(1%) Norepinephrine	α <sub>1,2,3</sub>		X	
	β <sub>1,2</sub>		X	
Serotonin	5HT <sub>1</sub>		X	
	5HT <sub>2</sub>		X	
	5HT <sub>3</sub>	X		
<b>Amino acids (abundant and widespread)</b>				
<u>Glutamate</u>	AMPA	X		EPSP
	NMDA	X		Ca <sup>2+</sup> second messenger
	Metabotropic		X	DAG/IP <sub>3</sub>
GABA (IPSP)	GABA <sub>A</sub>	X		IPSP
	GABA <sub>B</sub>		X	G protein → IPSP
Glycine (IPSP)		X		IPSP
<b>Peptides</b>			X	G protein-coupled (some tyrosine kinase)

ANIMAL PHYSIOLOGY, Table 12.2 © Sinauer Associates, Inc.

# Synaptic Plasticity

- Change synaptic efficacy
- Alter rate of NT production and release
- Learning and Memory
- Facilitation vs. antifacilitation/depression
- Retrograde messengers (i.e., NO)
- Calcium-dependent
  - Research on-going

11



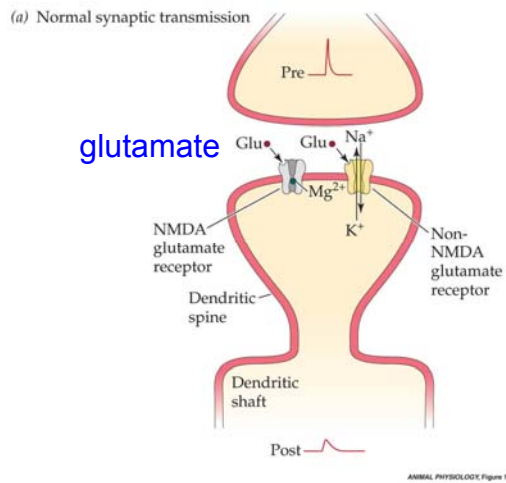
ANIMAL PHYSIOLOGY, Figure 12.24 © 2004 Sinauer Associates, Inc.

# Long-term Potentiation

- Often in Hippocampus
  - Site of Learning and Memory
- “Neurons that fire together wire together”
- NMDA glutamate receptors...

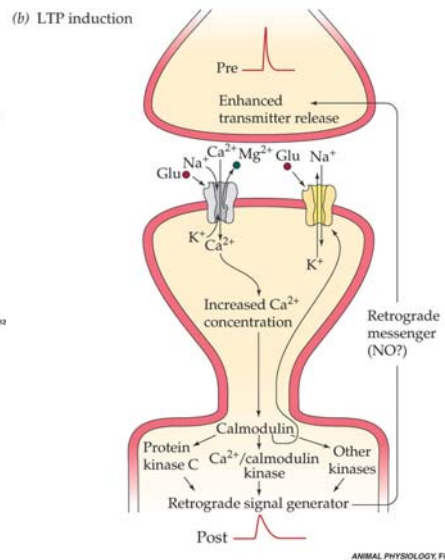
NMDA = N-methyl-D-aspartic acid

13



Hill et al. 2004, Fig 12.27

## Long term potentiation (LTP)



- NMDA glutamate receptors
  - role of Mg<sup>2+</sup>
  - voltage-dependent



Doogie Mice?



Genetic engineers upregulated production of **juvenile subunit of NMDA receptor** in adult mice (Doogie mice).

Ethical?

Should we do this in humans or other animals?

Under what conditions?

15

Vertebrate Physiology 437

Chapter 13

## Sensory Processes/Systems



16



## Sensing the Environment

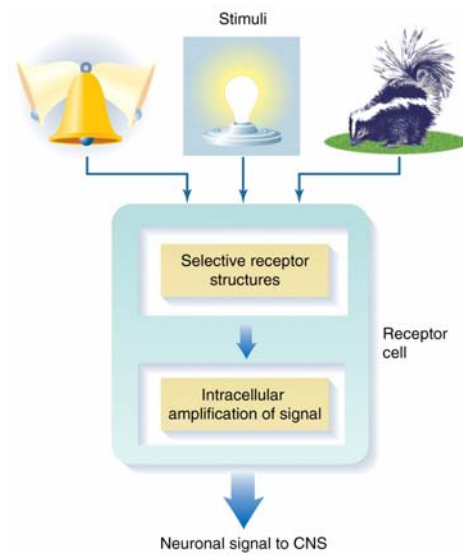
### Sensory Reception

- Environment
- Within body

### Integrated and Processed by NS

Sensory **Receptors** send signals to brain so perceive **sensations**

Sensory **Receptor** cells often organized into **organs**



7-1 Randall et al. 2002

## Properties of Receptor Cells

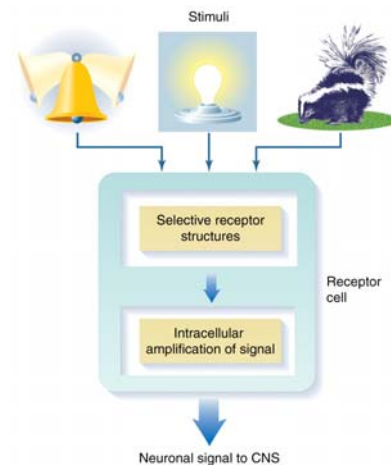
### Sensory **Modality**

Modalities include:

- vision, hearing, touch,
- taste, smell, **chemical,**
- thermal, proprioceptors**

**Qualities** within each modality

- e.g., Red or yellow;
- High or low-pitched

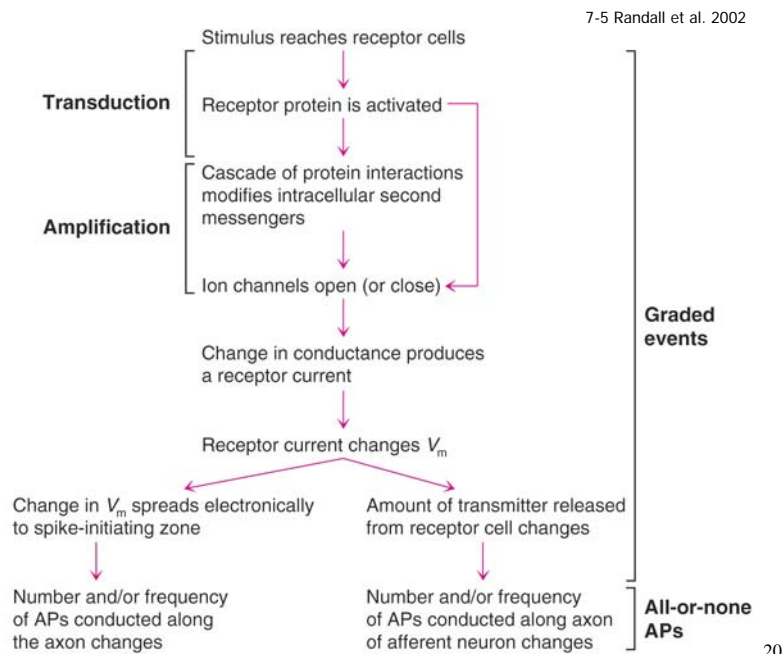


7-1 Randall et al. 2002

**TABLE 13.1** Classification of sensory receptors, based primarily on the kind of stimulus energy that excites them (Part 1)

Stimulus energy	Receptor modality	Stimulus perceived	Comments
Electromagnetic energy	Photoreceptors	Visible light Ultraviolet light Infrared radiation?	Some animals detect the polarity of visible and ultraviolet light Infrared detection may be thermal
	Electroreceptors	Electrical field or charge movement	
	Magnetoreceptors	Earth's magnetic field	Animals may sense polarity or angle of inclination of magnetic lines of force (see Chapter 16)
Thermal energy	Thermoreceptors	Hot Cold	"Infrared receptors" of rattlesnakes are actually sensitive heat receptors
Chemical energy	Chemoreceptors	Olfactory stimuli (distance chemoreceptors)	Chemical source is distant
		Taste (contact chemoreceptors) Internal chemoreceptors	Chemical source is nearby O <sub>2</sub> , CO <sub>2</sub> , H <sup>+</sup> , etc.
Mechanical energy	Mechanoreceptors	Touch, pressure	Skin or body surface
		Muscle length Muscle tension Joint position and movement	
		Sound (auditory stimuli) Balance and acceleration	Receptors detect gravitational or inertial forces
	Osmoreceptors	Osmotic pressure	Receptors detect mechanical stresses of osmotic swelling, etc.

19



## Mechanisms and Molecules

### Enzymatic Cascade to amplify

#### Threshold of Detection

e.g., 1 photon or hair cell  
movement of H diam.

Sour (pH; H+) and salt (Na+)  
move directly – no amplification

To measure quality need many receptors grouped  
into organ; different 'tunage' (e.g, wavelength of  
light or frequency of sound)

21

## Enhancing Sensitivity

### - Efferent Control

e.g., stretch receptors in muscle  
control length so can perceive stretch

### - Feedback Inhibition

Auto (helps keep in dynamic range)

vs.

Lateral...

22

## Properties of Receptor Cells

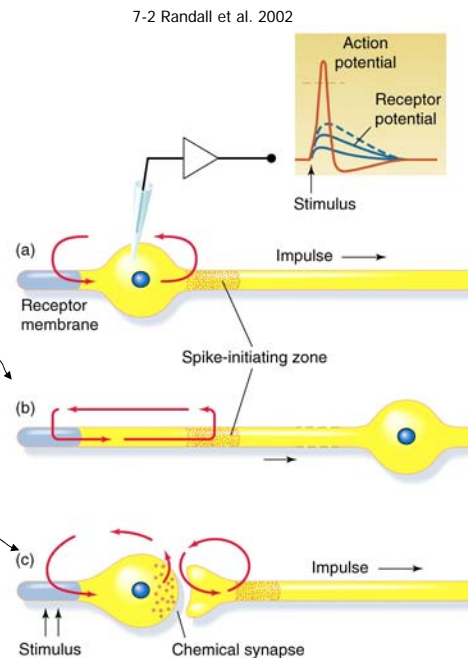
### Receptor Cells

- Specialized
- Selective for energy type and modality

-either is a neuron or  
-Synapses immediately on a neuron

(1° afferent neuron to CNS)

Stimulus modifies conformation of receptor



## Properties of Receptor Cells

### Transduction=

Stimulus energy converted to nerve impulse

### Example

Mechanoreceptors (touch)

- 1- Proteins respond to membrane distortion
- 2- Ion channels opened directly or indirectly
- 3- Current flows across membrane (often  $\text{Na}^+$ )

4- Vm changes (aka receptor potential changes)

5- Signal often amplified

6- AP sent or NT released causing AP

24

## Mechanisms and Molecules

### Sensory Adaptation

- orders of magnitude different stimulus strength
- often controlled via  $Ca^{++}$  availability
- local control or feedback from CNS

Type of stimulus received depends on *where* in CNS (~brain) AP arrives (Labeled lines).

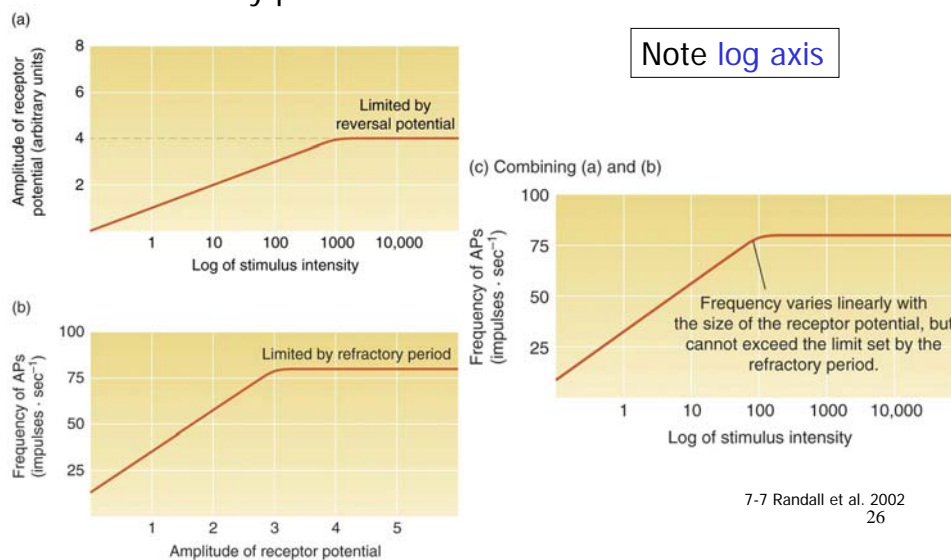
Rub eyes and see light!

Intensity signalled by frequency of APs, but...

25

### Stimulus Intensity and Dynamic Range

From lowest threshold, to upper limit imposed by refractory period:



## Dynamic Range

Shifting **range** of appropriate AP frequency

Detectable **light** intensity varies over **9 orders** magnitude

Detectable **sound** intensity varies over **12 orders** magnitude

## Range Fractionation

- Function of **sensory adaptation**
- Also recruit receptors with **different 'tunage'** or **sensitivity** (e.g., rods and cones in eye)

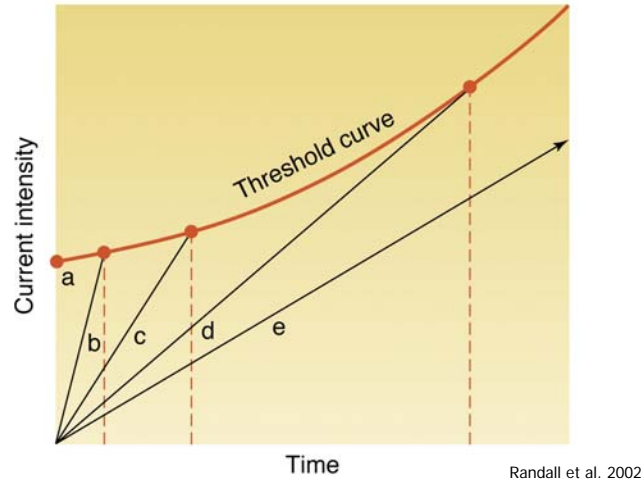
27

## Sensory Adaptation Possibilities:

1. Receptor cell mechanical **properties** may filter
2. Receptor cells may be **depleted** (e.g., visual pigments; need to be regenerated)
3. Enzyme cascade (during amplification) may be **inhibited** by (intermediate) product
4. **Electrical properties** change b/c  $\uparrow$   $[Ca^{++}]$
5. **Accommodation** of spike initiating zone
6. Sensory adaptation in **downstream neurons** (CNS)

28

## -Accommodation

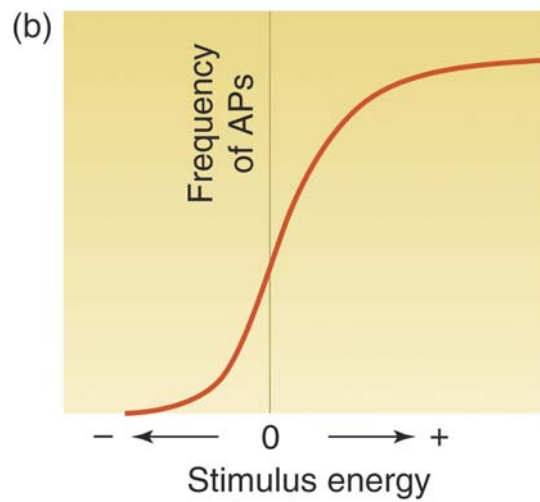


29

## Enhancing Sensitivity

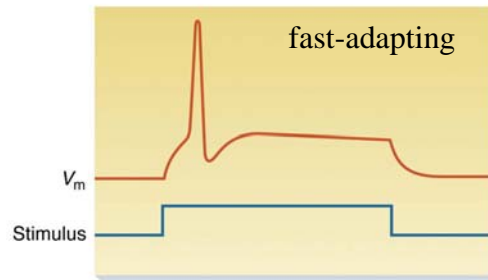
- Spontaneous basal activity
- Constant rate of APs
- Directionality if  
    ↑ or ↓ AP frequency

7-12 Randall et al. 2002

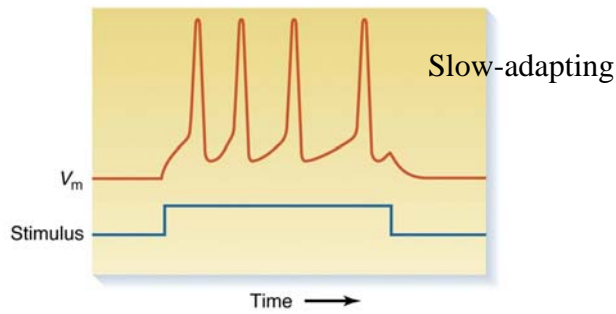


## Tonic vs. Phasic receptors

(a) Phasic response



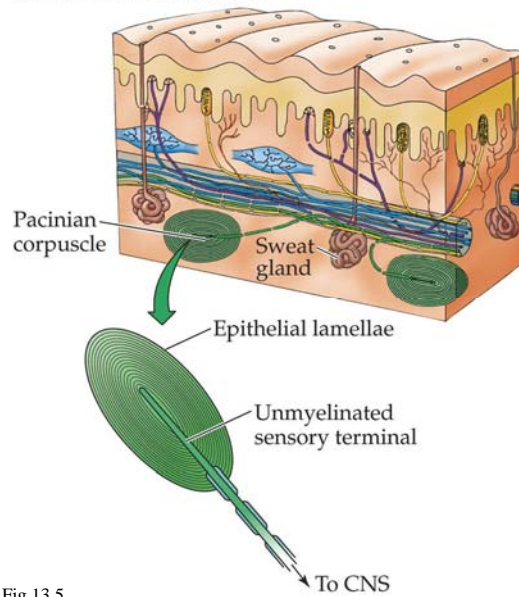
(b) Tonic response



5-19 Randall et al. 2002

31

(a) Location and structure

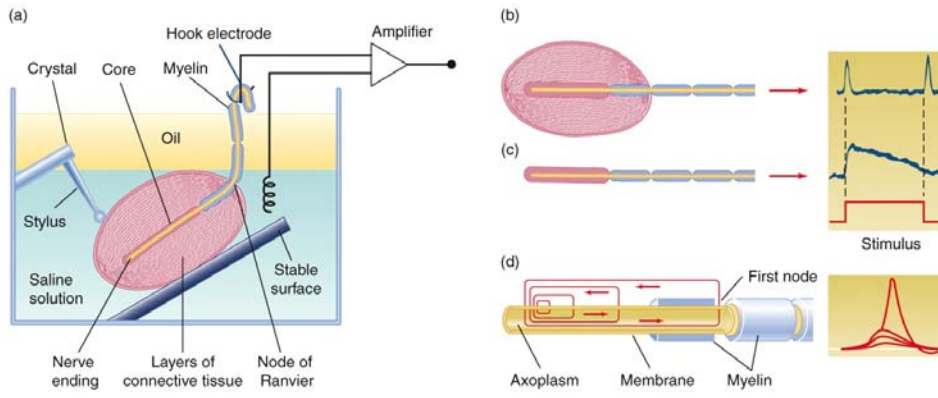


Hill et al. 2004, Fig 13.5

ANIMAL PHYSIOLOGY, Figure 13.5 (Part 1) © 2004 Sinauer Associates, Inc.

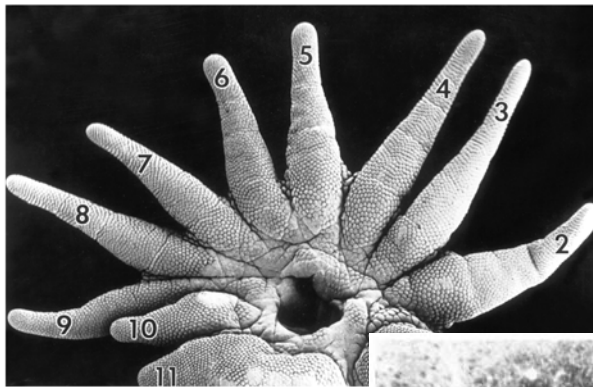


## Sensory Adaptation; Pacinian Corpuscle - Touch Example



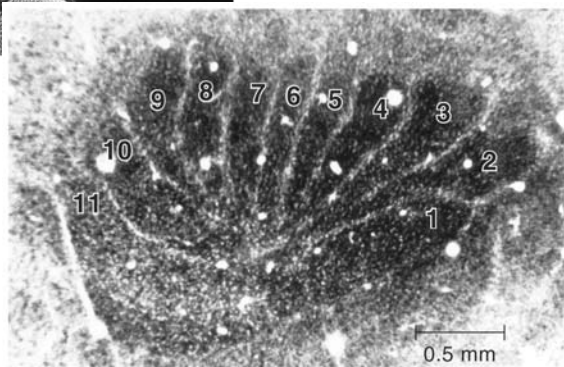
Movement of Oil between layers is what triggers APs  
Signal changes in pressure, not steady pressure

33  
7-10 Randall et al. 2002



### Star-Nosed Mole

(↑ # neurons, ↑ subtlety)  
(receptor field size?)



## External Chemoreception (Taste and Smell)

### -Taste

~ direct contact

### -Smell

~ distant signal source



-Chemoreception very sensitive

-*Bombyx* moth antenna example:

Male responds to female pheromone at low [ ] of  
1 molecule in  $10^{17}$  !

35

## Taste Chemoreception

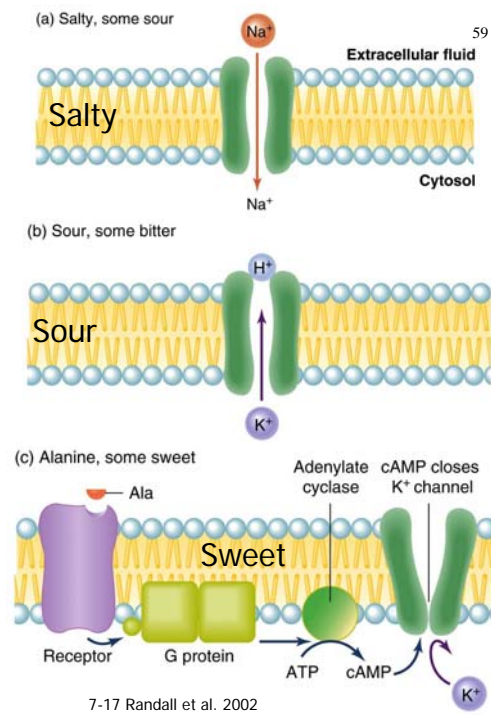
### -Taste

Usually oral cavity  
Some fish fins!

4-5 qualities:

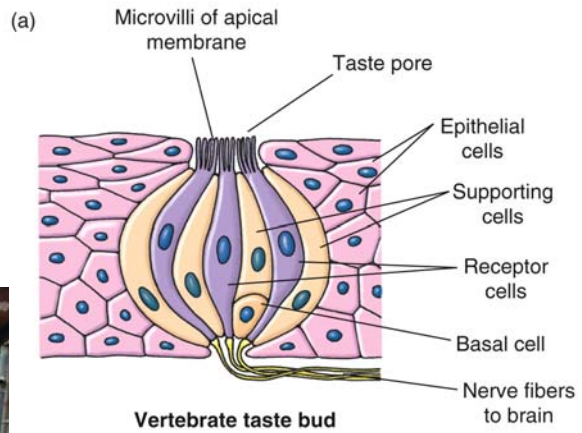
1. Salt
2. Sour
3. Sweet
4. Bitter
5. Umami  
("savory" or "meaty")

Differing Receptor  
Properties



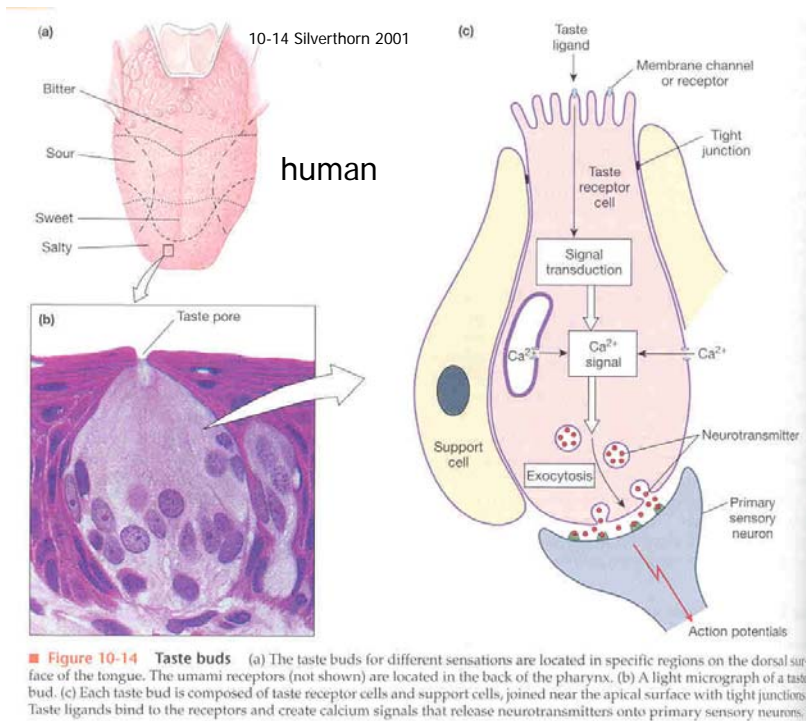
# Taste

- microvilli
- basal cells give rise to new receptor cells every 10 days

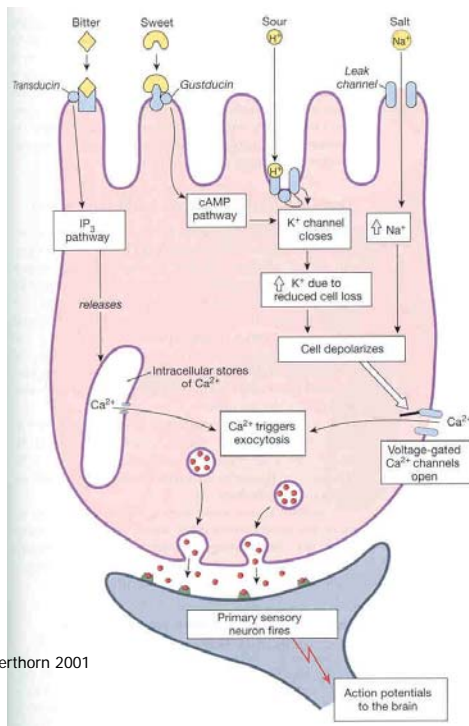


7-16 Randall et al. 2002

37



38



10-15 Silverthorn 2001



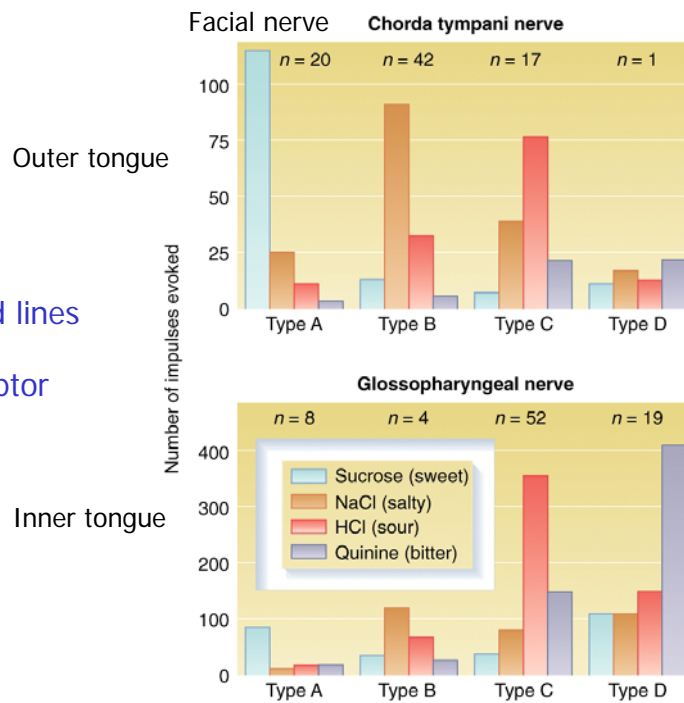
**Figure 10-15 Taste transduction**  
 Bitter and sweet ligand signal transduction uses G protein-coupled membrane receptors. Transducin releases  $Ca^{2+}$  from intracellular stores. Gustducin activates a cAMP second messenger pathway that closes  $K^+$  channels. Ionic ligands alter ion channels and depolarize the taste receptor, which allows  $Ca^{2+}$  entry from the extracellular fluid. For all taste ligands, the  $Ca^{2+}$  signal triggers exocytosis of neurotransmitter.

39

## Taste

-Quasi Labelled lines

- multiple receptor types/neuron



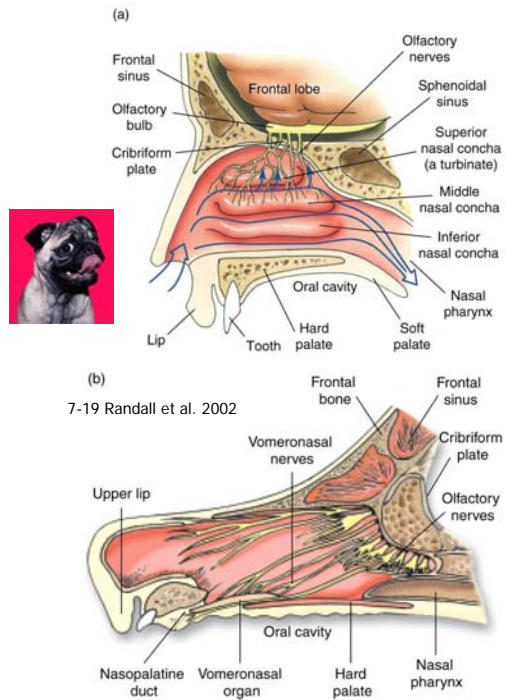
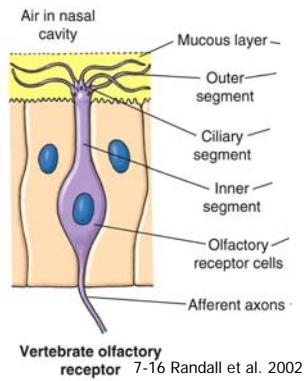
# Smell



41

## Smell/ Olfaction

- 1 **Nasal Cavity**  
-turbinates (↑s.a.)
- 2 **Vomeronasal organ**  
-usually **conspecific** communication



Smell/ Olfaction

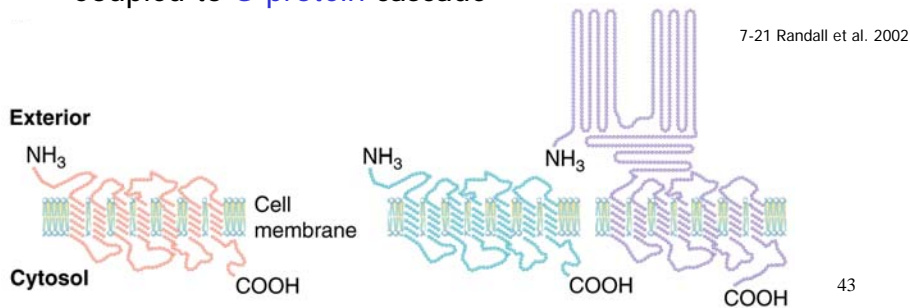
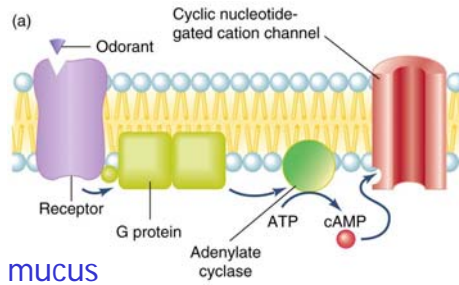
-Nasal and Vomeronasal:

-Epithelial tissue origin

-Cilia or Microvilli covered in mucus

-Receptor proteins with 7-transmembrane helices

-Coupled to G-protein cascade



Smell/ Olfaction

- Nasal and Vomeronasal:

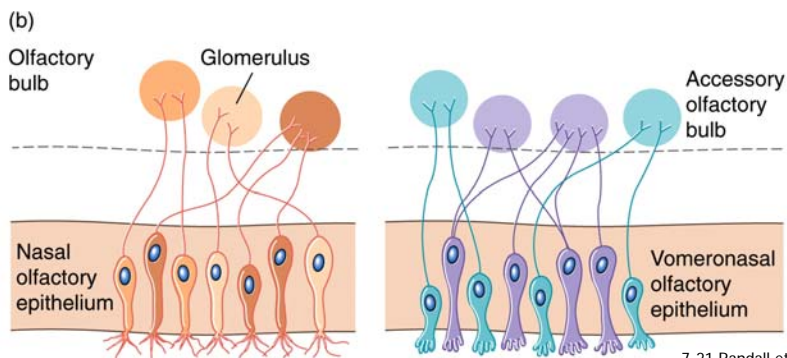


-Thousands of receptor proteins (general & special)

-but different for nasal and vomeronasal

-Receptor cells contain axons

- Glomeruli in olfactory bulb/accessory olfactory bulb



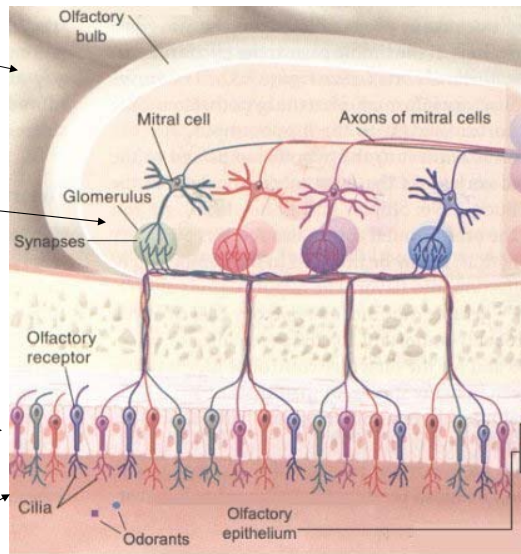
7-21 Randall et al. 2002

Olfactory bulb (info processing in brain)

Glomeruli (similar odor receptor synapses)

Sensory neurons (~ odor specific)

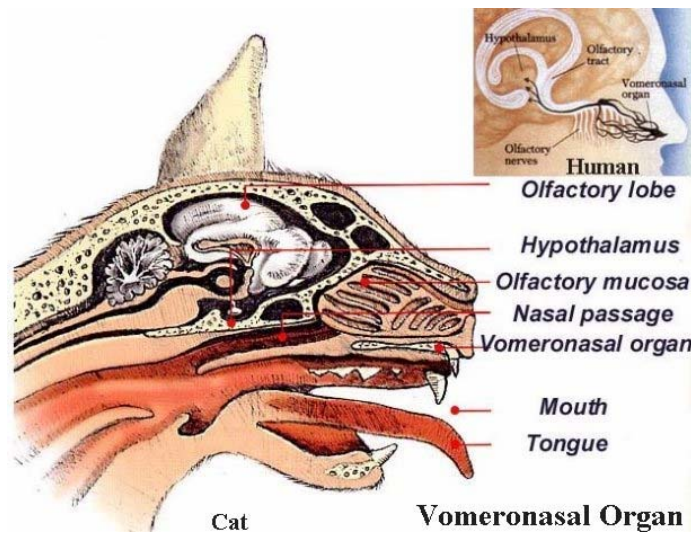
Mucus from epithelial glands



axons

dendrites

45



46

## Olfactory Neurons

In humans,  $10^7$  olfactory receptor neurons

In dogs,  $2 \times 10^8$

Human auditory nerve:  $10^4$

Human optic nerve:  $10^5$

47

end

48