

# Sensory Systems and Perception: Vision

Week 5

Martin Schrimpf

Slide credit: structure and content adapted from  
EPFL BIO-311 (P. Ramdya) and MIT 9.017 (R. Desimone)

# Learning Objectives – Week 5

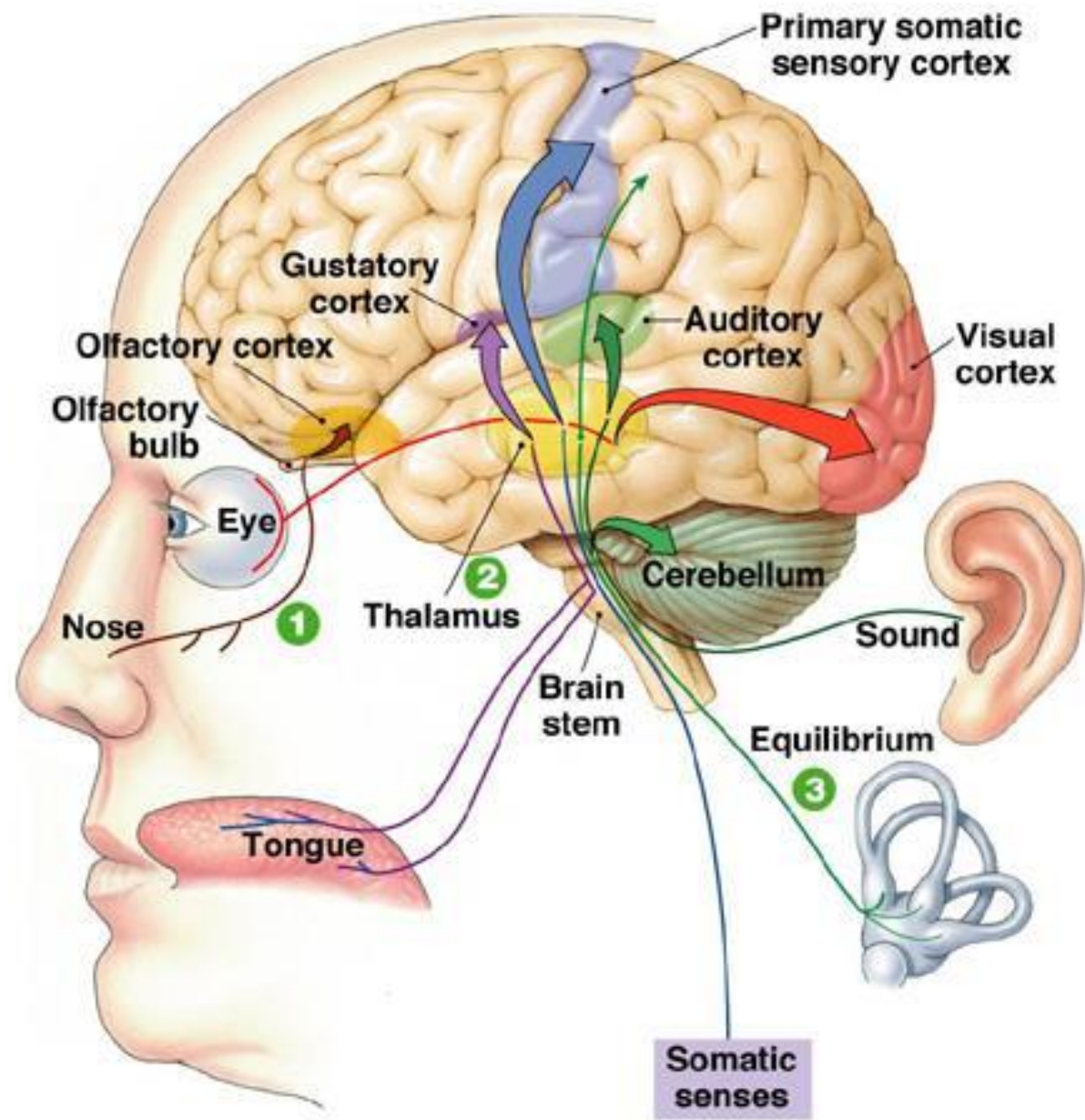
Describe the key components involved in vision.

Know what V1 responds to and the Hubel & Wiesel experiments.

Explain receptive fields.

Understand how a Gabor model relates to early vision.

Explain the visual hierarchy.

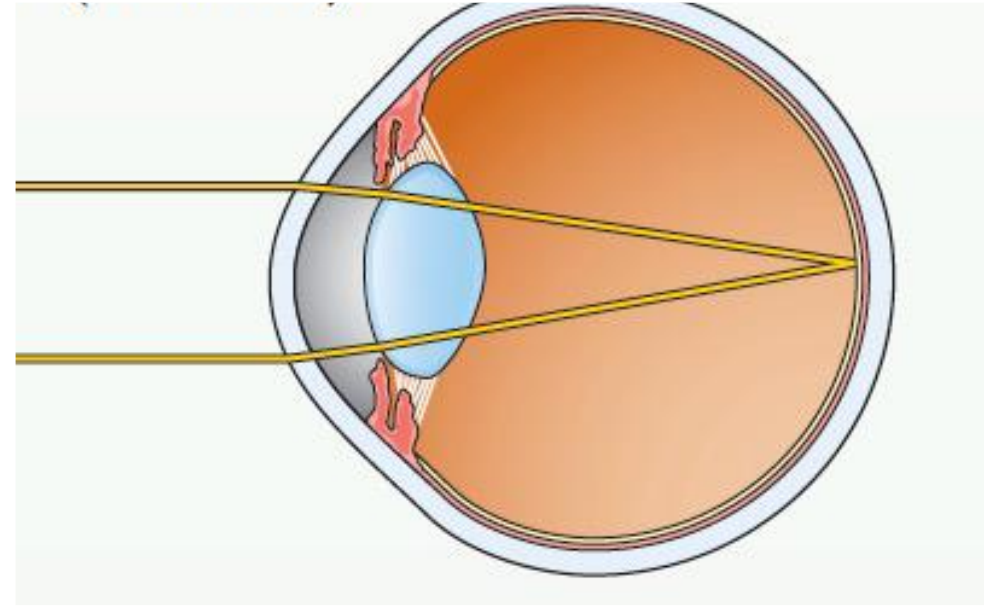
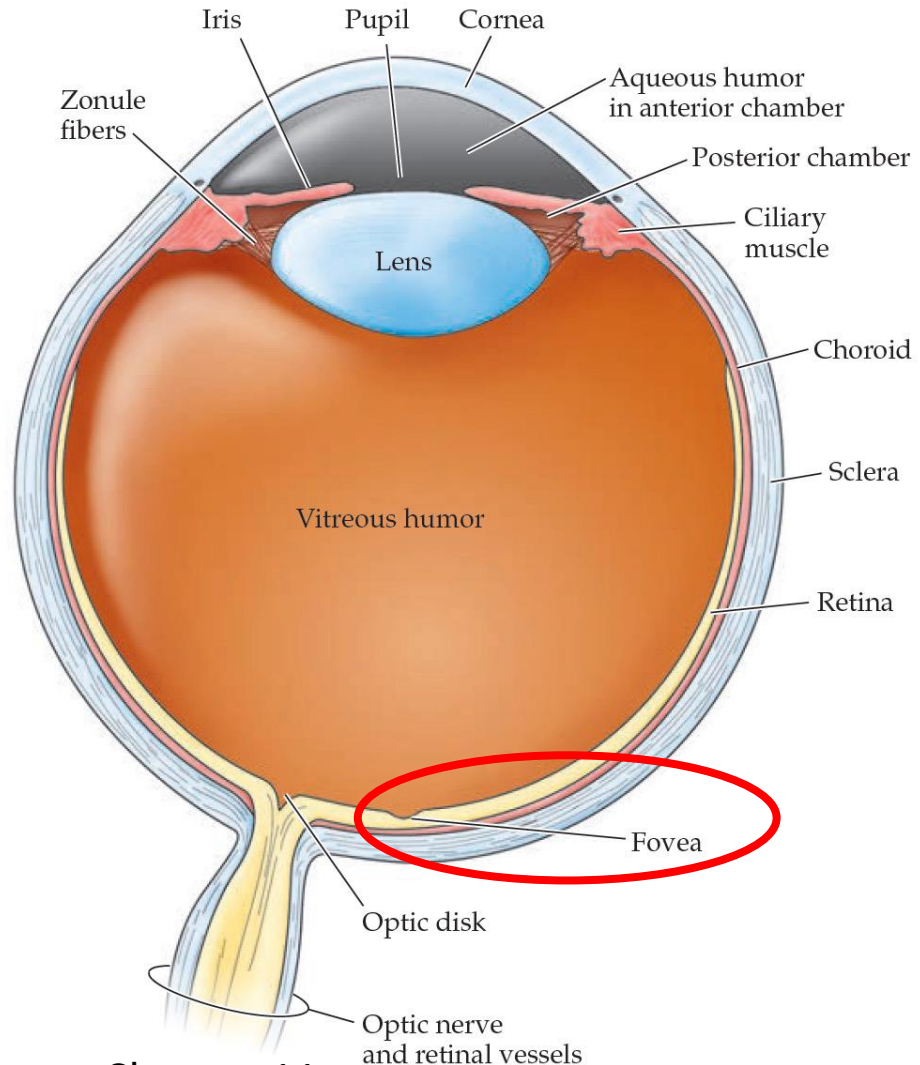


Copyright © 2007 Pearson Education, Inc., publishing as Benjamin Cummings.

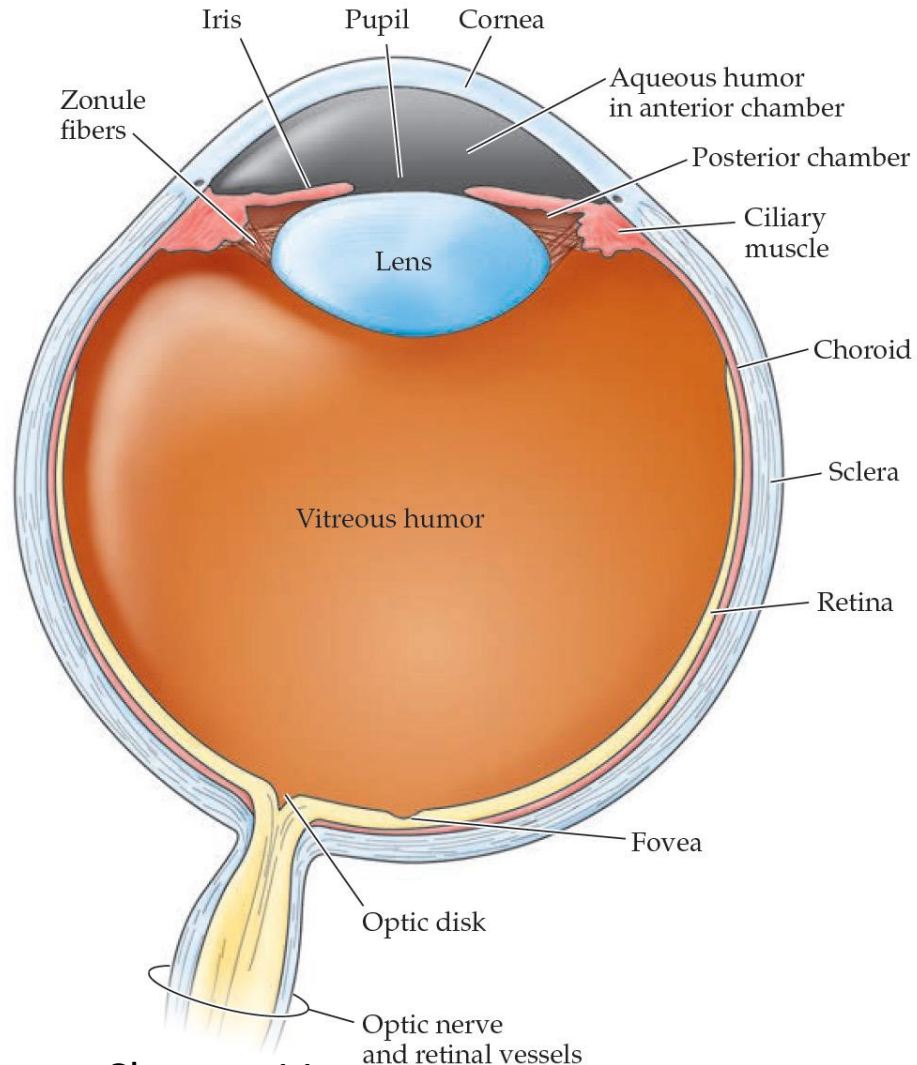
Fig. 10-4

Courtesy of Bob Desimone's class MIT 9.017

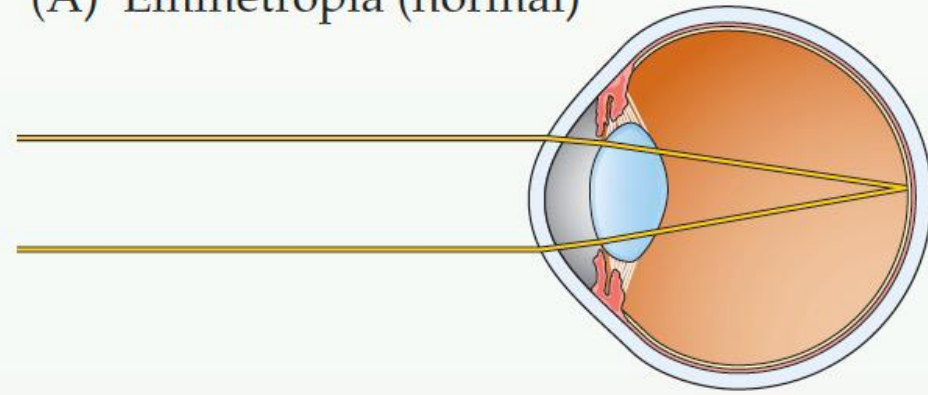
# Retina



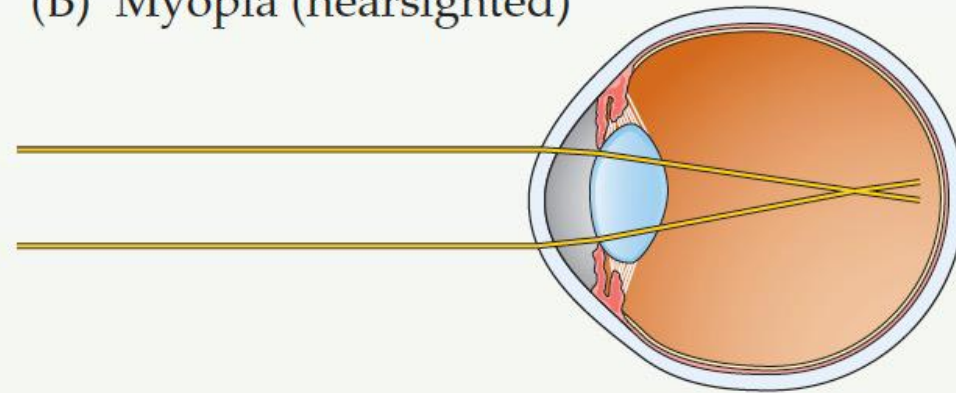
# Retina



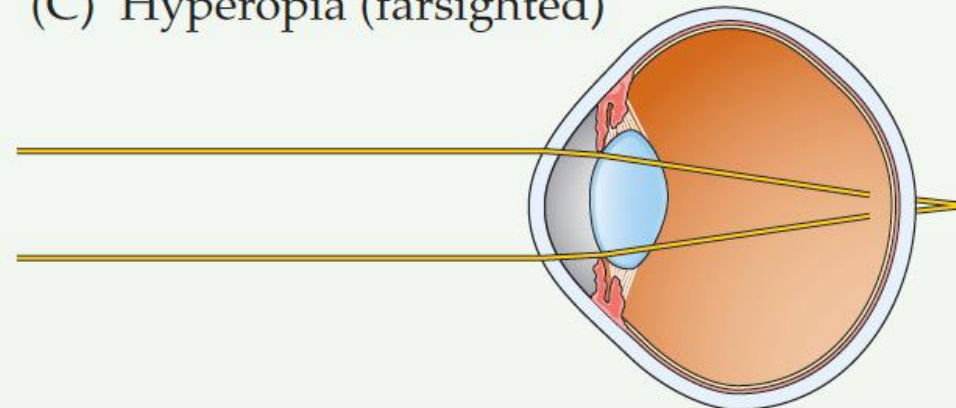
(A) Emmetropia (normal)



(B) Myopia (nearsighted)

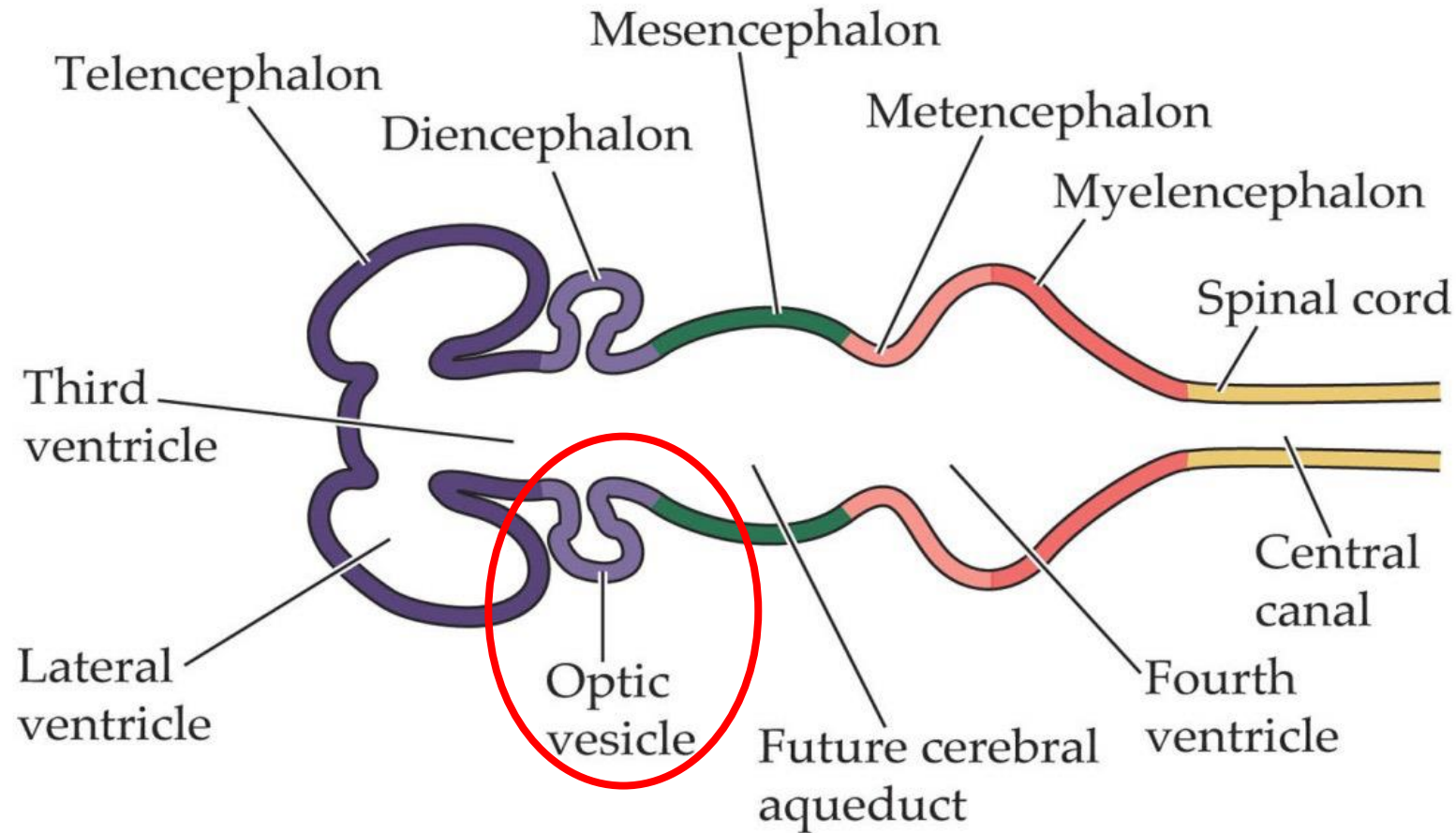


(C) Hyperopia (farsighted)

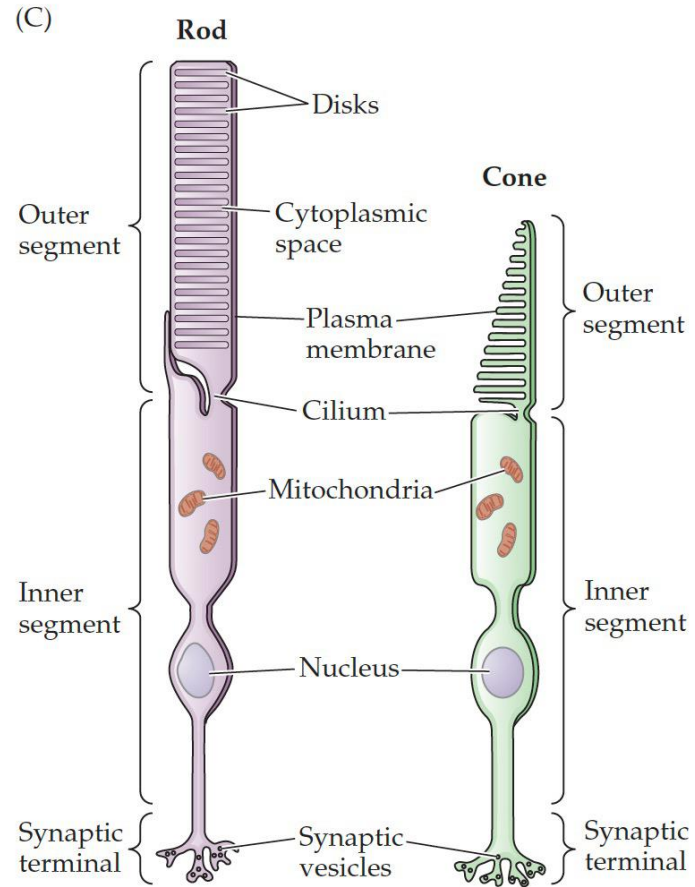




# Retina is part of central nervous system



# Photoreceptor cells: rods and cones



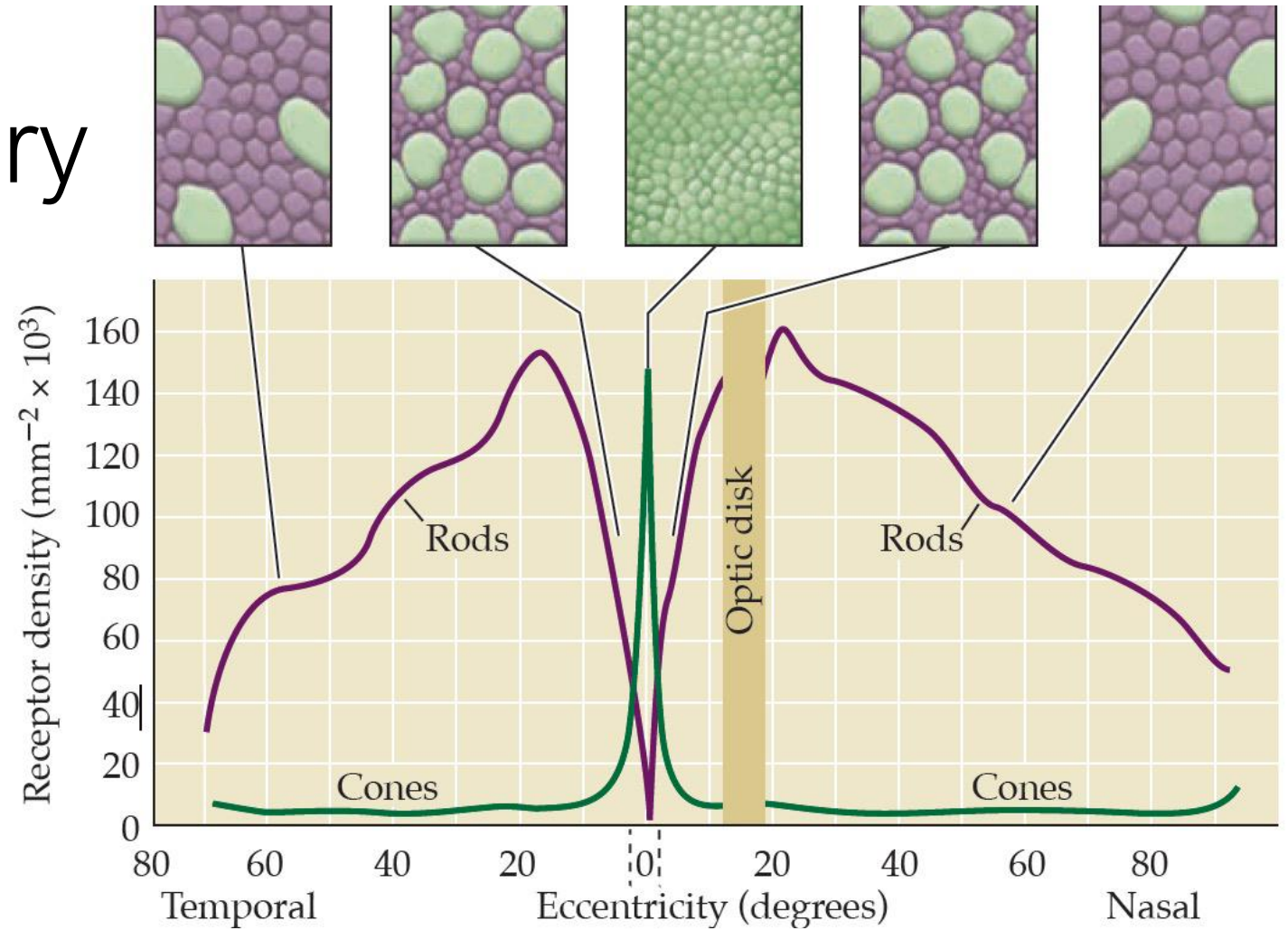
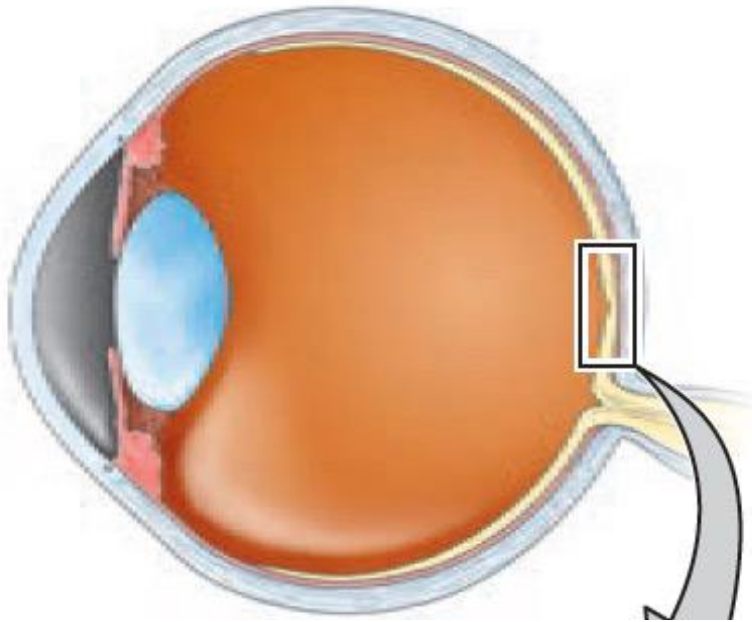
## Rods:

- Low spatial resolution
- High light sensitivity

## Cones:

- High acuity (spatial resolution)
- Less light sensitive

# Fovea vs periphery



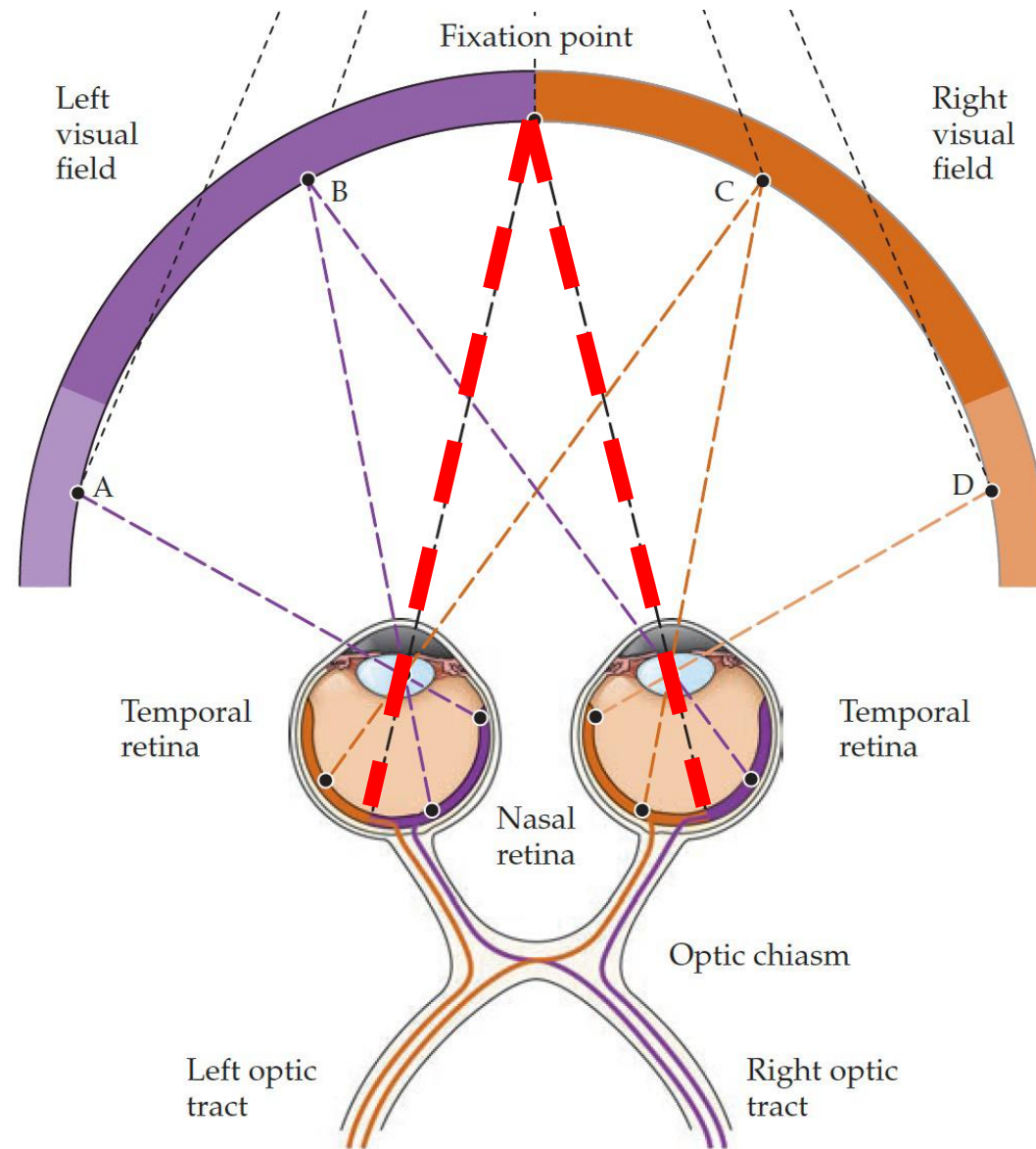
Fovea: high density of cones → better resolution

Periphery: rods > cones → worse resolution, better night vision

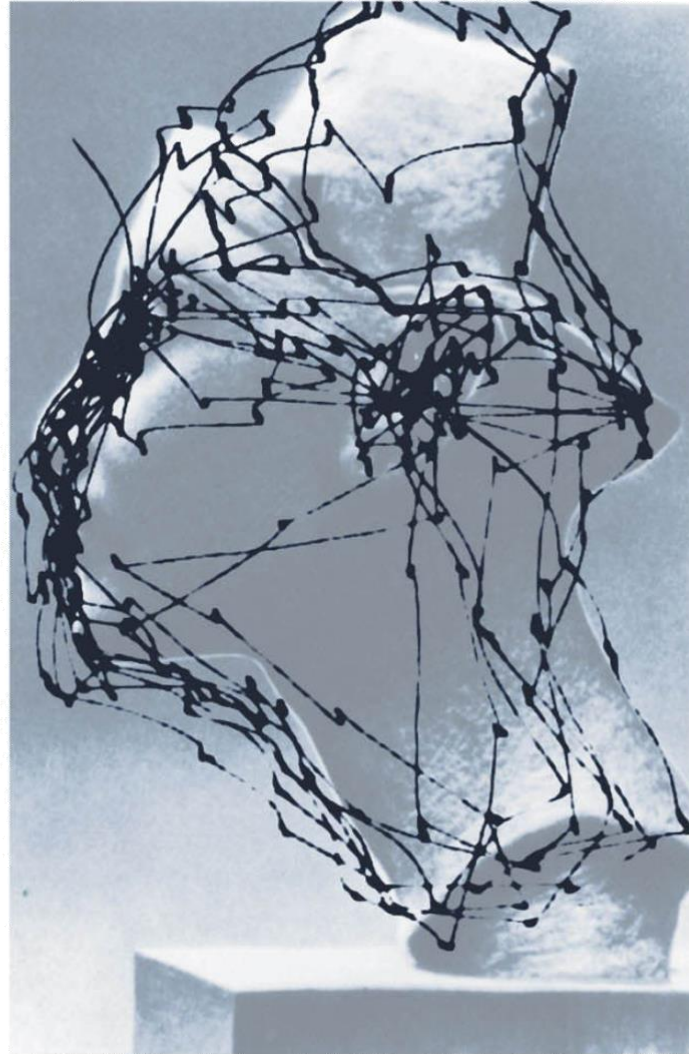
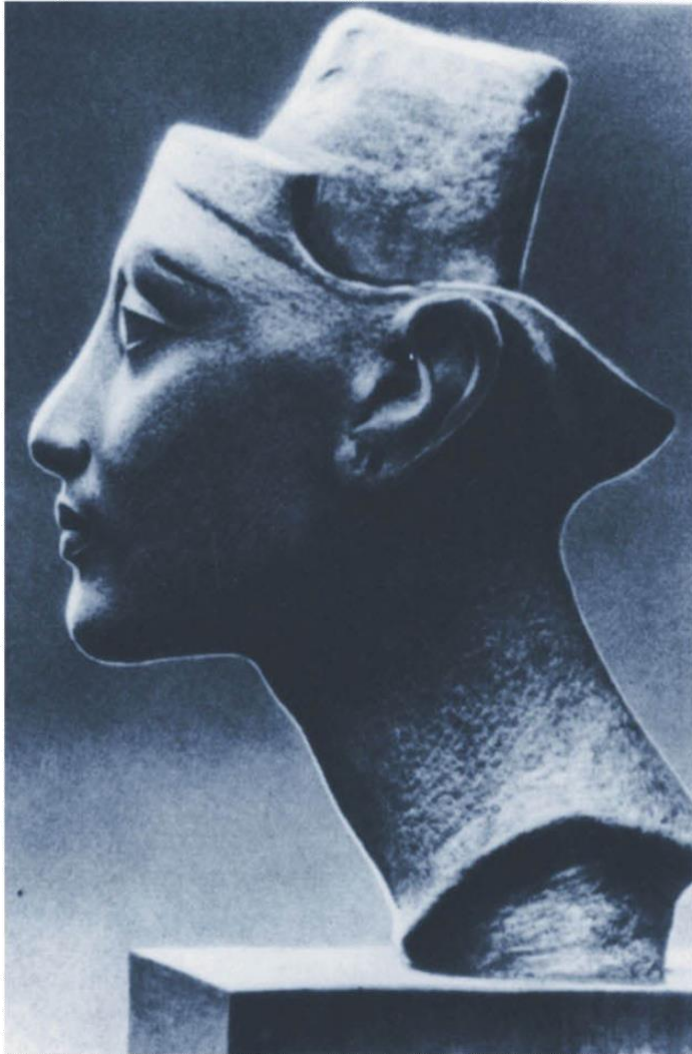
Optic disk: optic nerve to LGN → blindspot



# Fixation point falls on the fovea



# Foveation via saccades



To sample many parts of image at high resolution:

- Saccade to new point
- That part of the image can now be processed with high resolution (fovea)
- Repeat

# Blind spot

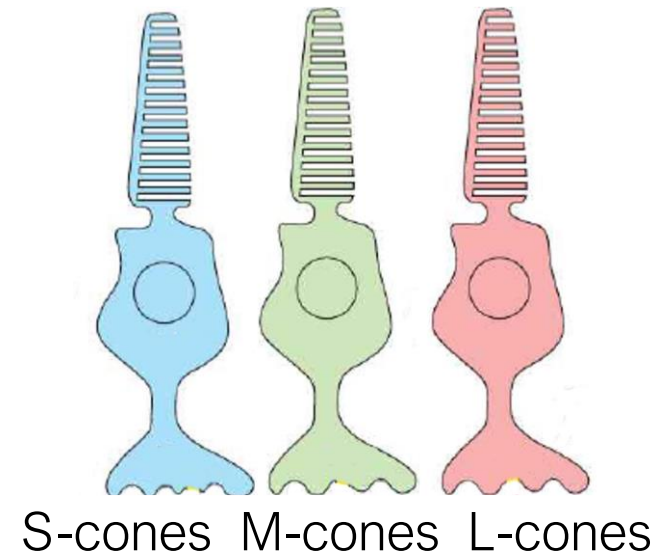
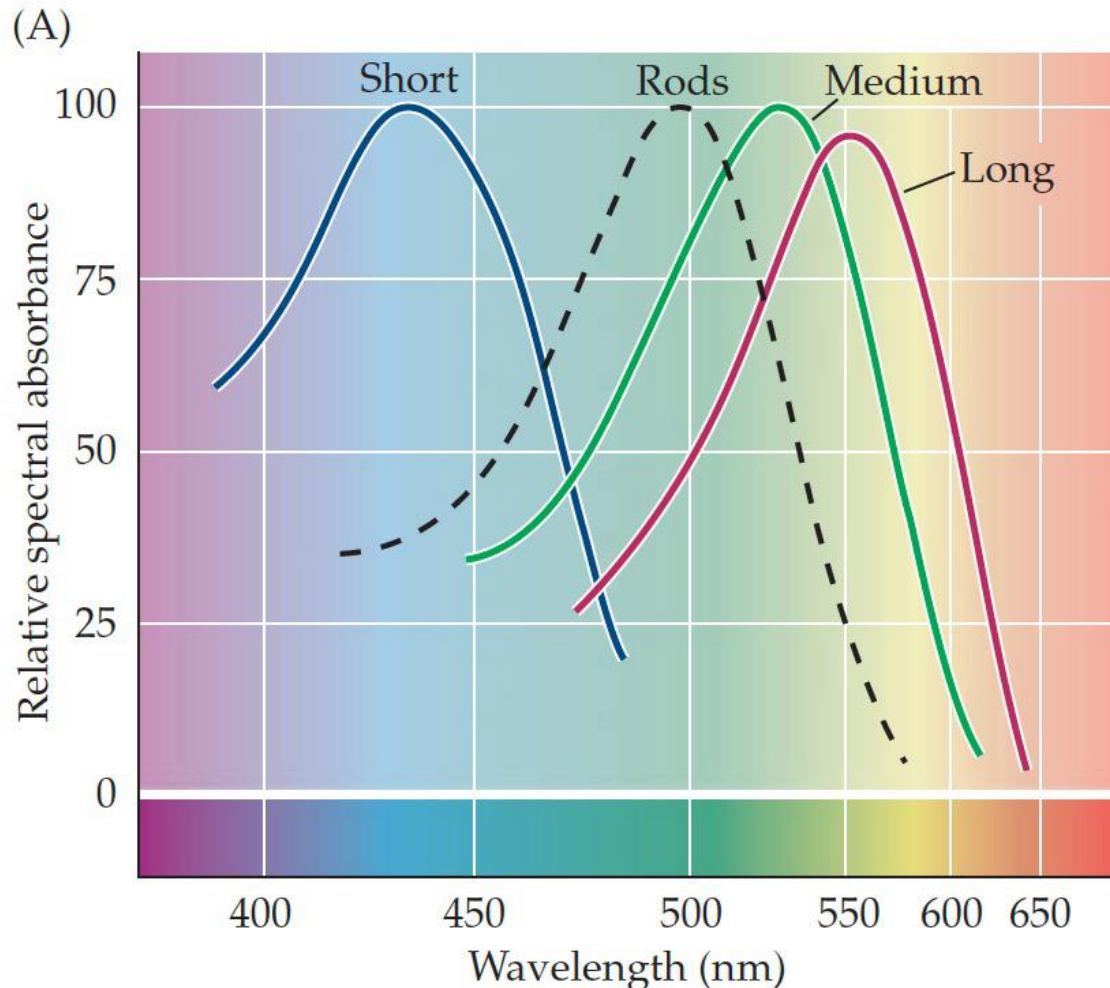
A demonstration of the blind spot:

Courtesy of Pavan Ramdya's class EPFL BIO-311

- Fixate on the cross (+) with your left eye (having your right eye closed)
- Move the image backward / forward around ~20 cm from the eye
- The circle should disappear (its image is on the blind spot!)
- The line will appear continuous (the image of the break is on the blind spot!)



# Color vision is realized by 3 types of cones with different spectral sensitivities

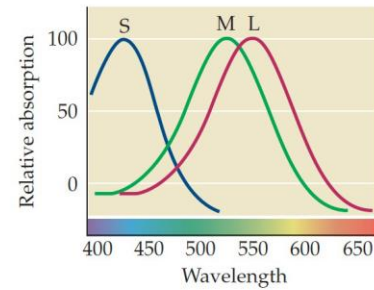


Identify specific colors via combination of cone activity.  
~No color vision in periphery!



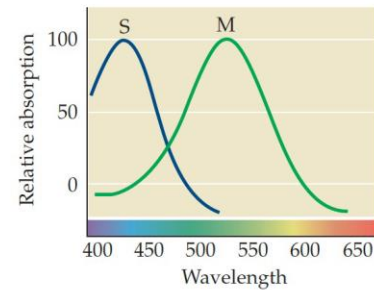
# Color blindness (dichromacy)

(A) Normal (trichromat)



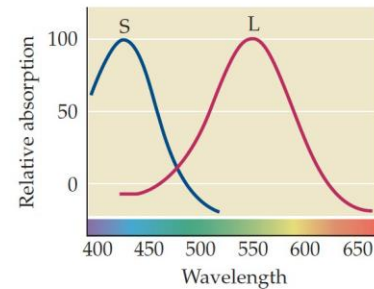
Neurotypical

(B) Protanopia



Loss of L-Cones → Red-Green color blind

(C) Deuteranopia



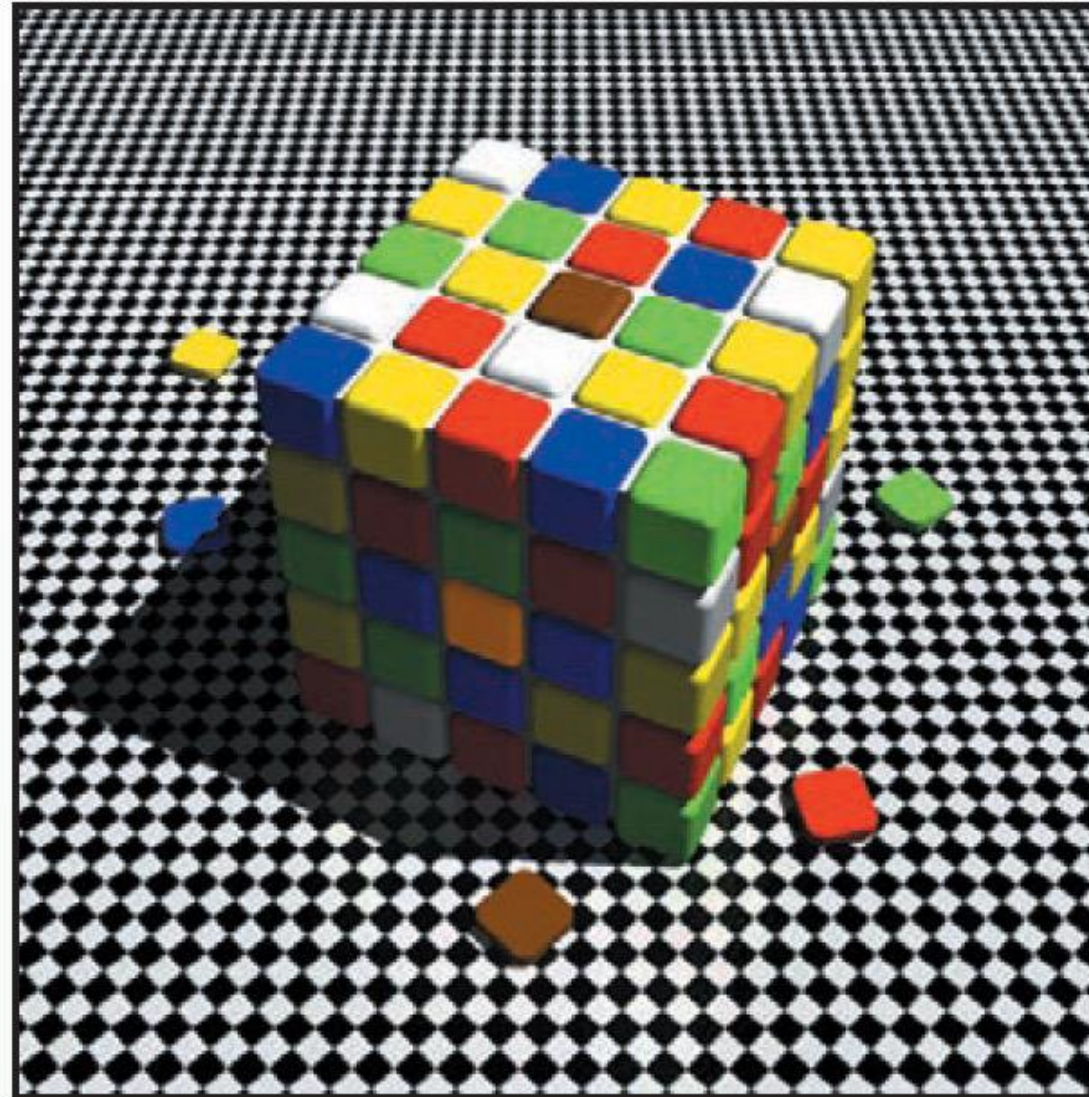
Loss of M-Cones → Red-Green color blind

Also: Tritanopia. Loss of S-Cones → Blue-Yellow (rare)



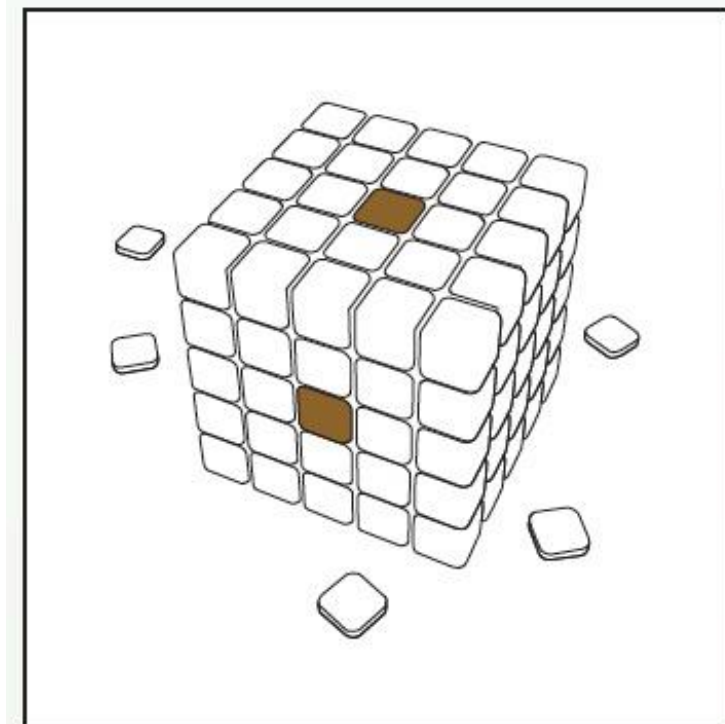
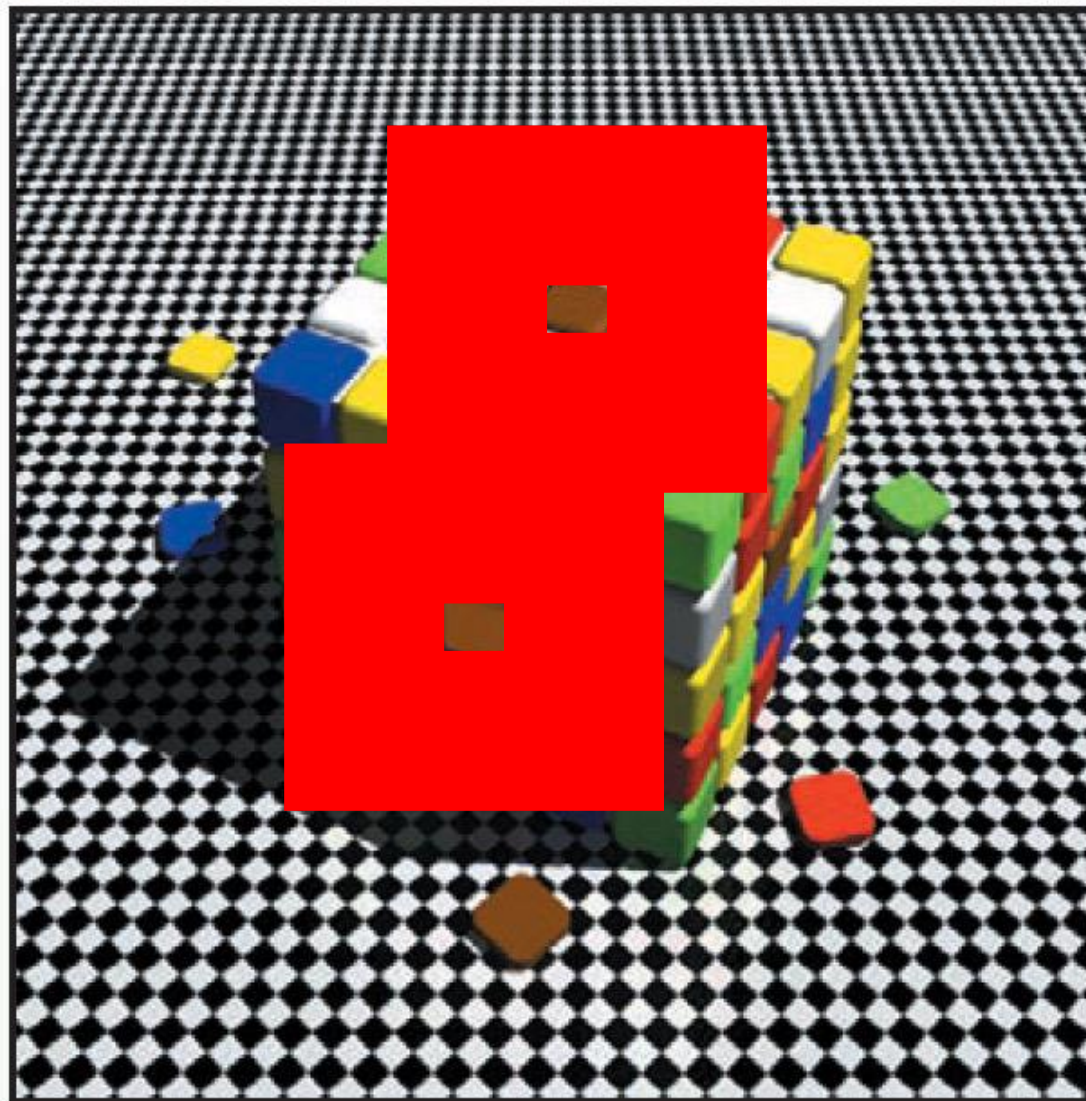
# Visual Context

What is the color  
of the middle  
squares?



# Visual Context

What is the color of the middle squares?





# The dress

Blue and black?

or

White and gold?



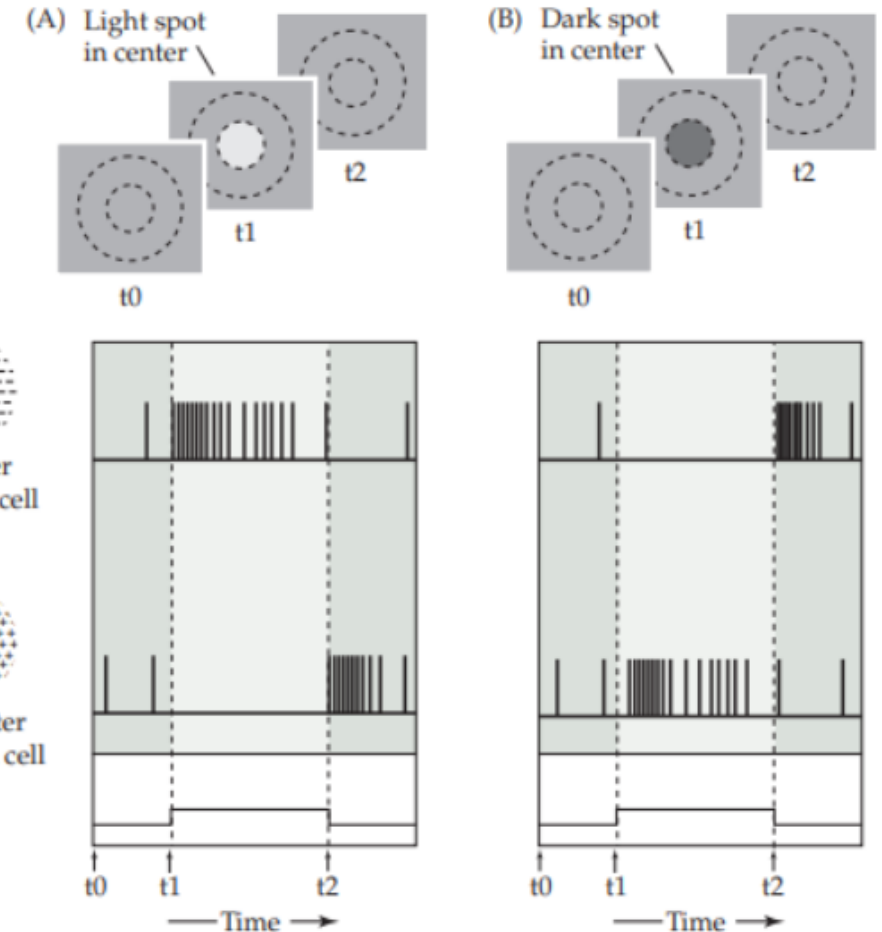
# A simple form of context in retinal ganglion cells: ON- and OFF-center

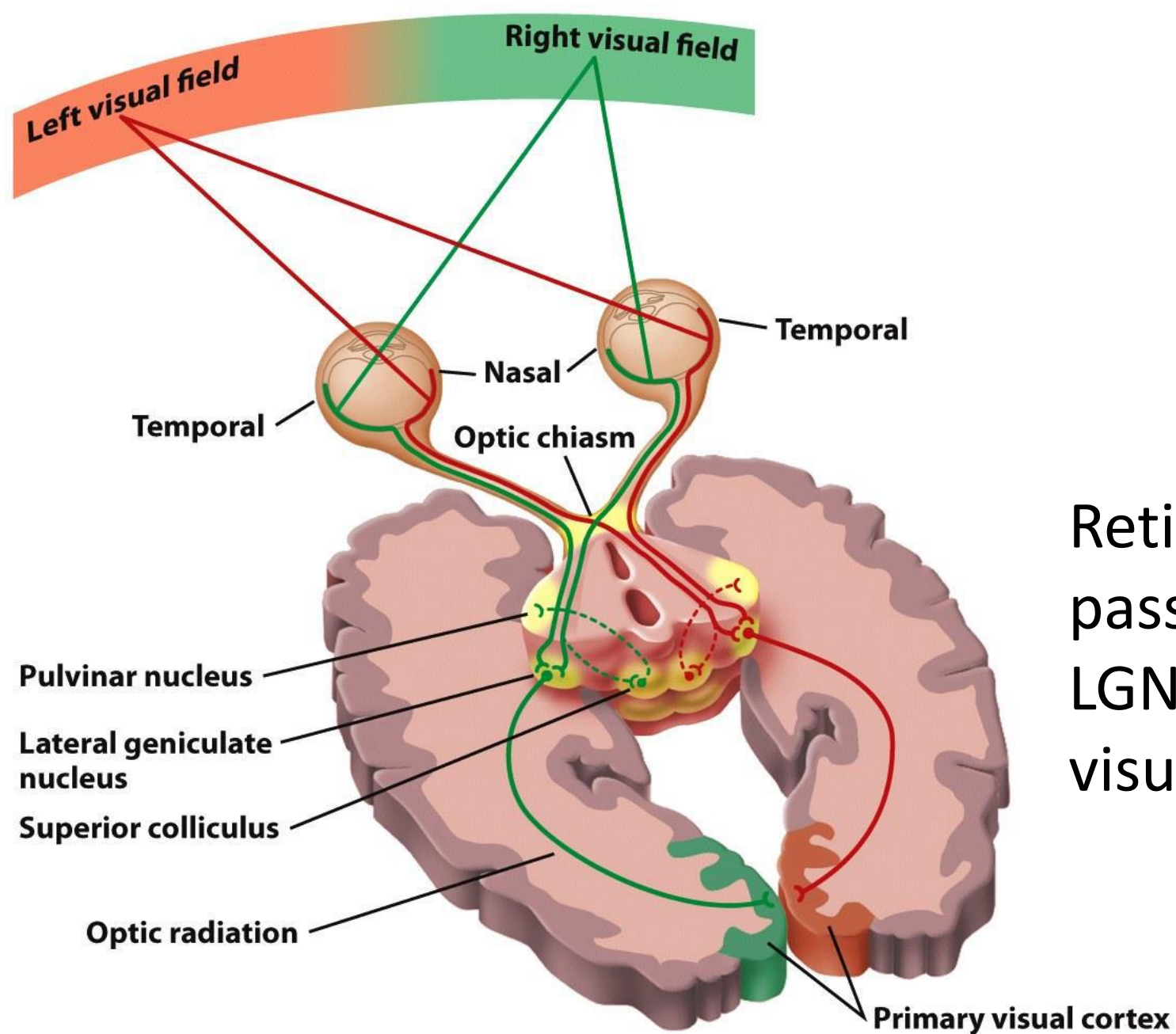


"ON-center":  
↑ AP- frequency  
in the center of the receptive field (RF)



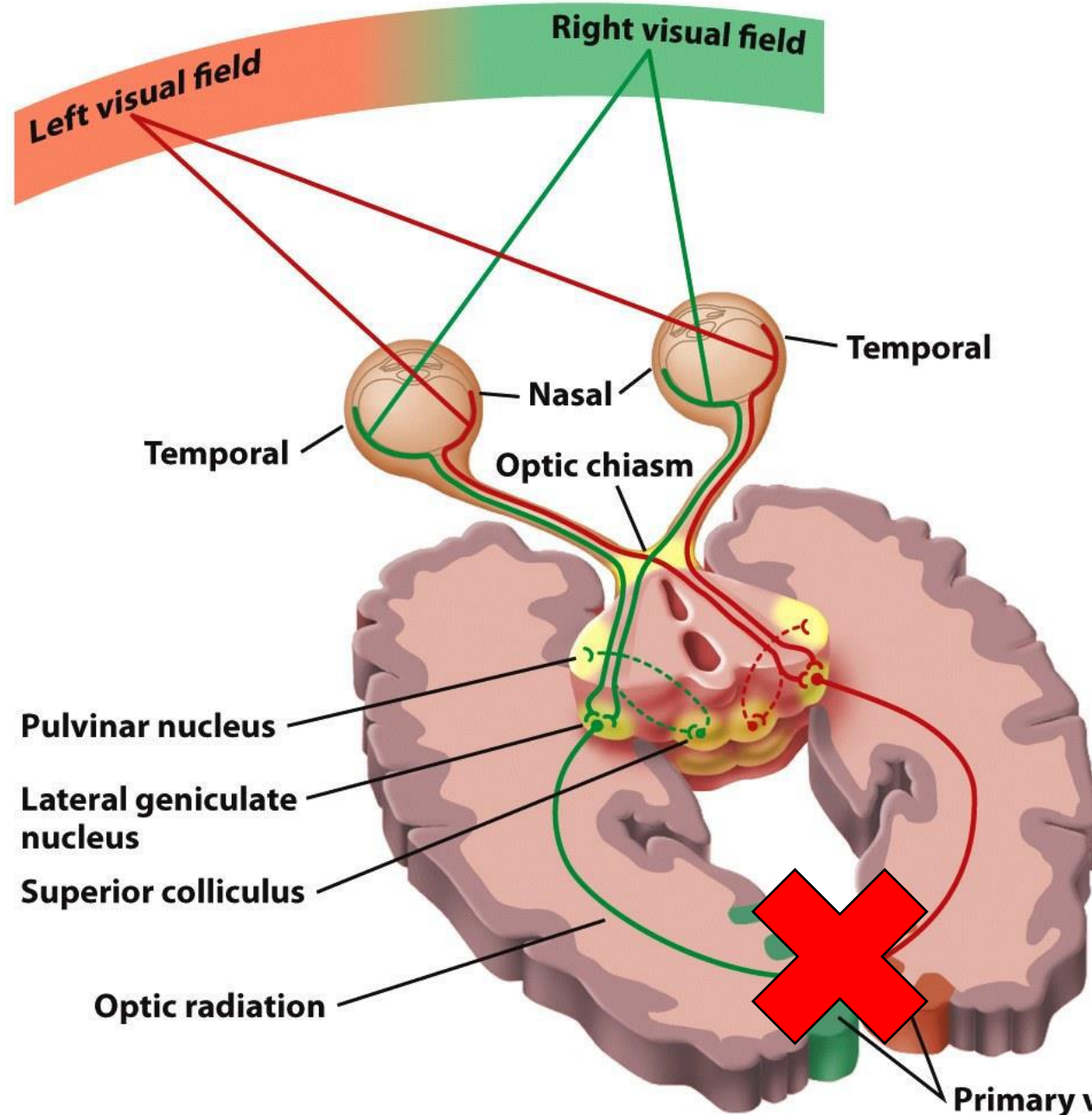
"OFF center"  
↓ AP-frequency  
in the center of the RF





Retinal input  
passes through  
LGN into primary  
visual cortex



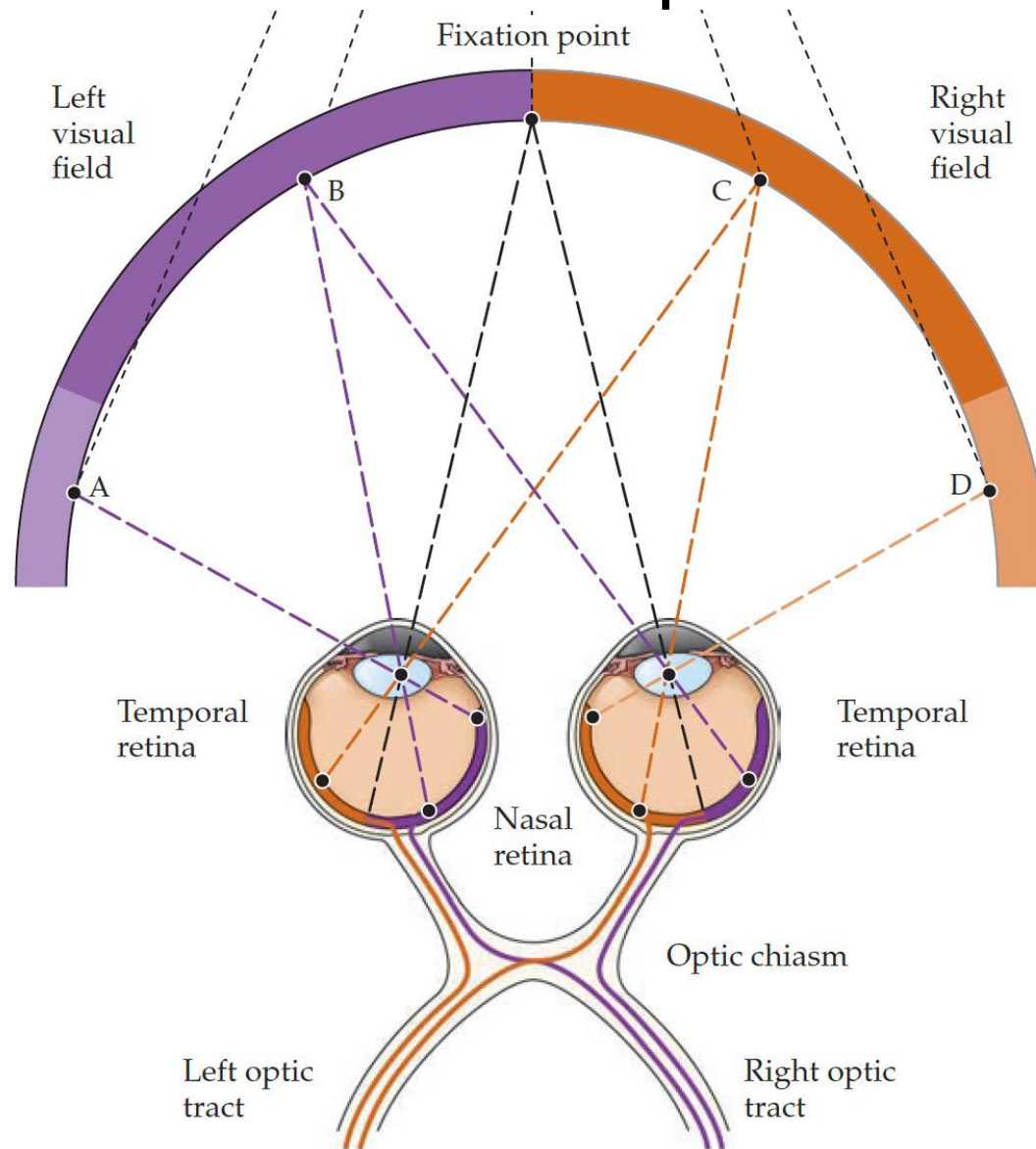


## Blindsight?

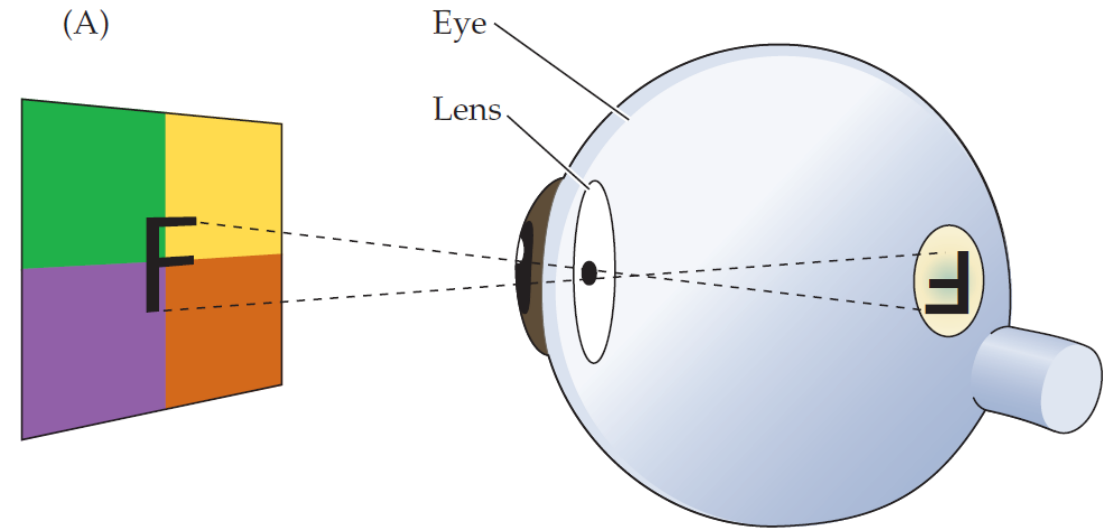
Some visual function without input into primary visual cortex

Courtesy of Bob Desimone's class MIT 9.017

# Contralateral processing along optic chiasm



Visual input is inverted – but your brain takes care of that



- Each eye perceives left **and** right visual field
- **Left** optic tract propagates only **right** visual field
- **Right** optic tract propagates only **left** visual field
- Contralateral processing: each hemisphere processes sensory input from the opposite side of the body

# NB: contralateral processing and split-brain patients

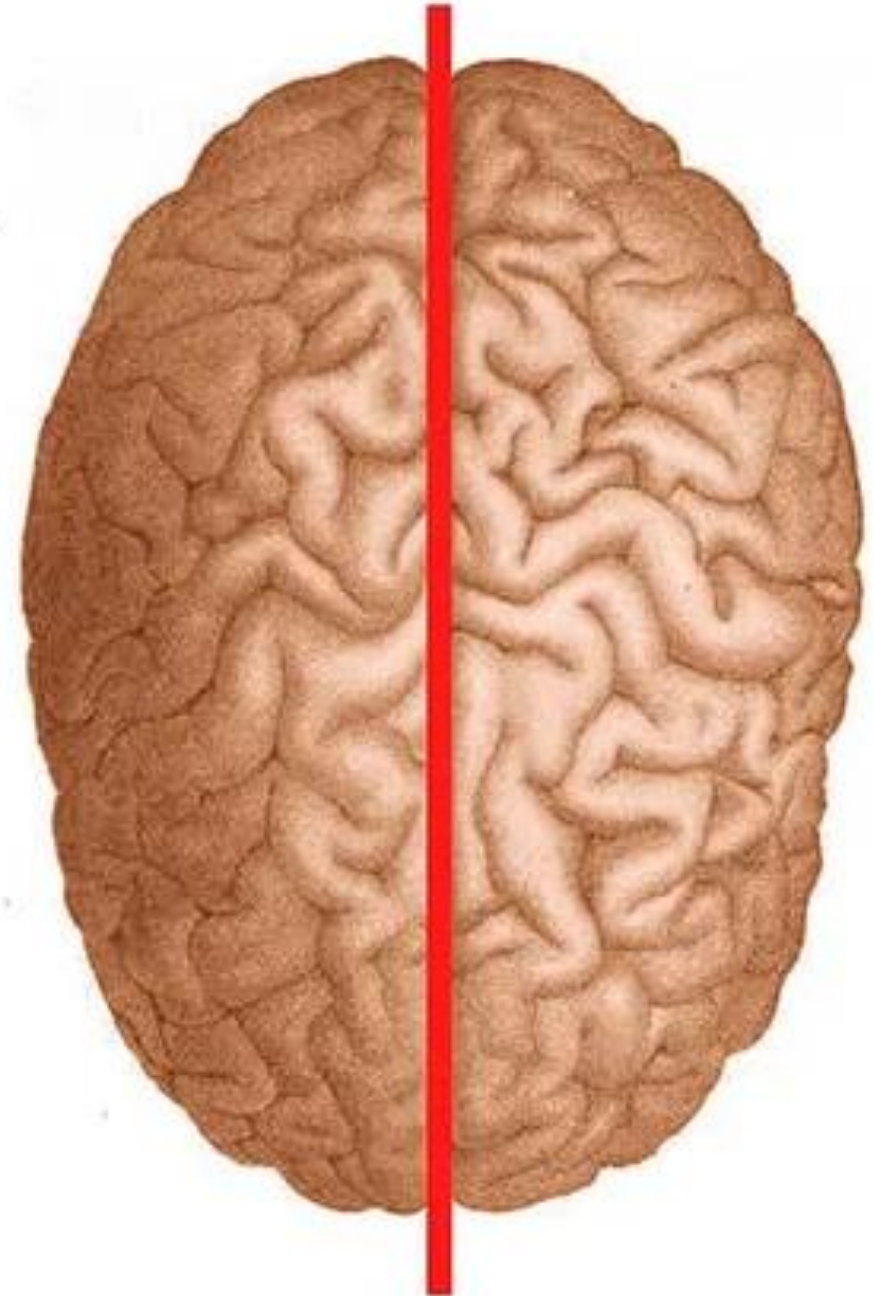
Two hemispheres are connected via corpus callosum.

Sever this connection as a last resort to treat epilepsy.

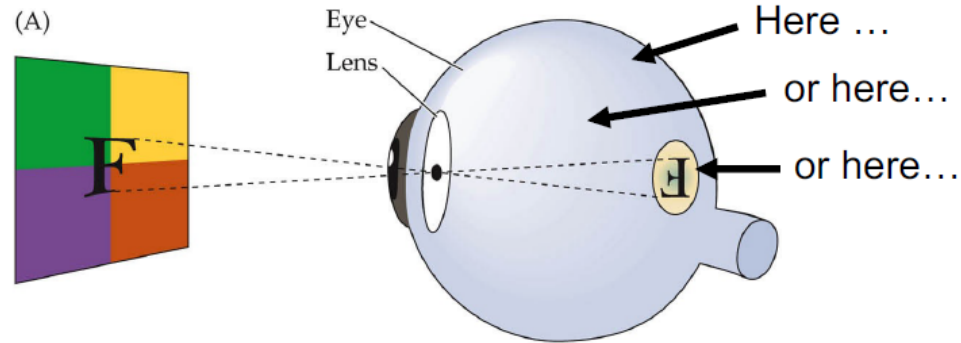
Effect in some patients: **alien-hand syndrome**, e.g.

- Ask patient to pick up glass (language ~ left hemisphere)
- Right hand will attempt to pick up glass (LH → right side of body)
- Left hand will interfere (left side of body → RH → did not receive language input)

Seen as evidence for dual consciousness after corpus callosotomy.

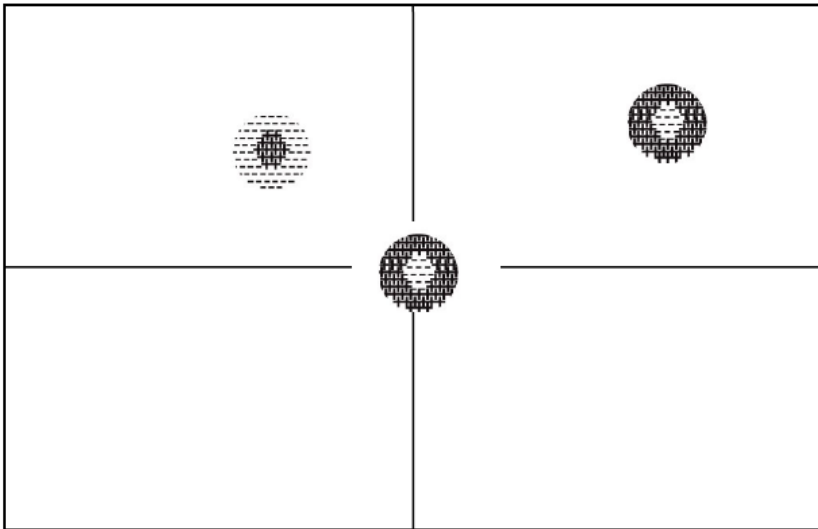


# Receptive fields



For each neuron: the part of the visual field for which it is most sensitive

Field of view:



If you record from a retinal ganglion cell (RGC).  
They fire APs with generally two types of responses:



"ON-center":  
↑ AP- frequency  
in the center of the receptive field (RF)



"OFF center"  
↓ AP-frequency  
in the center of the RF

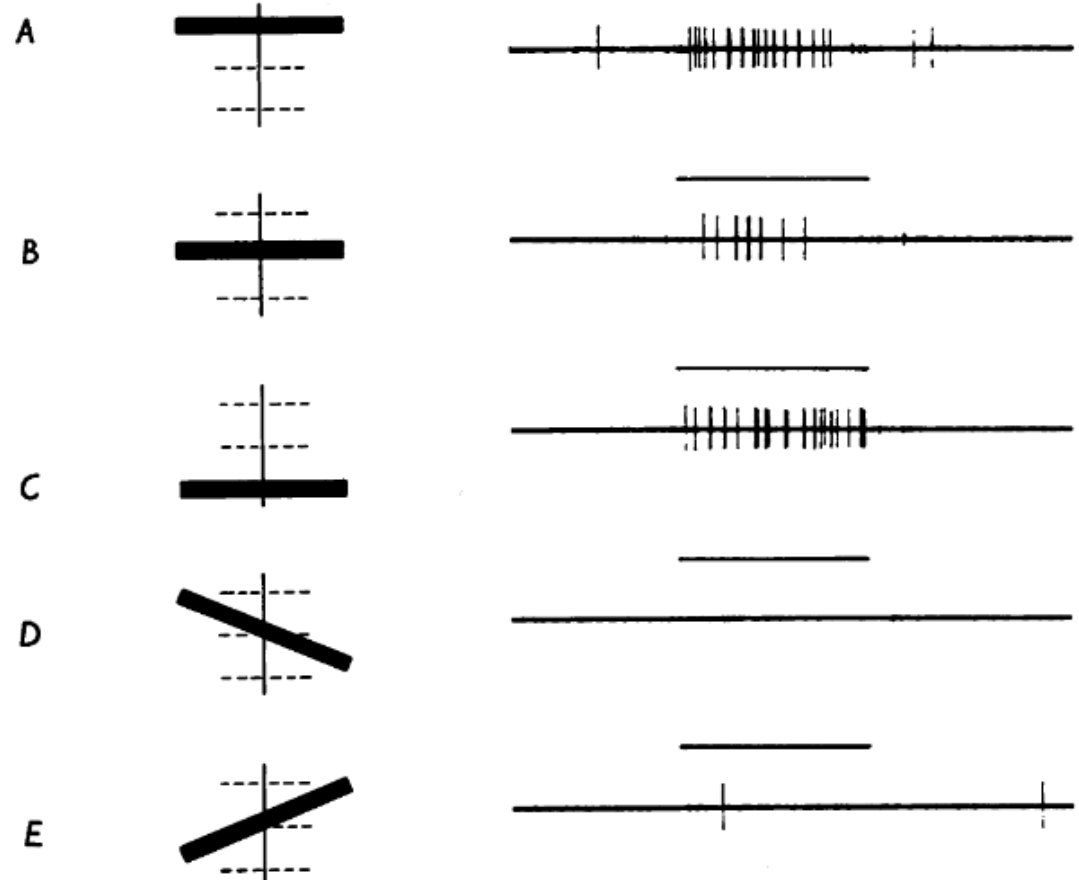
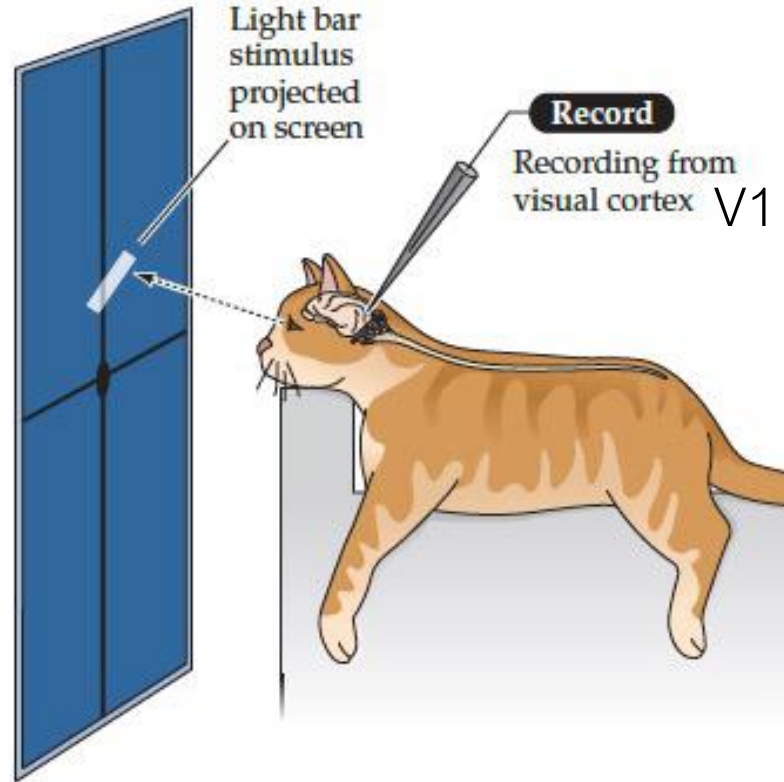


# V1

Receptive fields:  $\sim 1^\circ$  visual angle.

Textbook function: edge detection.

(A) Experimental setup





# V1

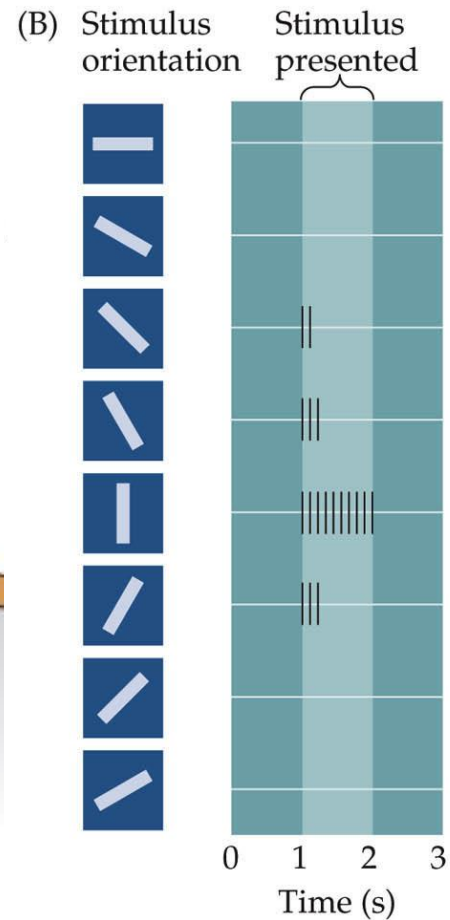
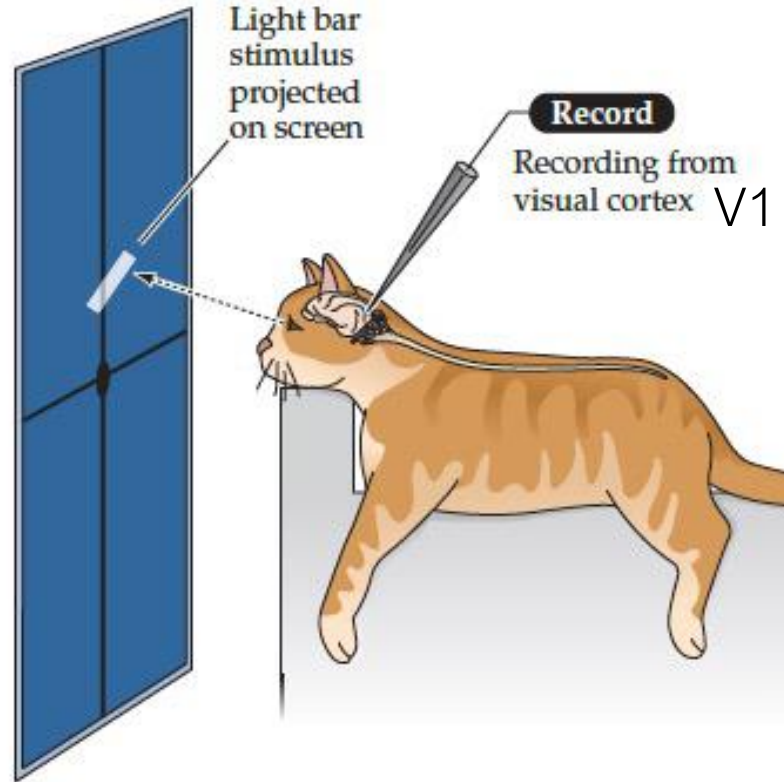
Receptive fields:  $\sim 1^\circ$  visual angle.

Textbook function: edge detection.

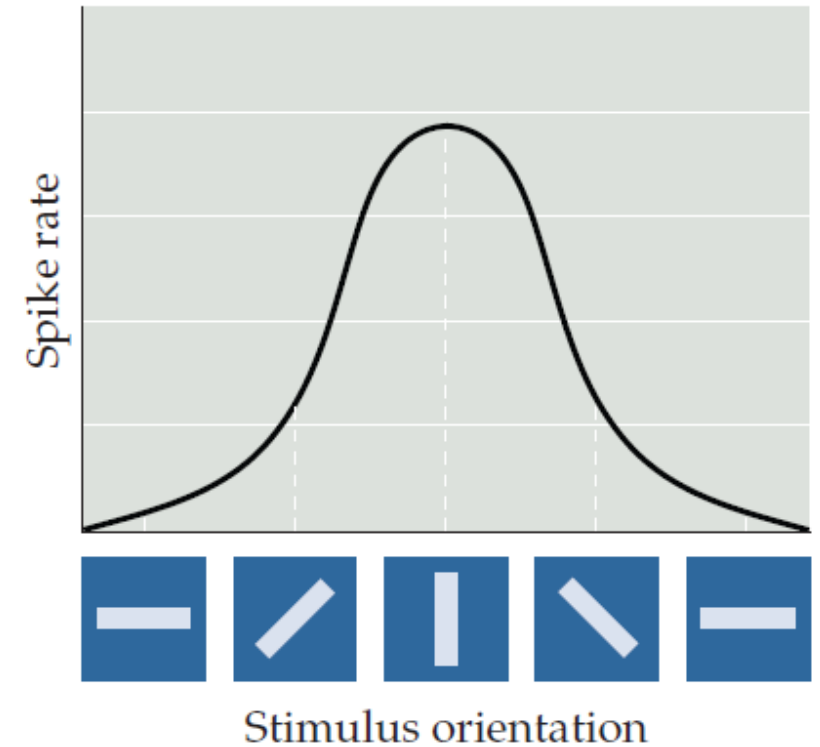
Hubel and Wiesel 1962

Nobel Prize 1981

(A) Experimental setup



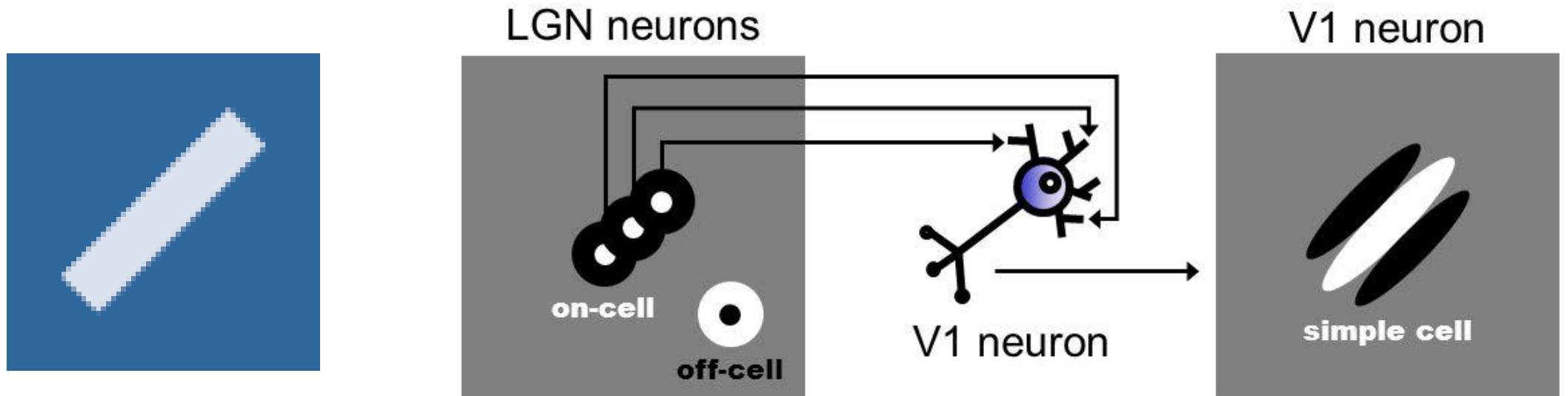
(C)



# V1

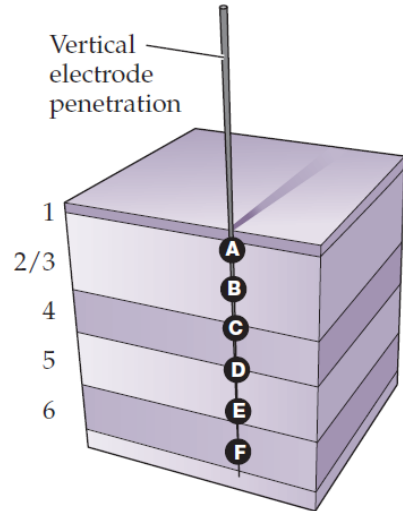
Receptive fields:  $\sim 1^\circ$  visual angle.

Textbook function: edge detection.

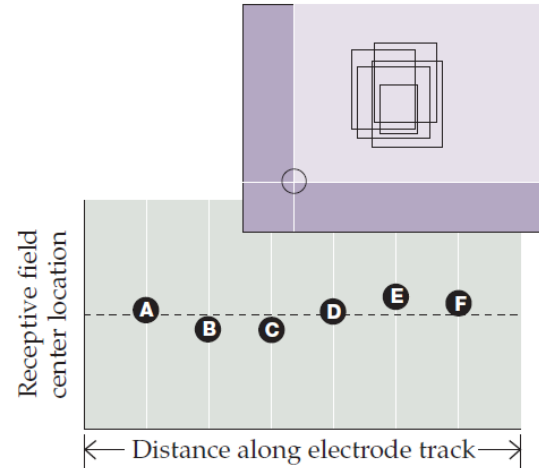


# V1 columnar organization

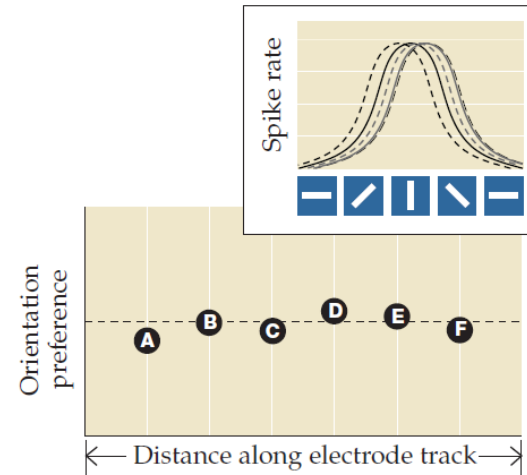
(A)



Receptive field position

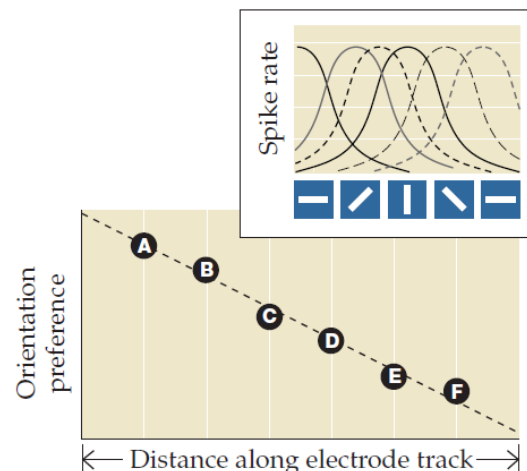
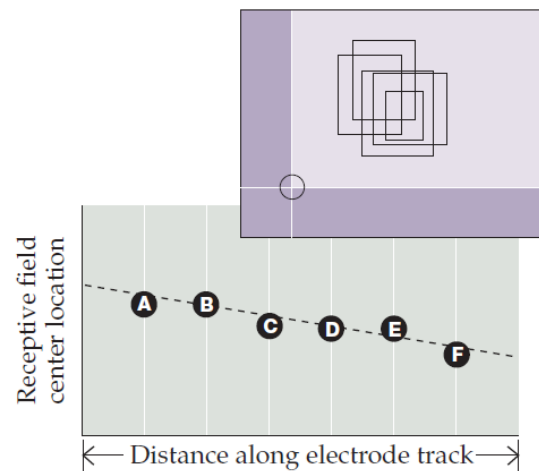
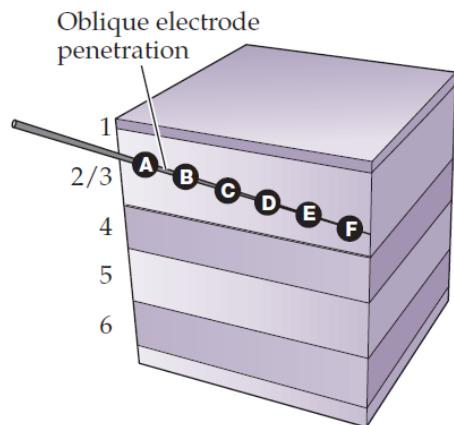


Orientation tuning curves



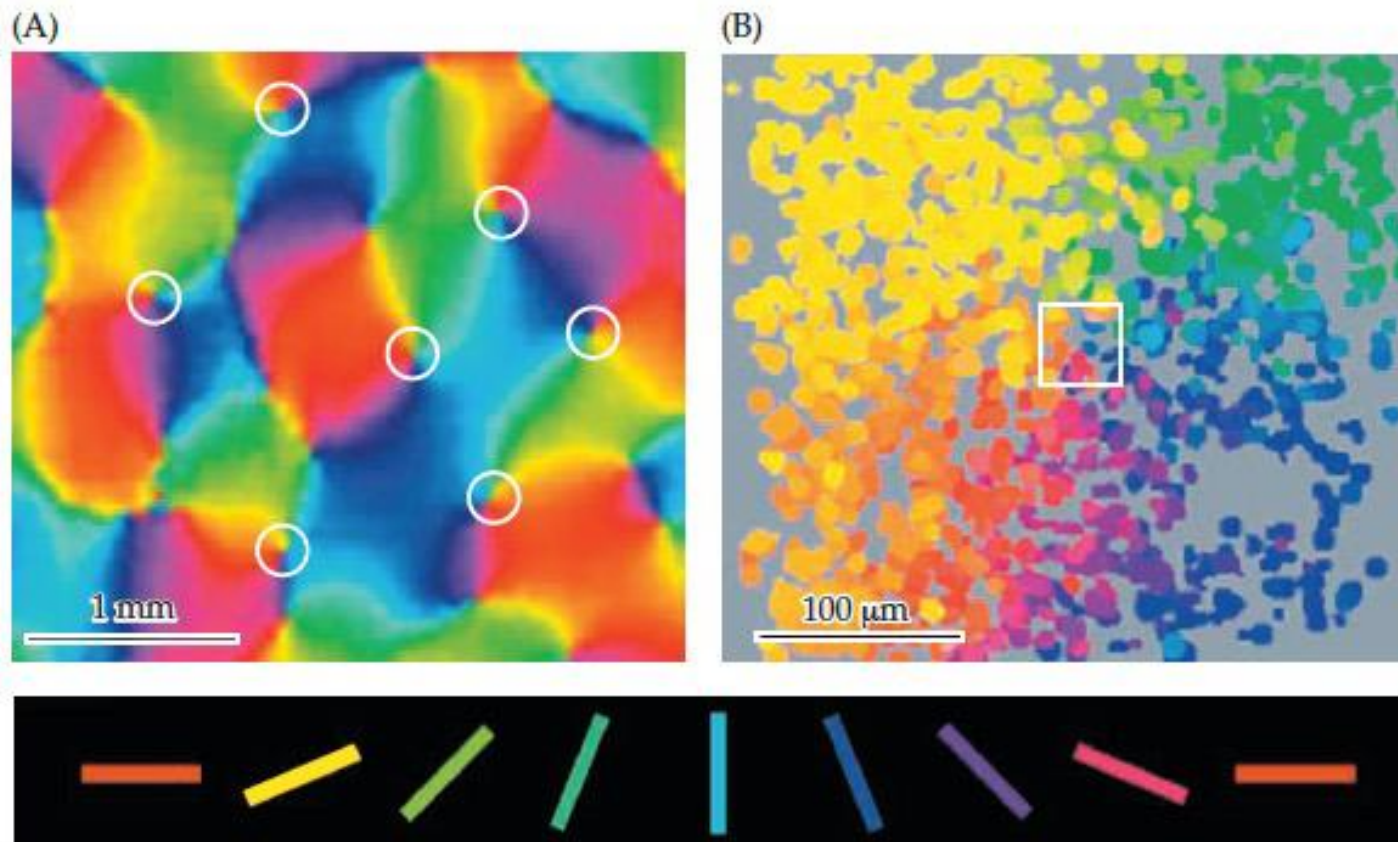
Layers in cortical micro-circuit typically have highly similar receptive fields  
→ **Organized in columns**

(B)



Adjacent columns typically have adjacent receptive fields  
→ **Organized in maps**

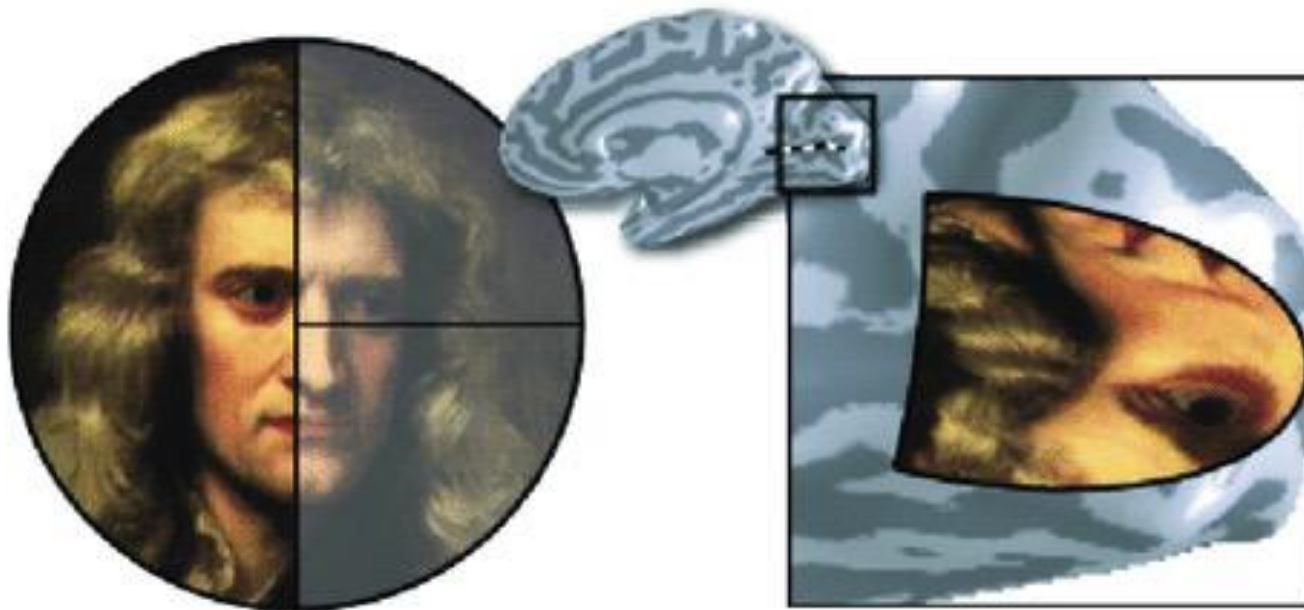
# V1 columnar organization: pinwheels



Nearby V1 neurons tend to respond to similarly oriented bars, organized in a pinwheel structure

**FIGURE 12.10** Functional imaging reveals orderly mapping of orientation preference in the primary visual cortex.

# V1 retinotopy

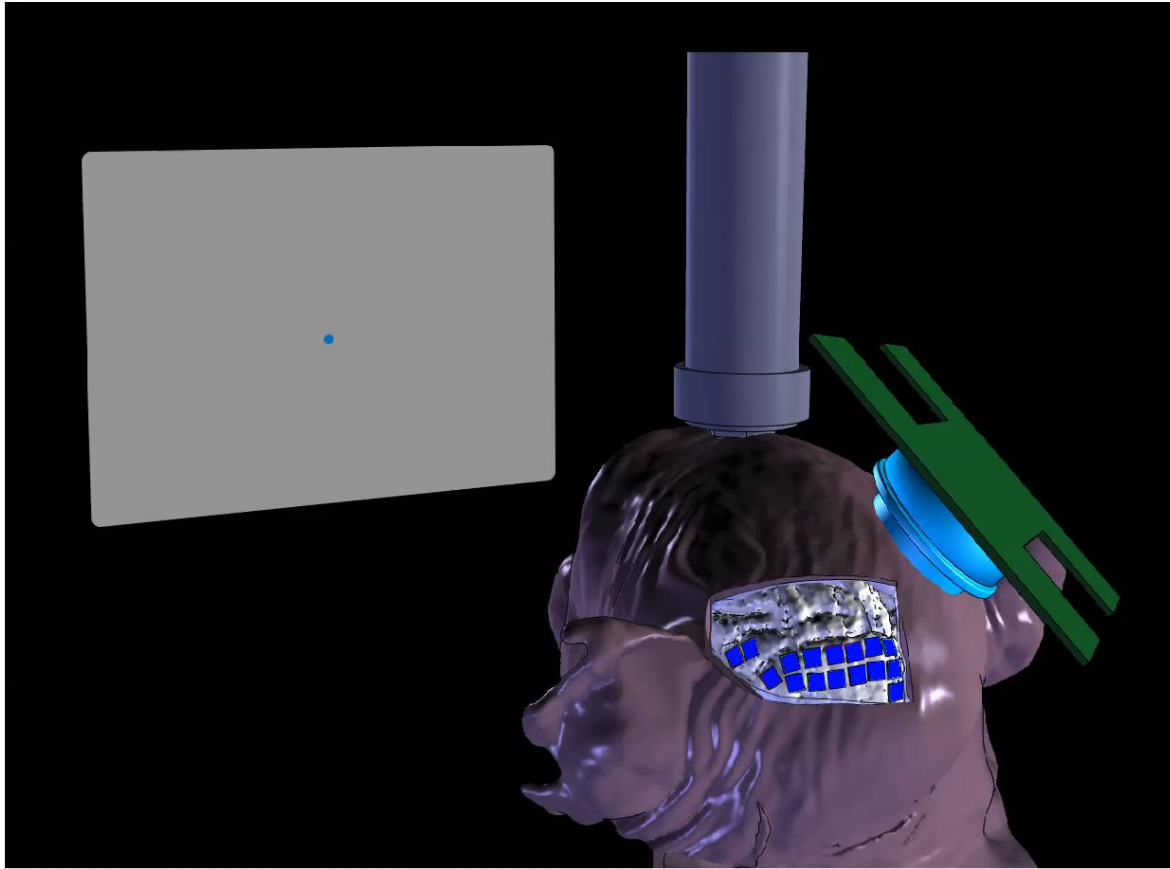


Mapping of visual input  
from retina to neurons

In V1: relatively well-  
preserved spatial  
organization of inputs.



# V1 retinotopy allows “painting on cortex” with electrical micro-stimulation



*Beauchamp et al. 2020*

*Fernández et al. 2021*

# Computational model of V1: Gabor filters

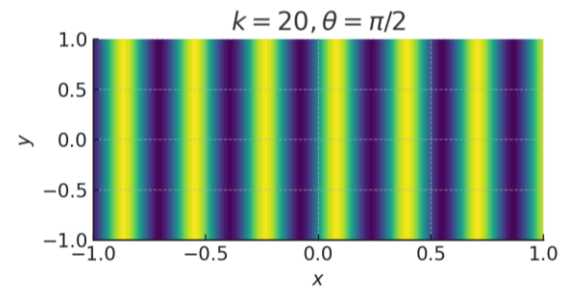
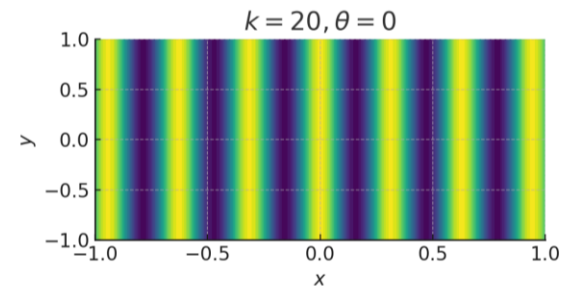
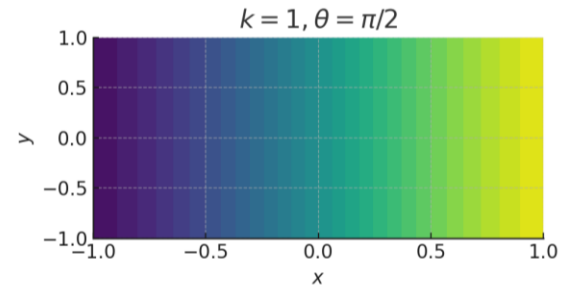
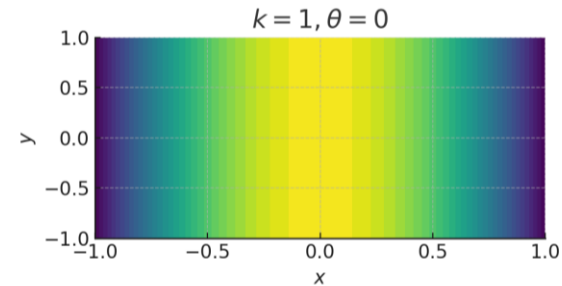
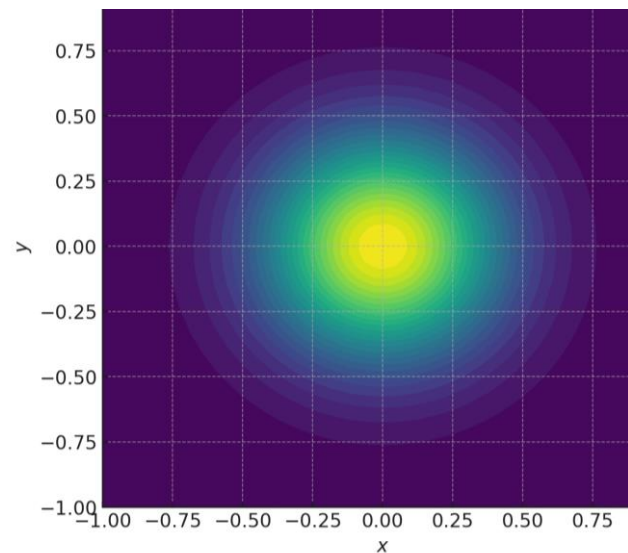
We can approximate the spatial receptive field of a simple cell with the Gabor function:

$$f(x, y) = \frac{1}{2\pi\sigma_x\sigma_y} \cdot \exp\left(-\frac{x^2}{2\sigma_x^2} - \frac{y^2}{2\sigma_y^2}\right) \cdot \cos(kx - \varphi)$$

receptive field location

spatial phase

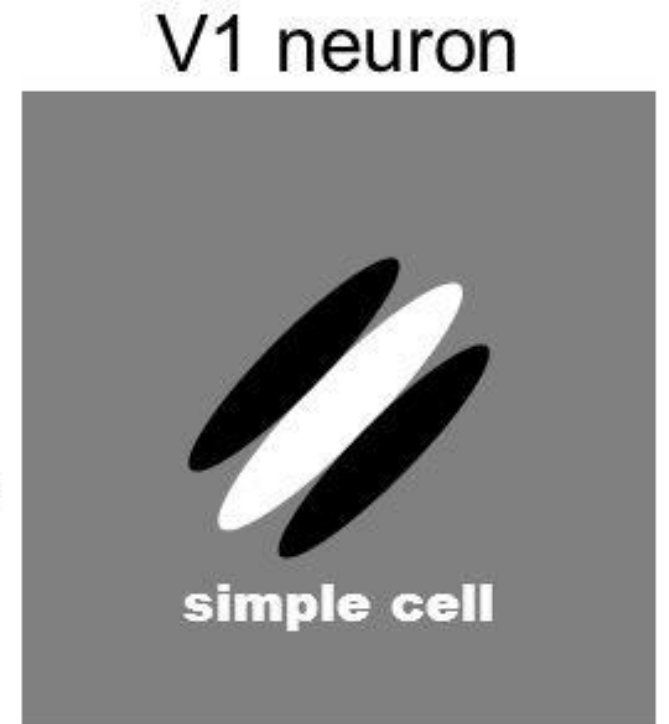
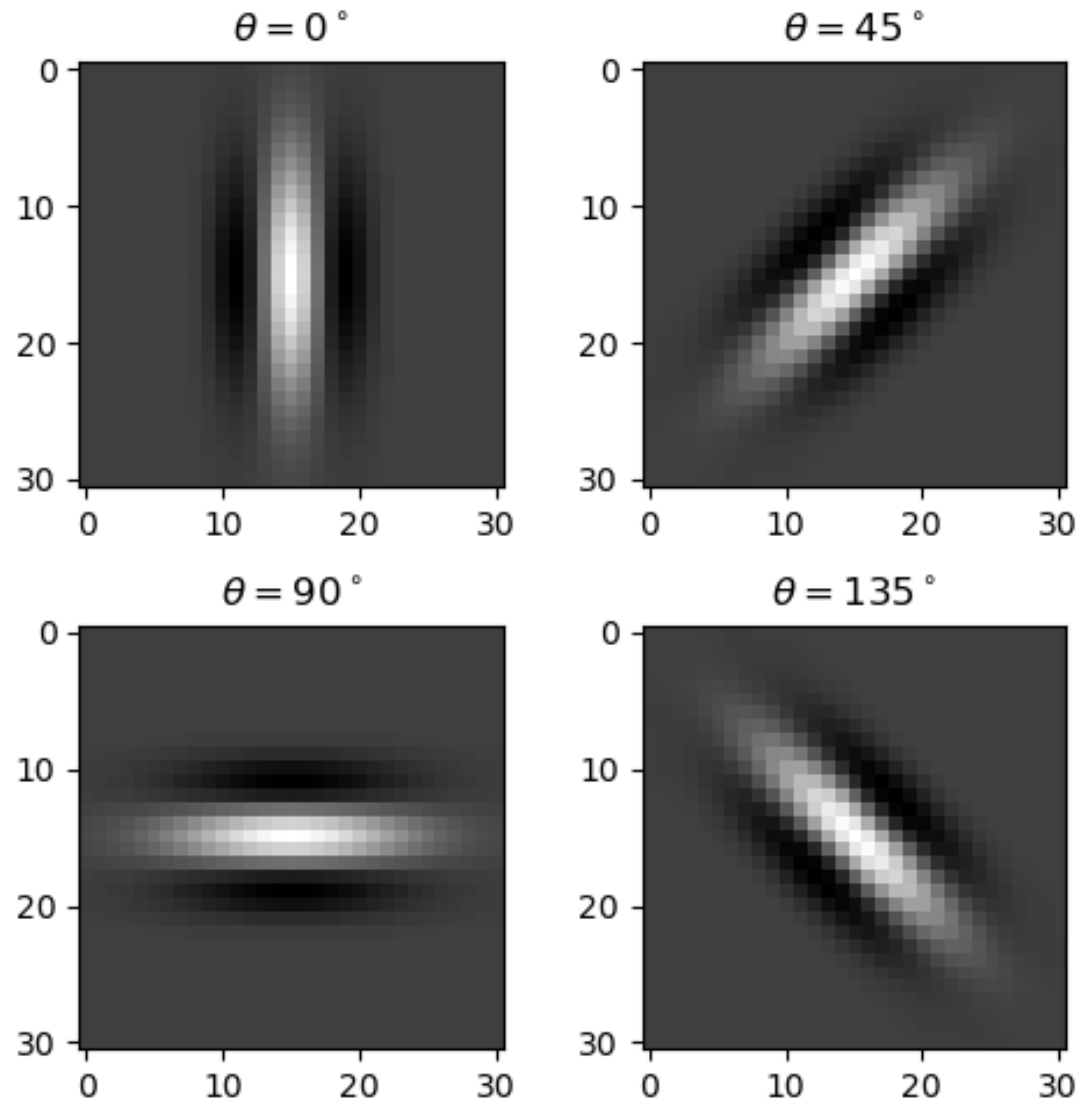
no rotation with this formulation, need x+y dependency



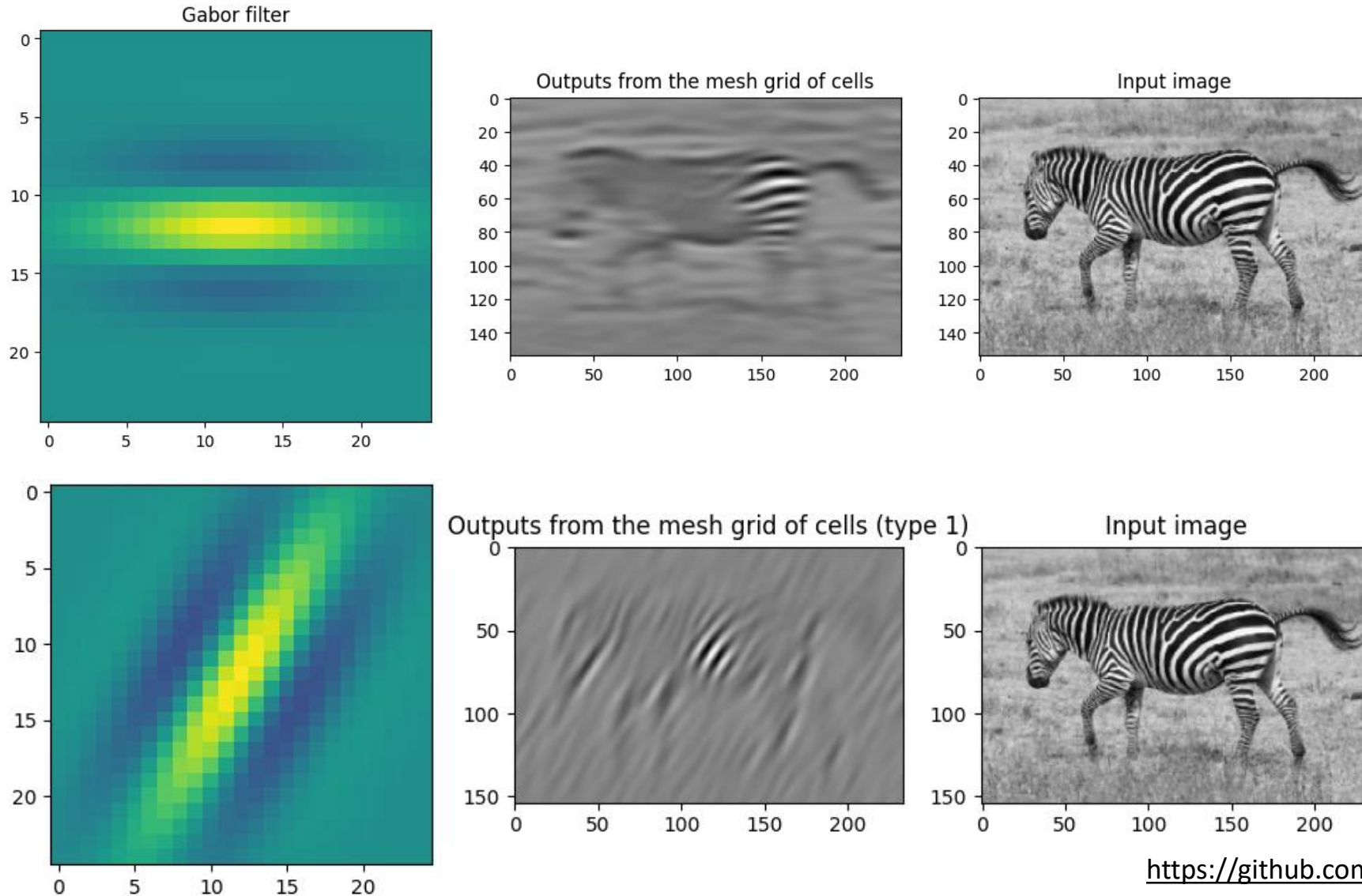
with:

- $\sigma_x, \sigma_y$  determine the spatial receptive field in x and y direction, respectively
- $k$  is the preferred spatial frequency (i.e. the spacing of light/dark bars)
- $\varphi$  is the preferred spatial phase

# Gabor model: example filters



# Gabor model: example filters applied





# Visual hierarchy

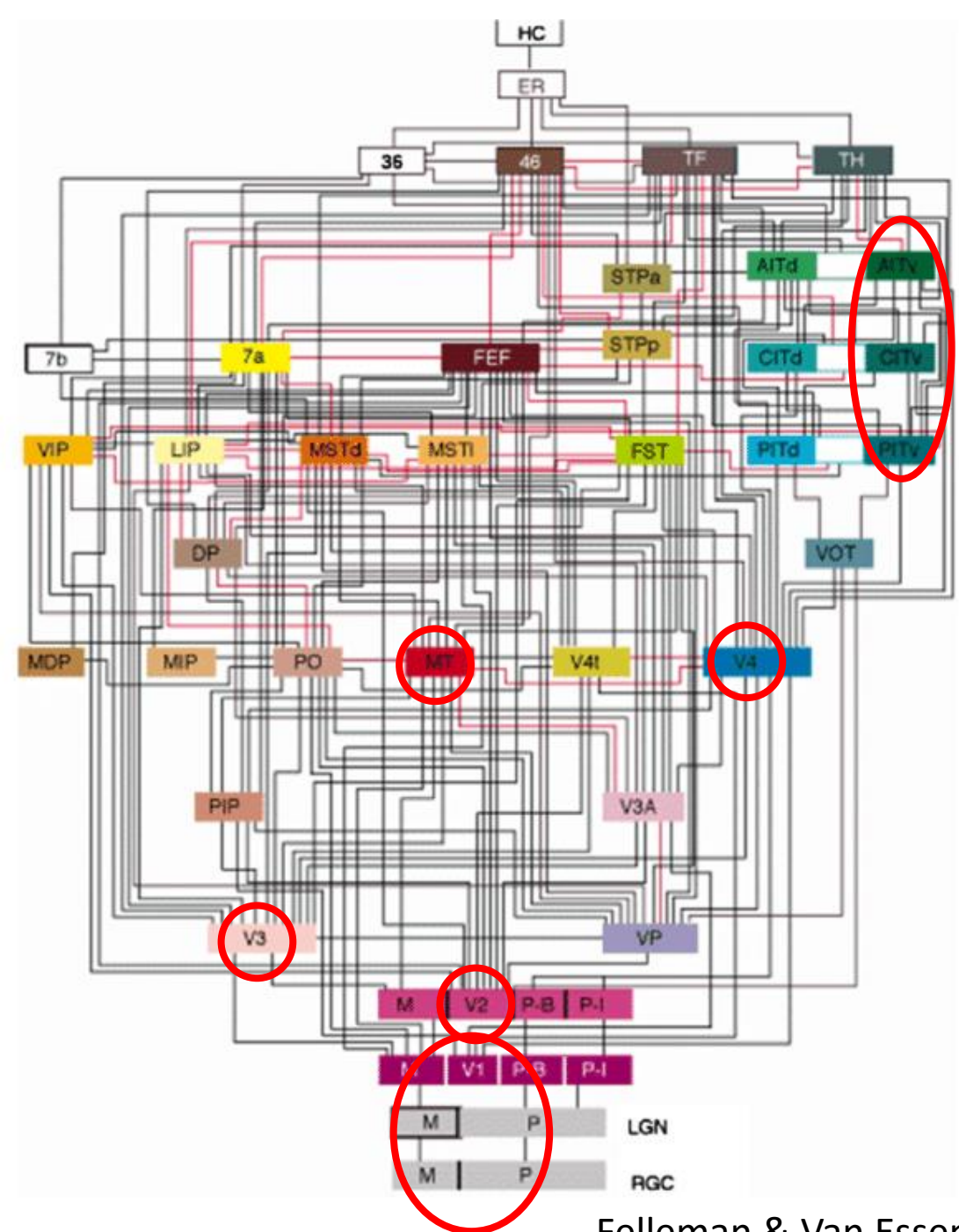
## Subcortical structures

- Inputs from retinal ganglion cells (RGC)
- Project to lateral geniculate nucleus (LGN)

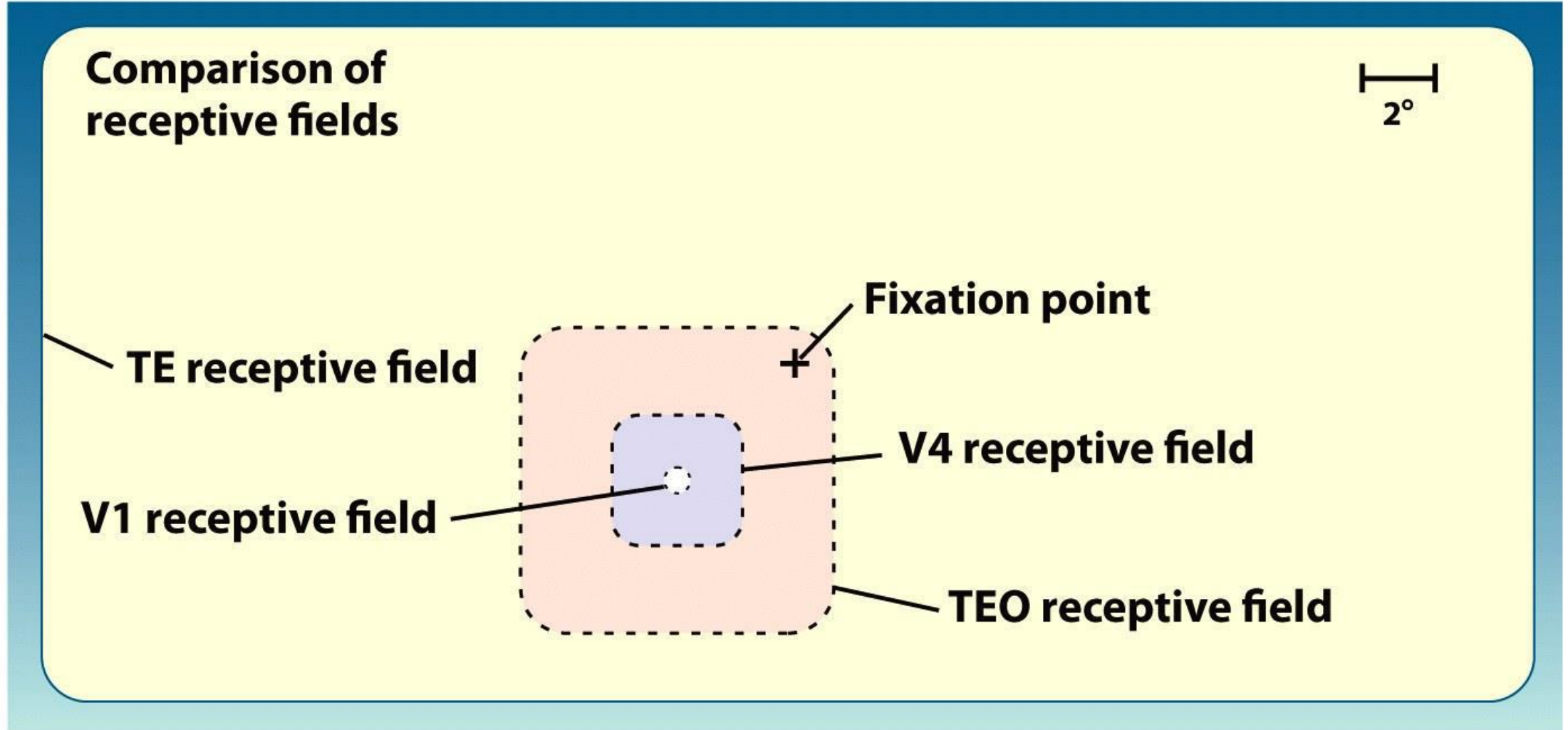
## Cortex

- To V1
- Dorsal stream
  - V3
  - MT (V5)
- Ventral stream
  - V2
  - V4
  - IT (TEO)

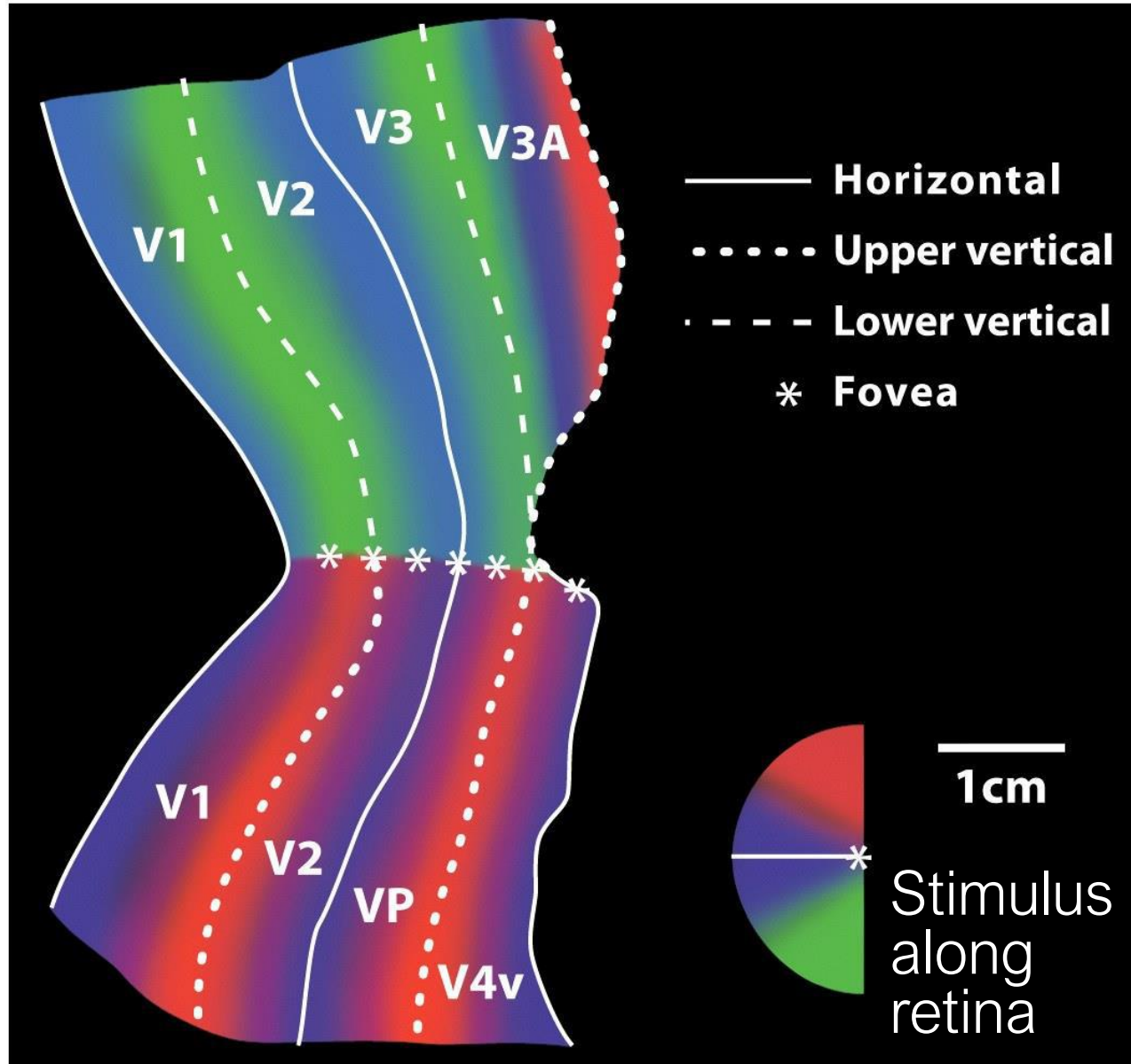
...and to a whole lot of other areas!



# Beyond V1: receptive field sizes increase along the visual hierarchy



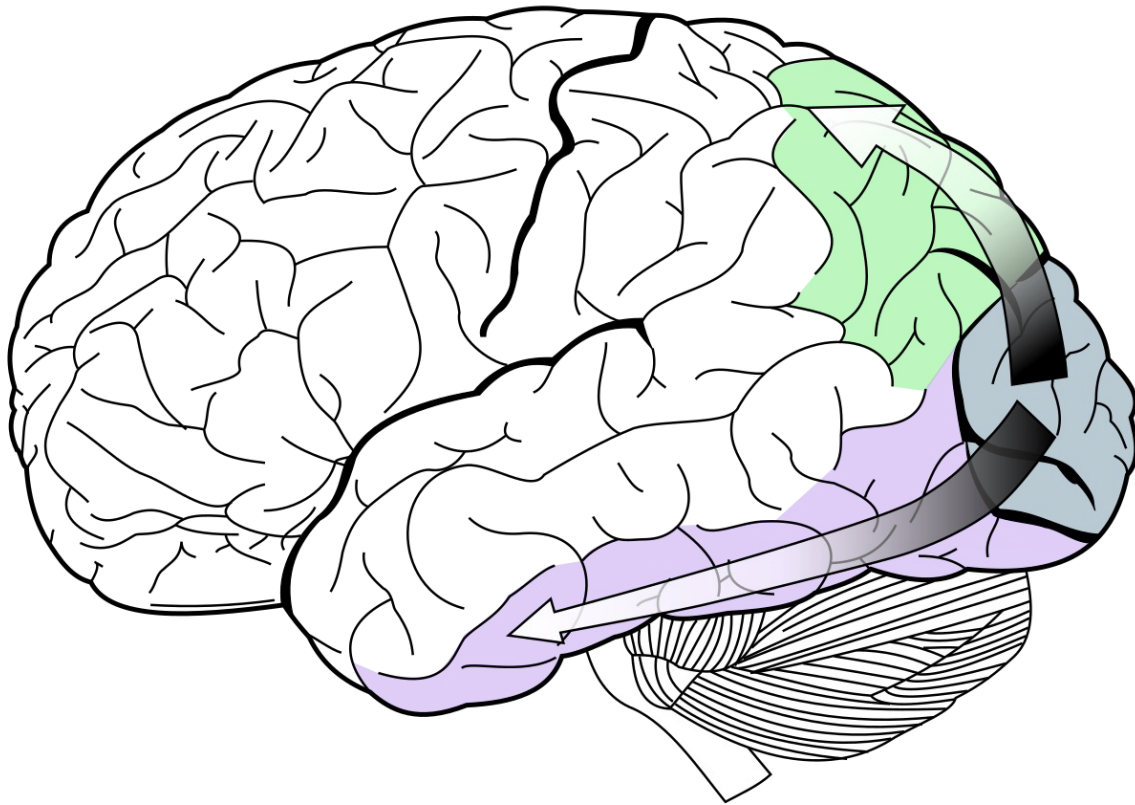
# Retinotopy



Mapping of visual input from retina to neurons

- Foveal input processed centrally
- Peripheral input along edges to subsequent areas

# Two-streams hypothesis

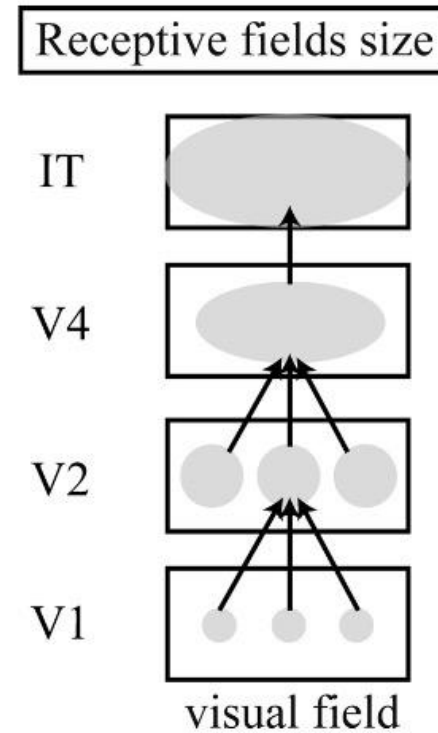
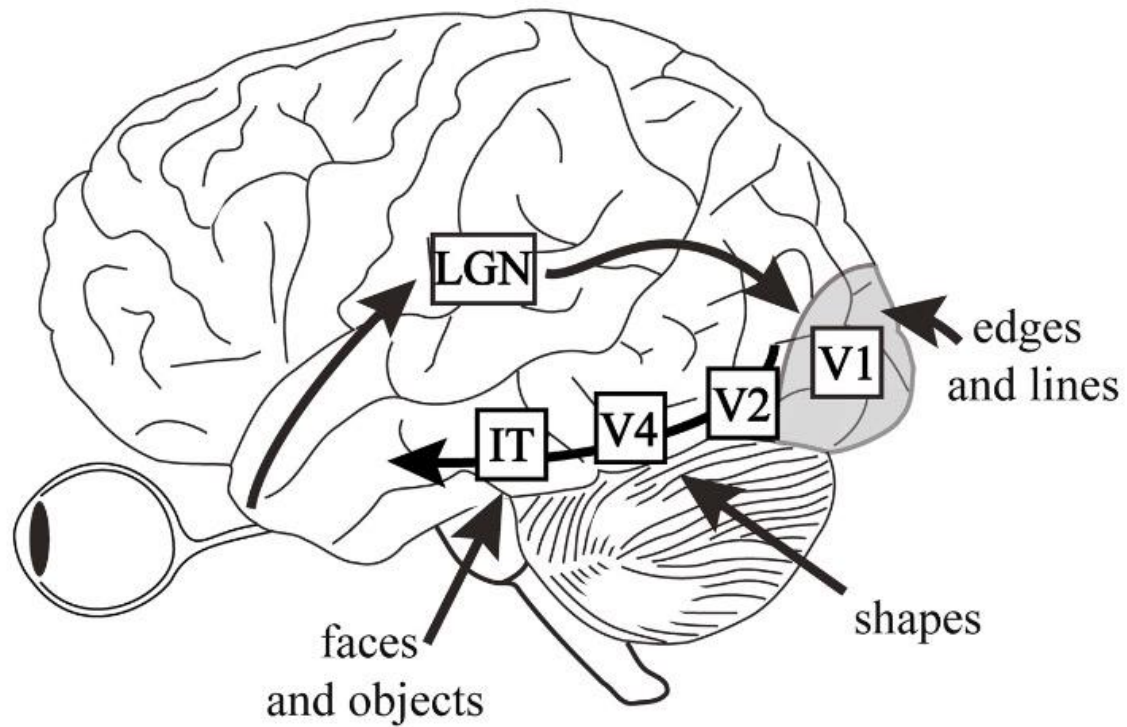


**Dorsal** pathway:  
spatial vision  
“where?”

**Ventral** pathway:  
object recognition  
“what?”



# Visual ventral stream

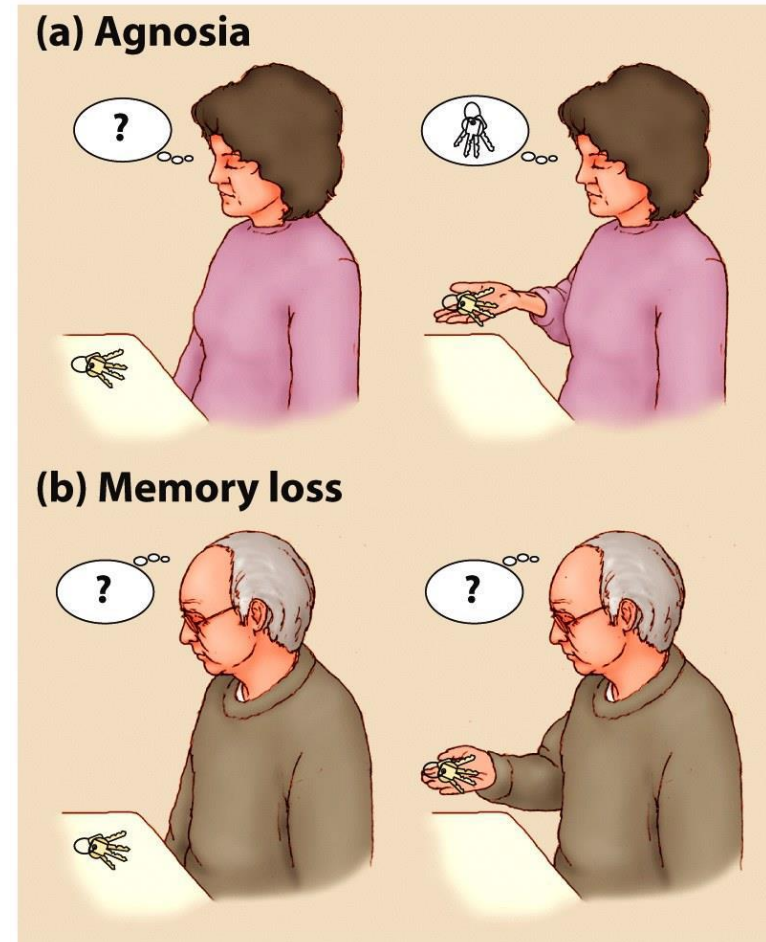


# Visual agnosia: Inability to recognize objects despite normal vision

To diagnose an agnosic disorder, it is essential to rule out general memory problems.

(a) The patient with agnosia is unable to recognize the keys by vision alone but immediately recognizes them when she picks them up.

(b) The patient with a memory disorder is unable to recognize the keys even when he picks them up.



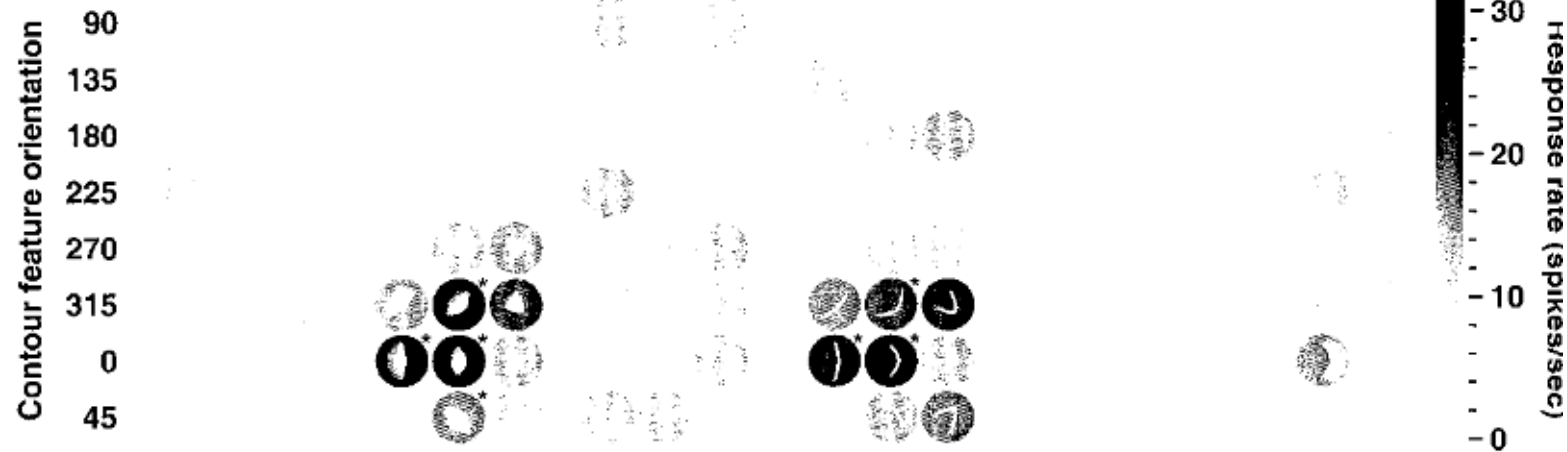
# Ventral stream: V2, V4

## V2

- RF size:  $\sim 1.4^\circ$
- Function:  $\sim$ combined edges

## V4

- RF size:  $\sim 5^\circ$
- Function:  $\sim$ curvatures



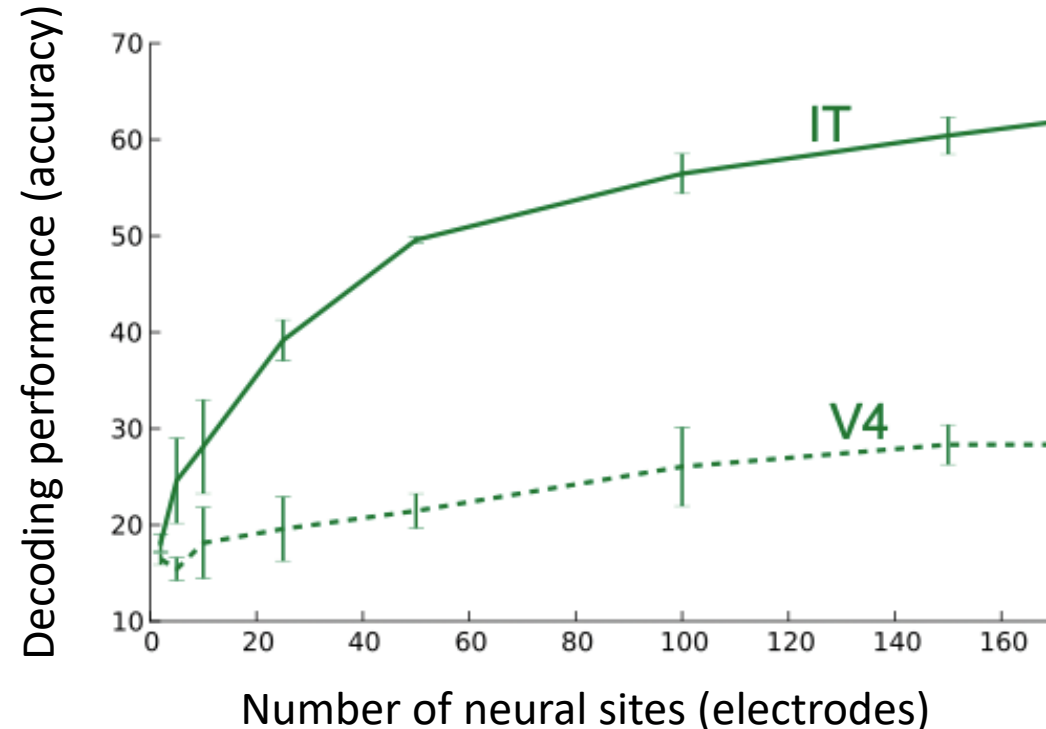
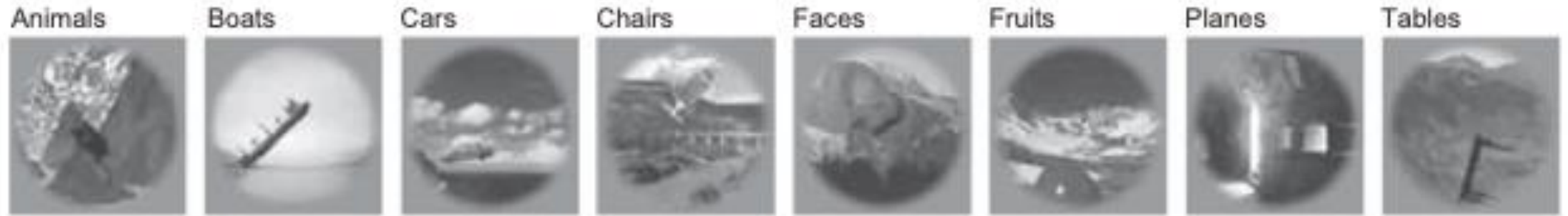
# Ventral stream: IT

- RF size:  $\sim 12^\circ$
- Function: objects
- Directly supports object recognition (correlational and causal evidence)



# IT linearly provides object category

**a** Testing image set: 8 categories, 8 objects per category



Majaj & Hong et al., JNeuroSci 2015  
Hong & Yamins et al., NatNeuro 2016

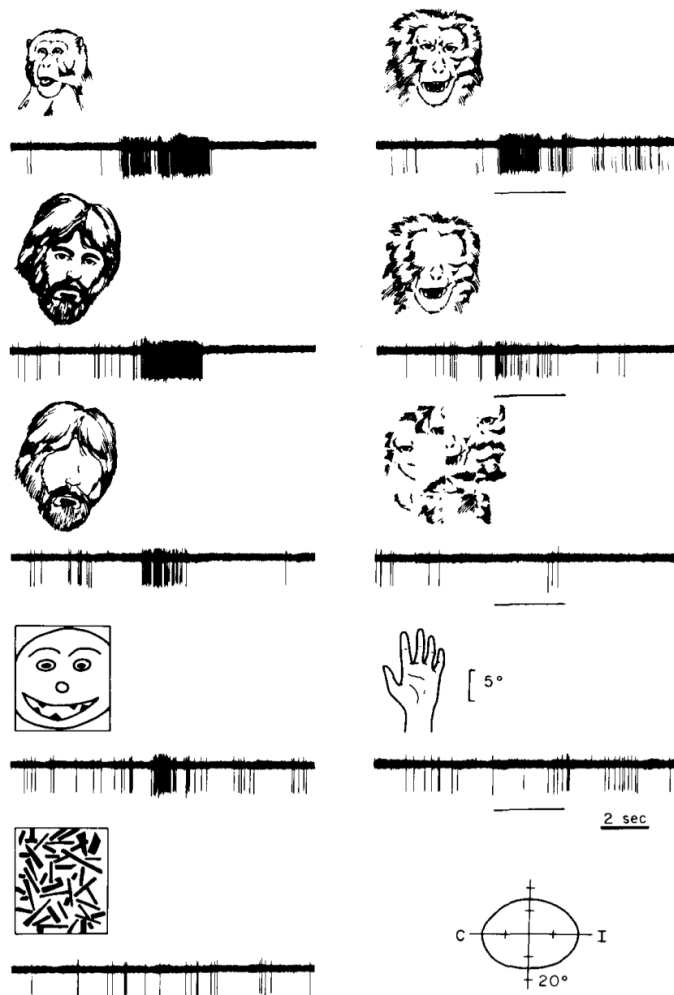
# Ventral stream: IT Face cells

Single neuron preference for faces

Face patches

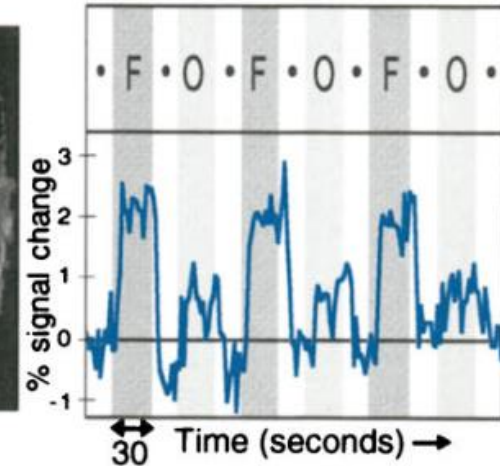
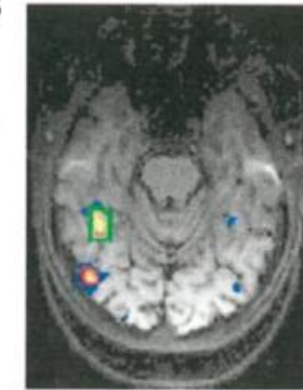
Kavli Prize in  
Neuroscience 2024

Nancy Kanwisher will  
visit EPFL on April 4!

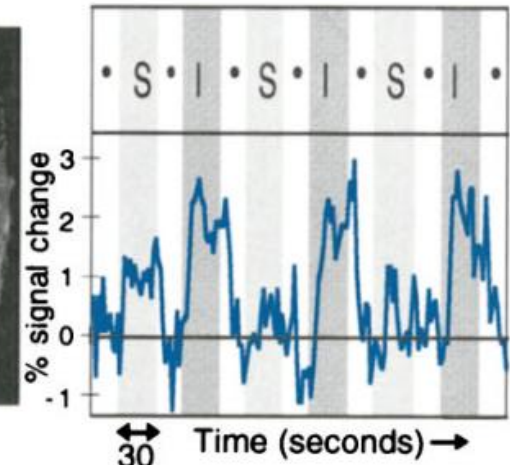
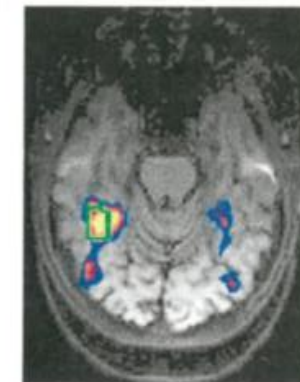


Bruce, Desimone, Gross 1981

3a. Faces > Objects



3b. Intact Faces > Scrambled Faces







Kanwisher et al. 1997

# Stimulating face-selective sites distorts face perception





# Stimulating face-selective sites *induces* face perception

				
Stim. Elec. 181-182	<i>...just for the very first second... I saw an eye, an eye, and a mouth.</i>	<i>How do I explain this? Just like the previous one, I see an eye, an eye, and a mouth, sideways.</i>	<i>Your face completely changed...I don't know what's going on. Your eyes change.</i>	<i>Hm. Am I just imagining things? Can you do it again?... OK, just as I thought, I see a face.</i>
Stim. Elec. 177-178	<i>The left side of the box looks like a rainbow.</i>	<i>If I look at the ball, the rainbow is there, wider than before, and blinking.</i>	<i>If I look at the face, this side looks like a rainbow and glowing.</i>	<i>It's kind of the same, this half is colorful.</i>

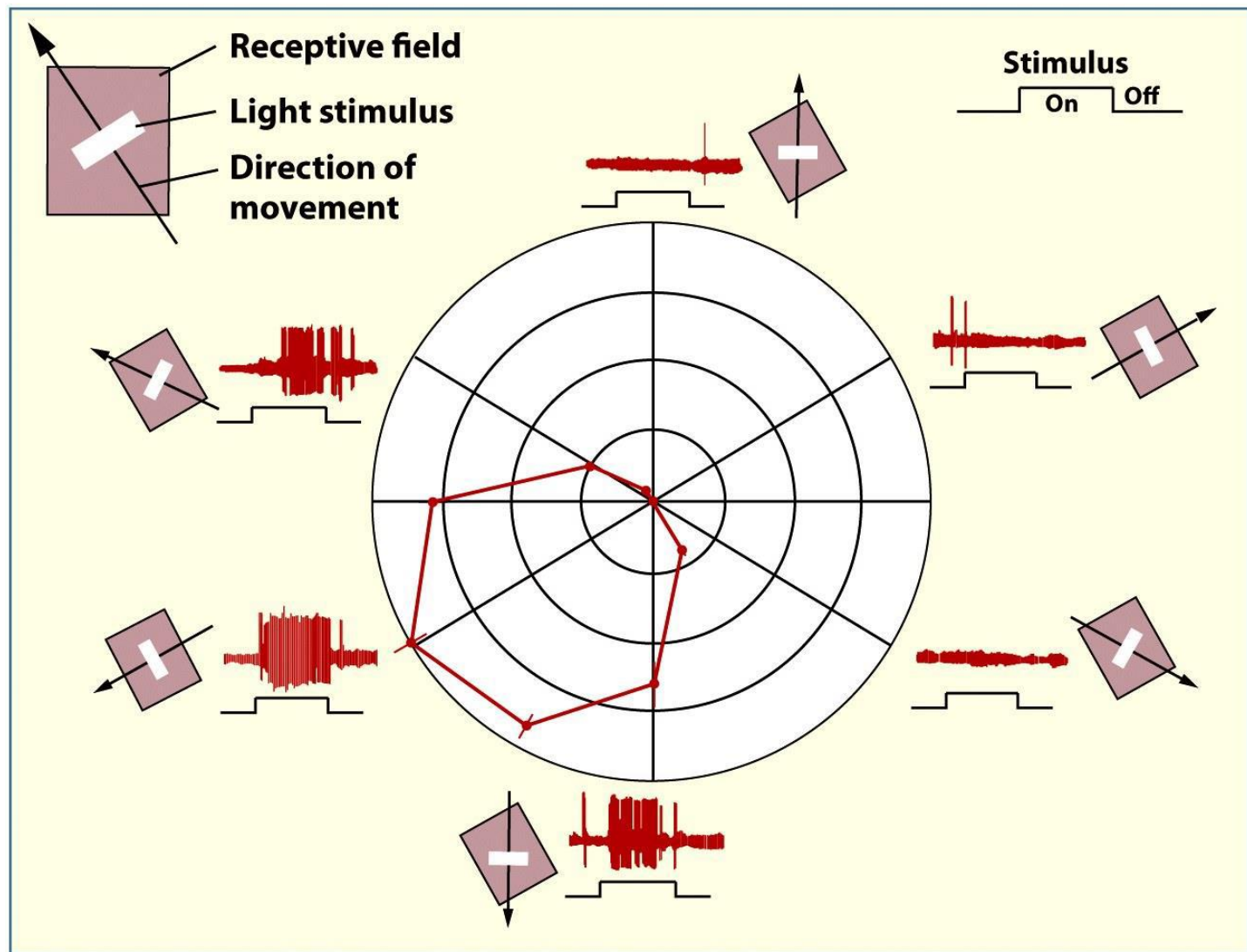
**Fig. 3.** Transcript excerpts from patient's report during electrical stimulation of electrodes 181–182 (FFA) and 177–178 (color-preferring site) while viewing a box, a ball, the experimenter's face, or a kanji character. For full transcript see [Transcript of Entire Stimulation Session](#); for excerpted videos see [Movies S1](#) and [S2](#).



# Dorsal stream: MT

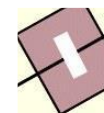
- Highly selective for motion direction and speed
- Invariant to object form or texture
- Thought to underlie motion recognition

# Dorsal stream: MT motion preference



single-neuron  
direction tuning

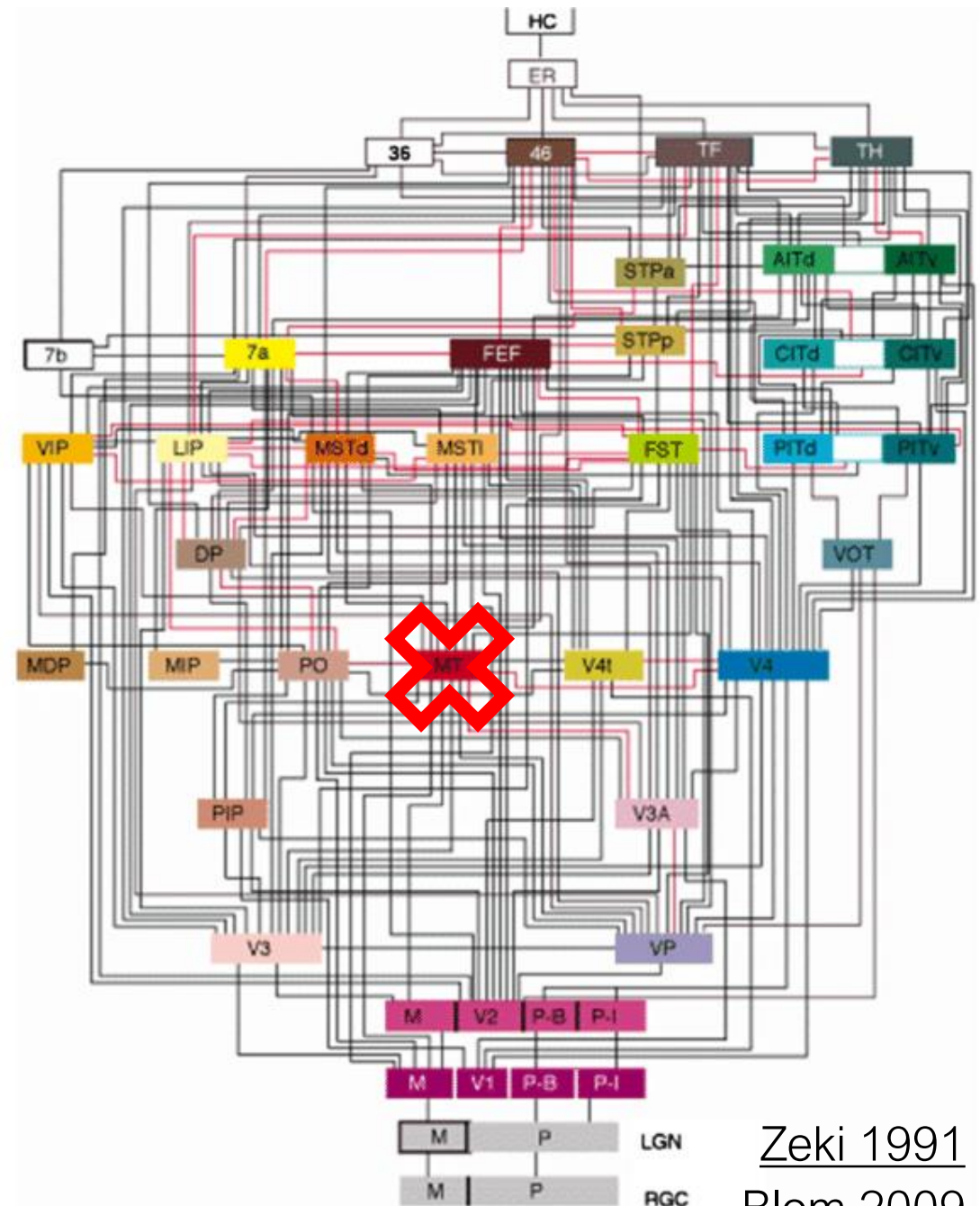
Same bar orientation  
evokes different  
responses depending  
on movement



# Akinetopsia

Damage to MT (middle temporal visual area, also called V5)

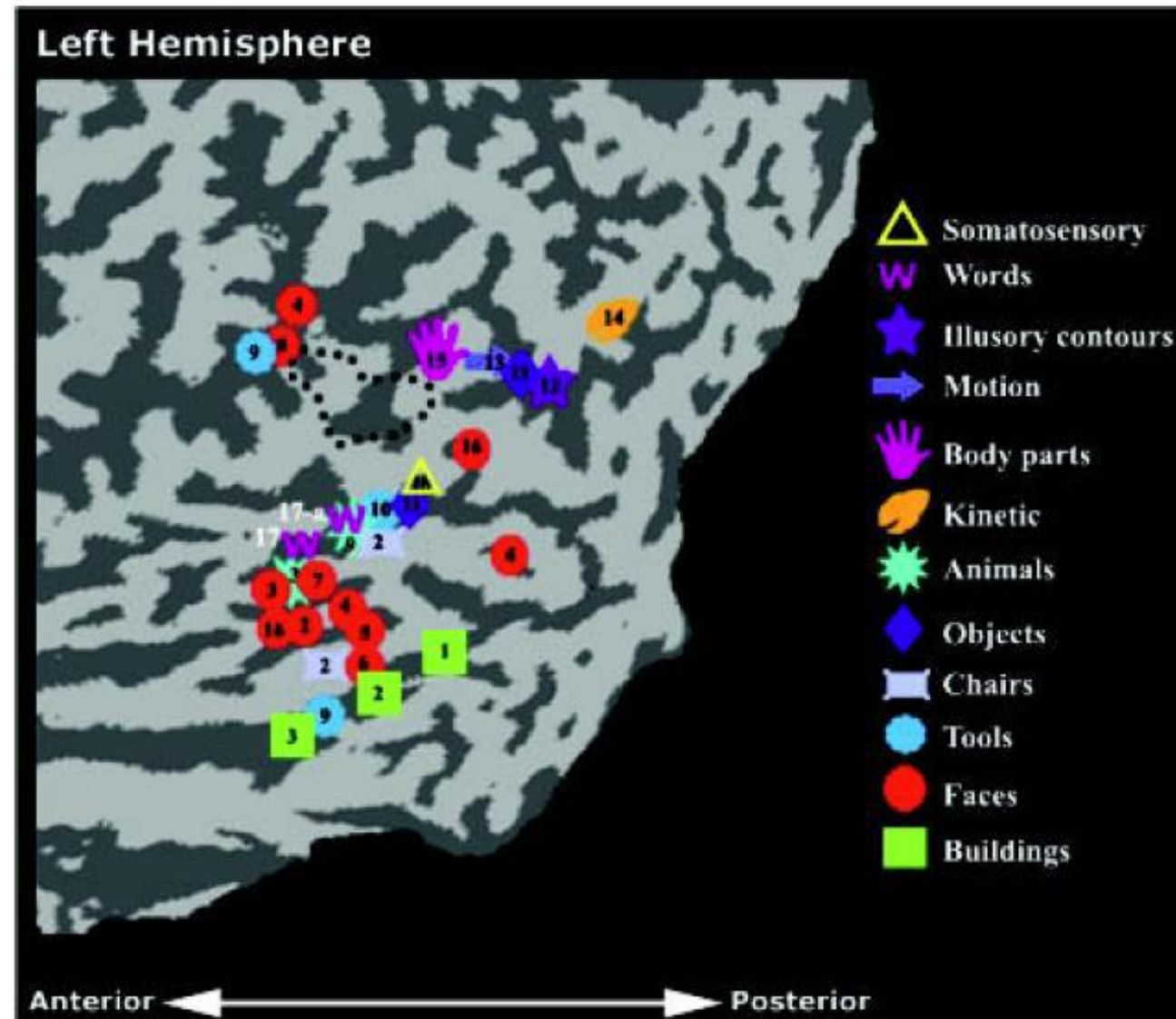
- inability to perceive motion
- but: intact ability to perceive stationary objects



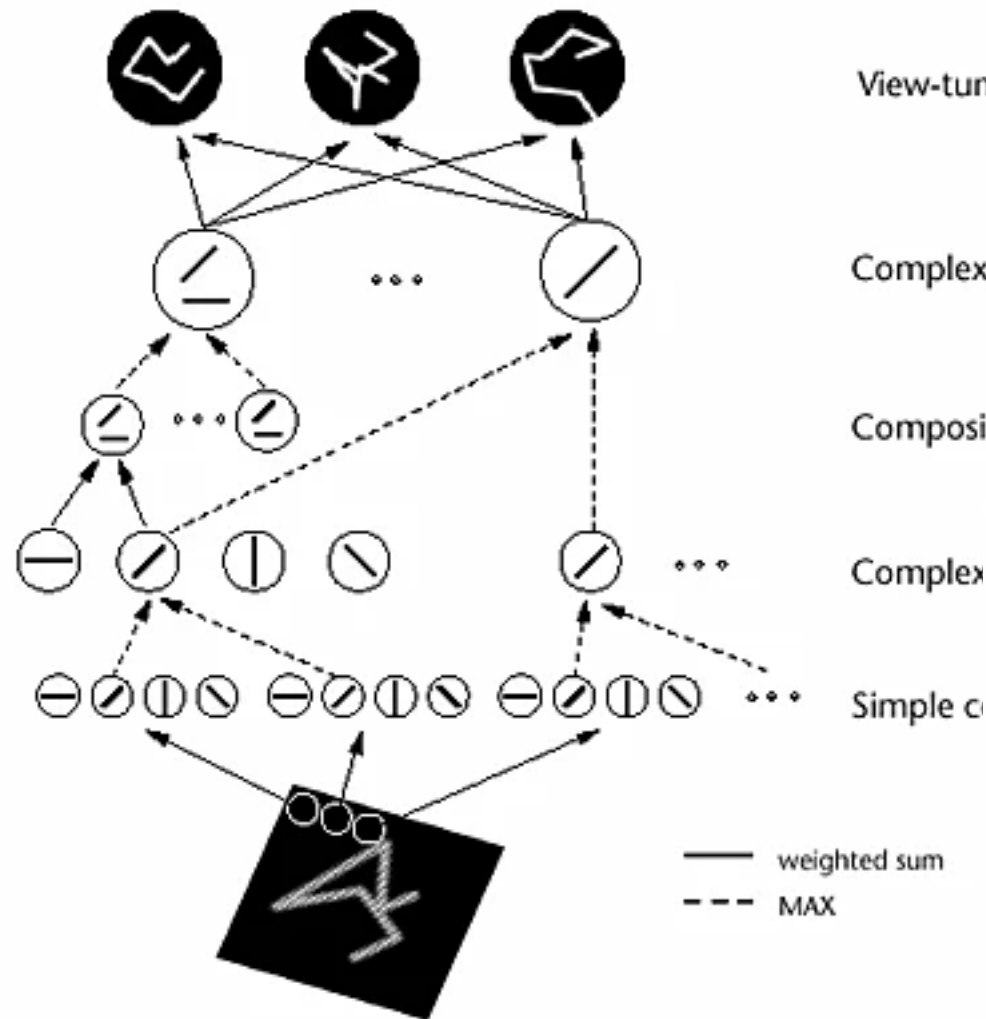
Zeki 1991

Blom 2009

# Swiss army knife of visual areas



# Computational Models of Vision: HMAX



View-tuned cells

Complex composite cells (C2)

Composite feature cells (S2)

Complex

Simple c

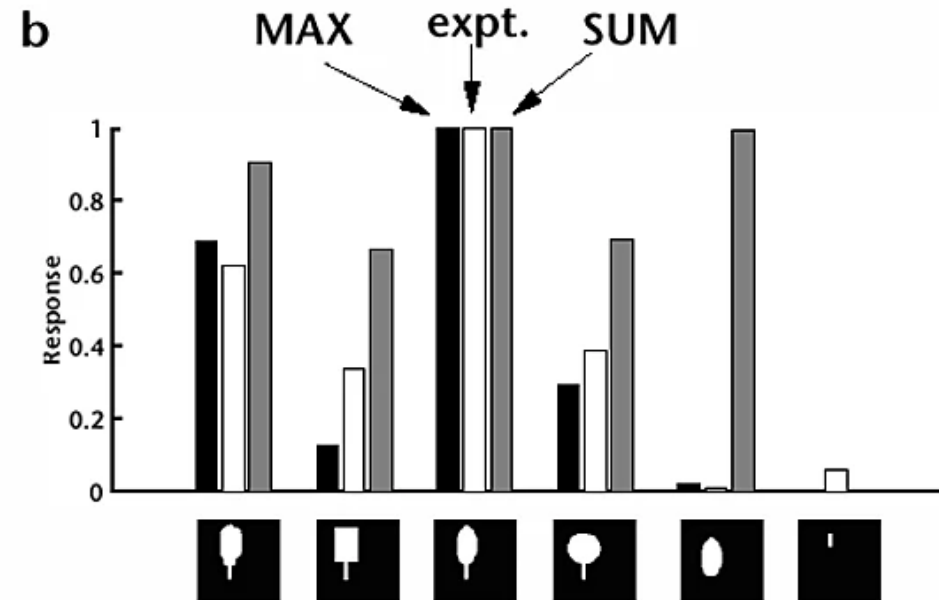
— weighted sum  
- - - MAX

Hierarchical feedforward architecture

Simple cells: linear operations

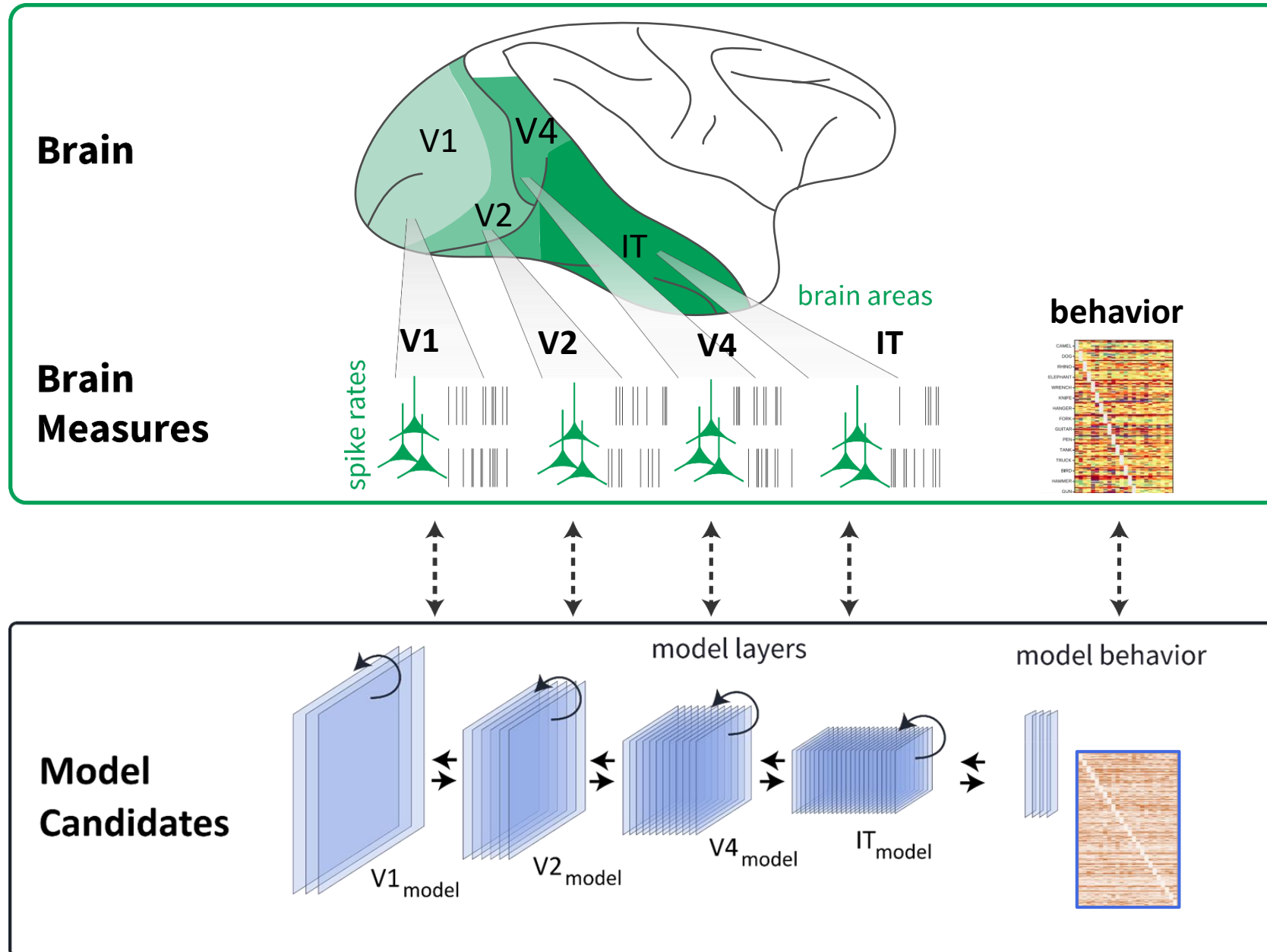
Complex cells: nonlinear pooling  
(max-pool)

Replicates higher-order effects in IT





# Computational Models of Vision: Deep Nets



# Vision conditions

Achromatopsia: see world devoid of color



Synesthesia: associate letters with numbers



Prosopagnosia: inability to recognize faces



# Prosopagnosia

Which is easier?



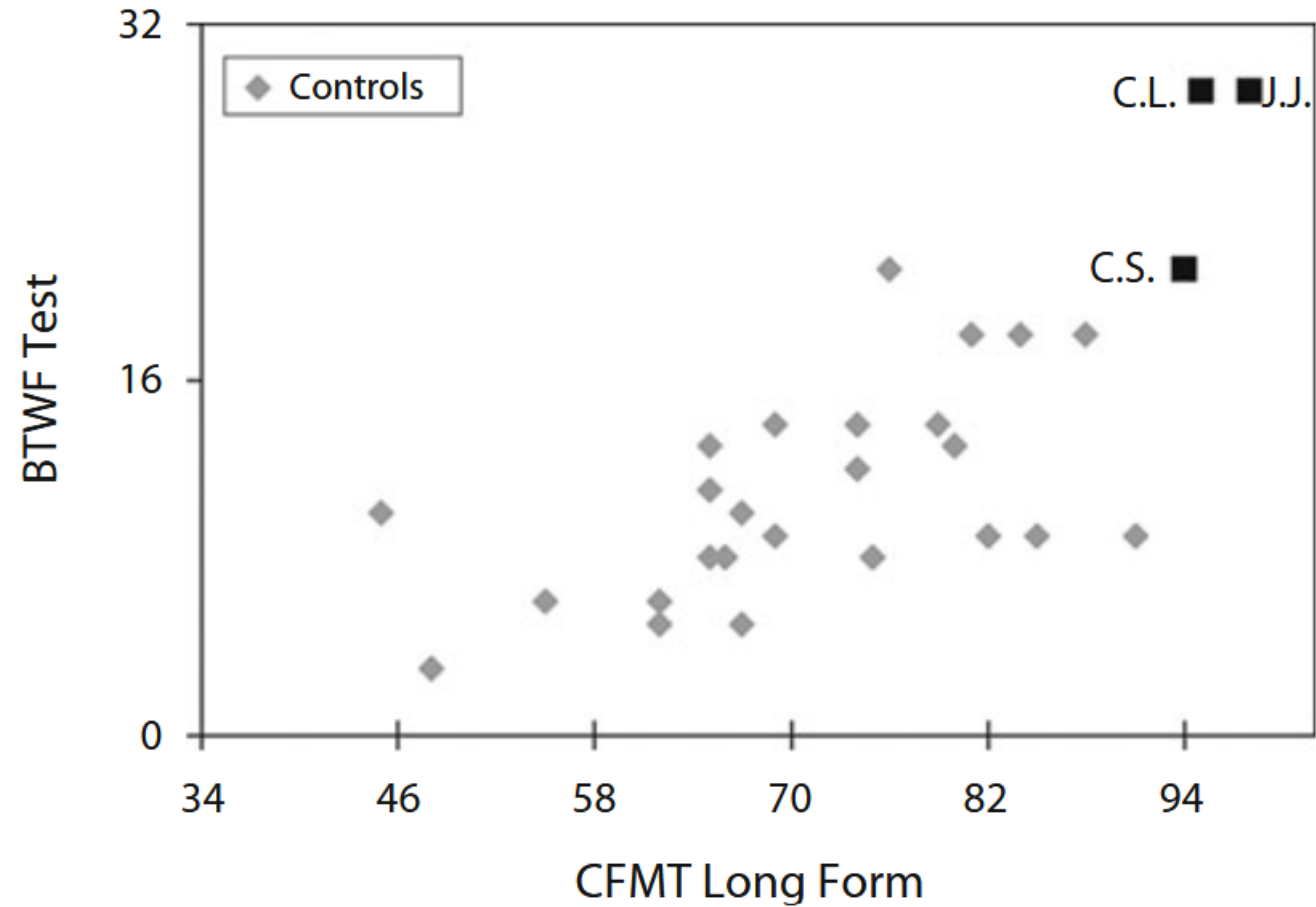
Which is easier?



Neurotypical individuals find it easier to recognize faces in their usual orientation. People with prosopagnosia have no difference in performance.

# Super-recognizers

- Extraordinary face recognition ability
- Opposite of prosopagnosia
- Improved face matching performance compared to average population



# Questions

- Name three structures involved in vision in the brain
- What is a receptive field?
- What types of stimuli do neurons in V1 respond to?
- How does the Gabor model relate to V1?
- What two pathways does V1 feed into and what are thought to be their key functions?
- How are IT and MT different from one another?