

Sensory Systems and Perception: Vision

Week 5

Martin Schrimpf

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

Learning Objectives – Week 5

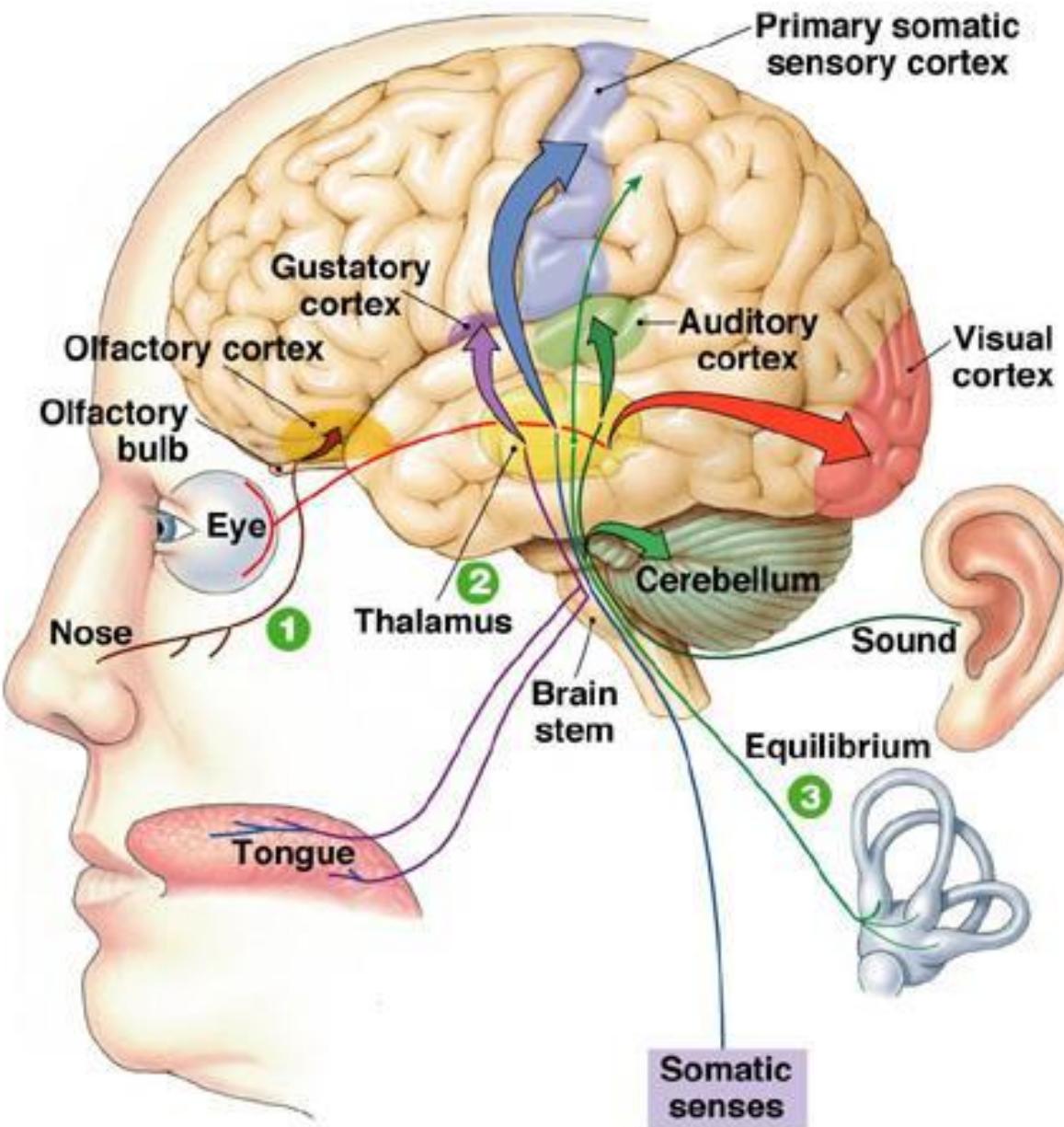
Describe the key components involved in vision.

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.

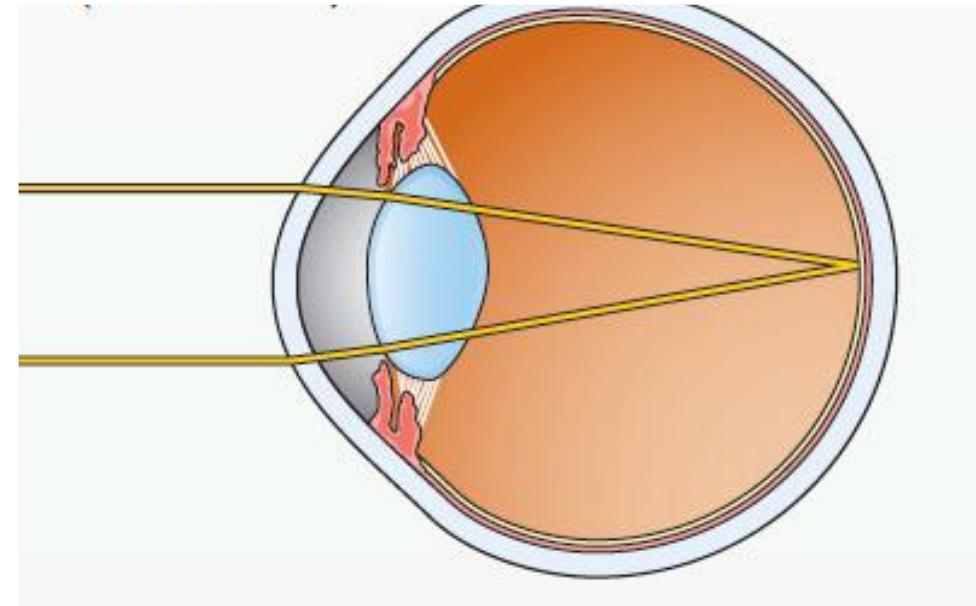
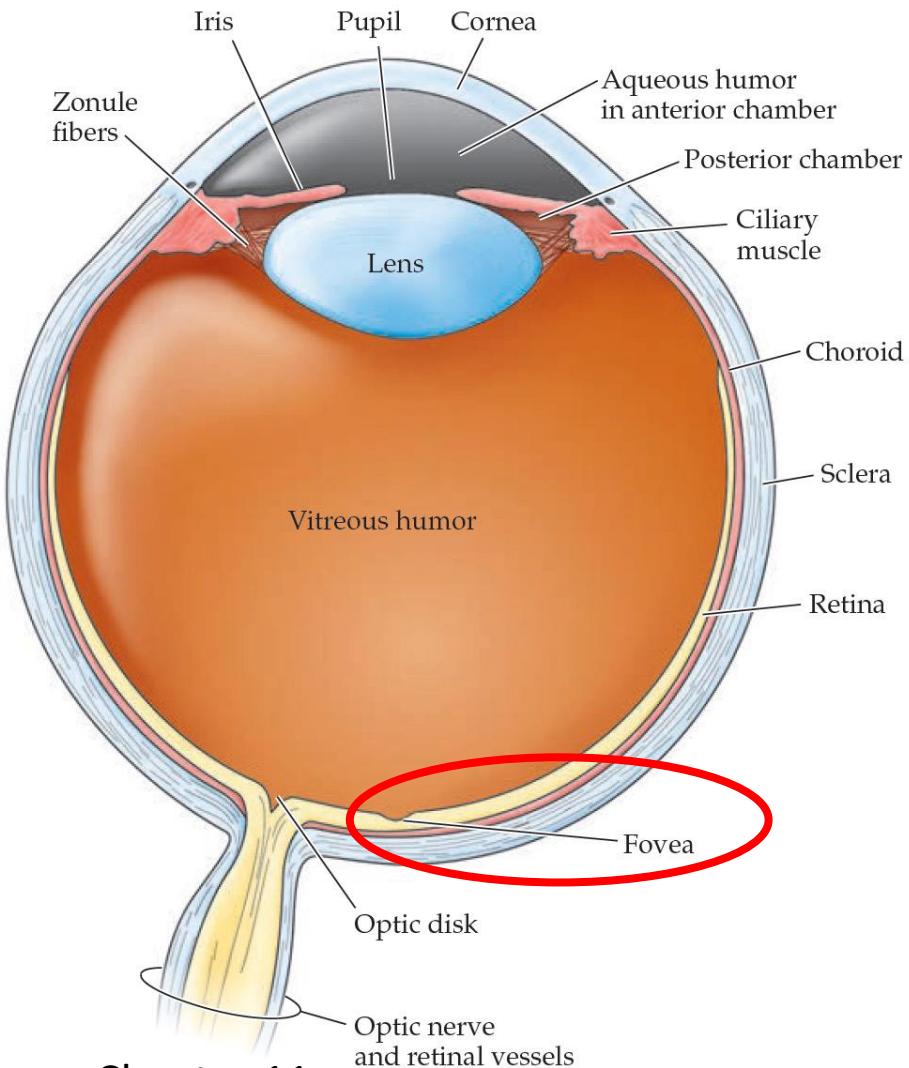


Copyright © 2007 Pearson Education, Inc., publishing as Benjamin Cummings.

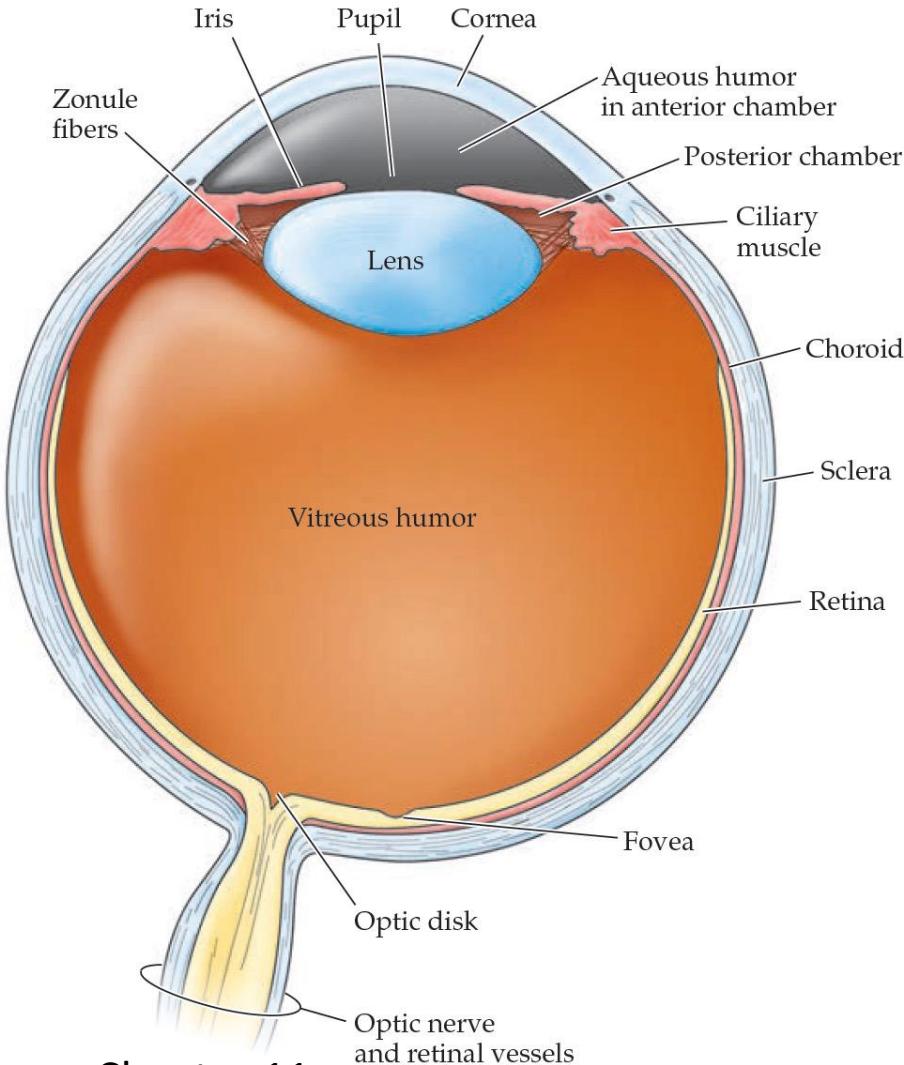
Fig. 10-4

Courtesy of Bob Desimone's class MIT 9.017

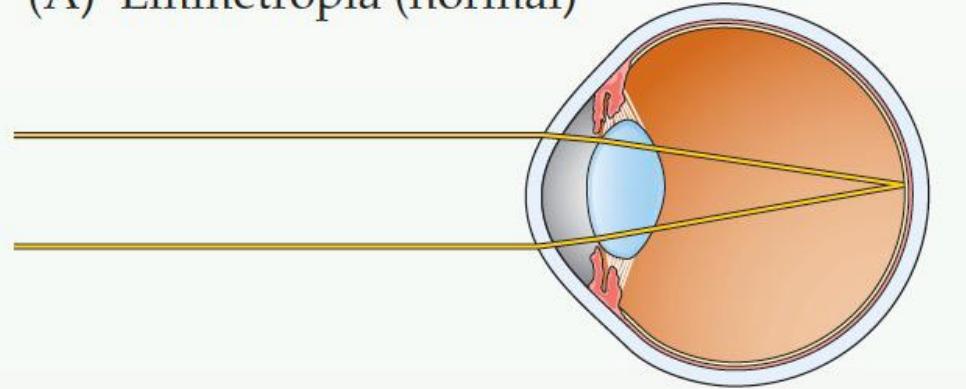
Retina



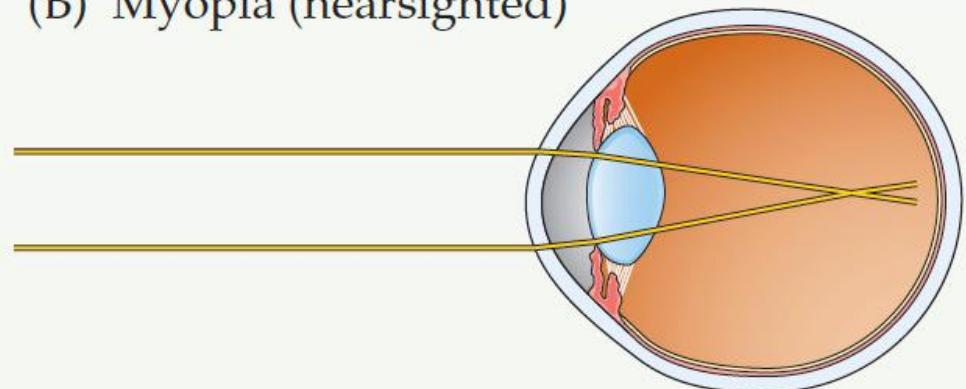
Retina



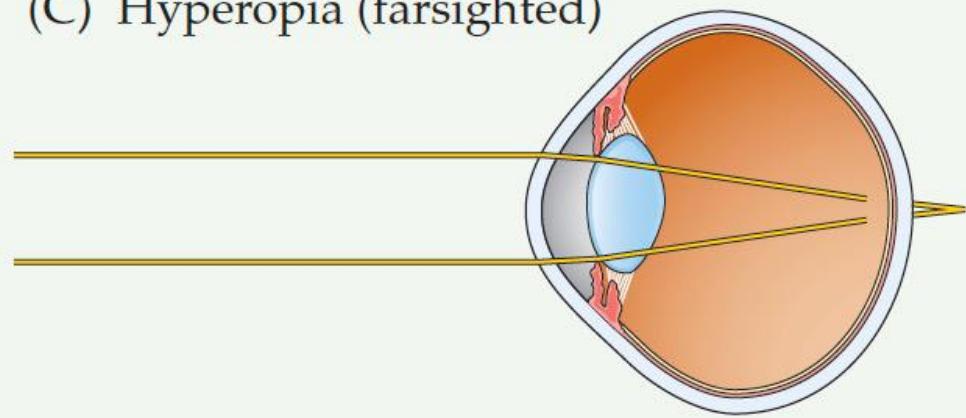
(A) Emmetropia (normal)



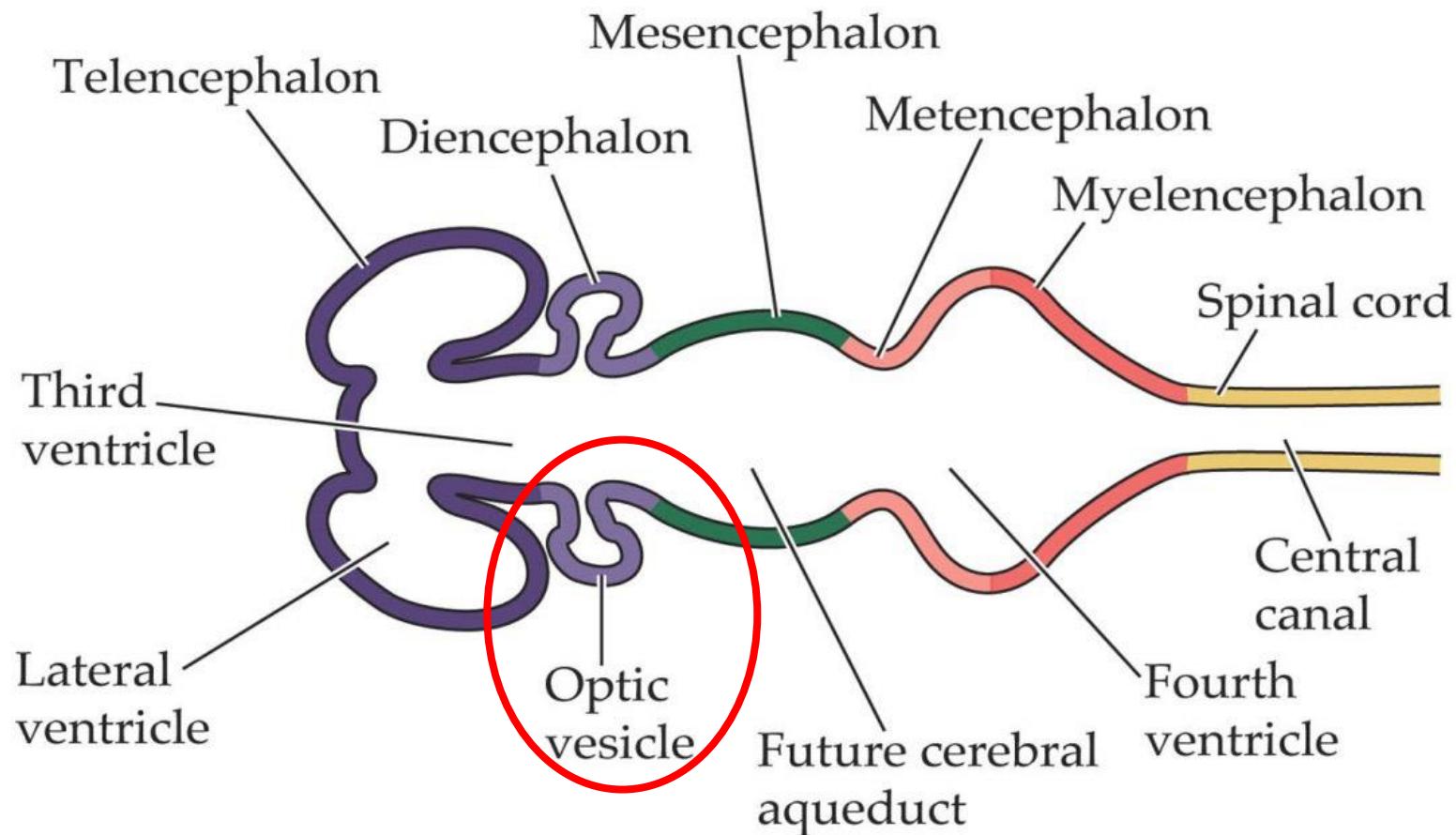
(B) Myopia (nearsighted)



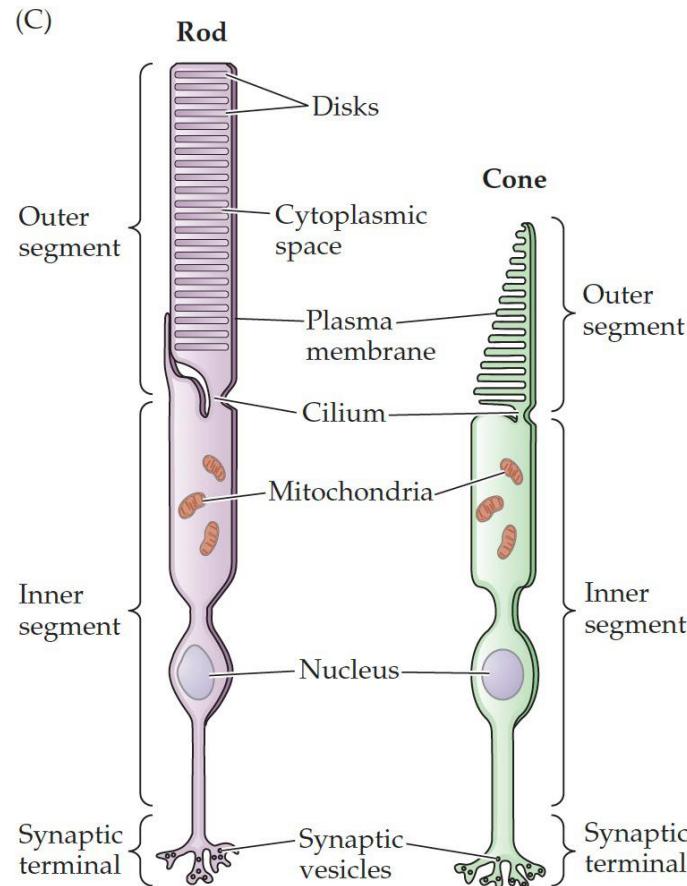
(C) Hyperopia (farsighted)



Retina is part of central nervous system



Photoreceptor cells: rods and cones



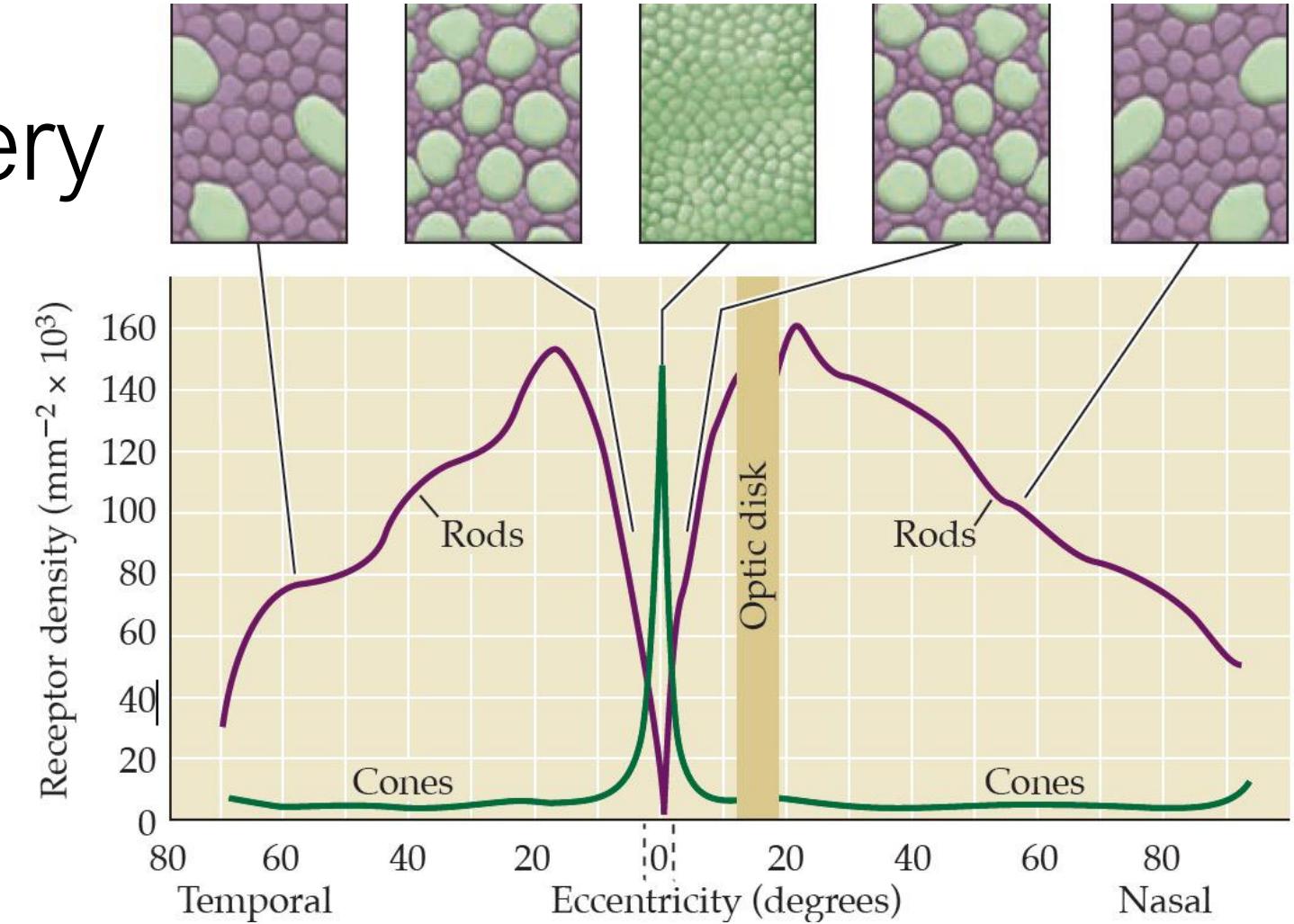
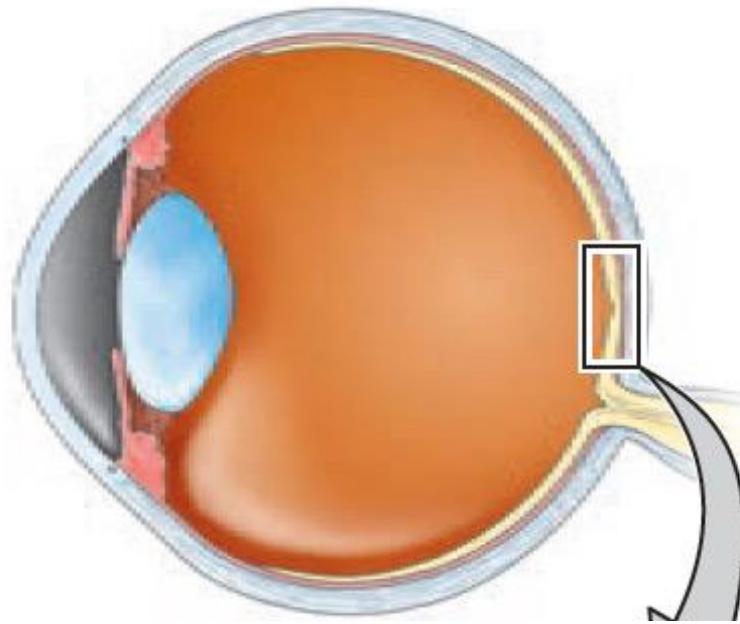
Rods:

- Low spatial resolution
- High light sensitivity

Cones:

- High acuity (spatial resolution)
- Less light sensitive

Fovea vs periphery

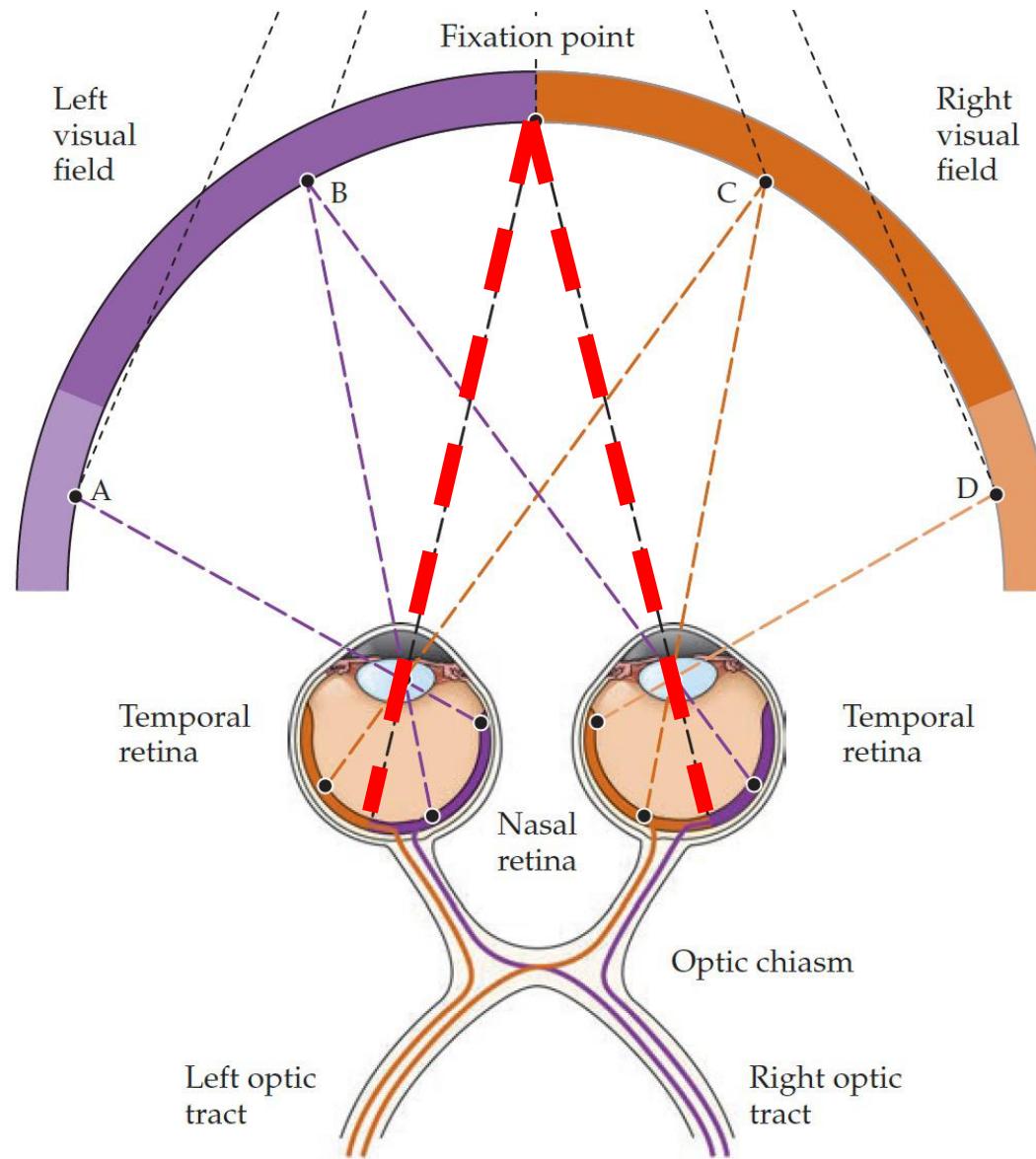


Fovea: high density of cones → better resolution

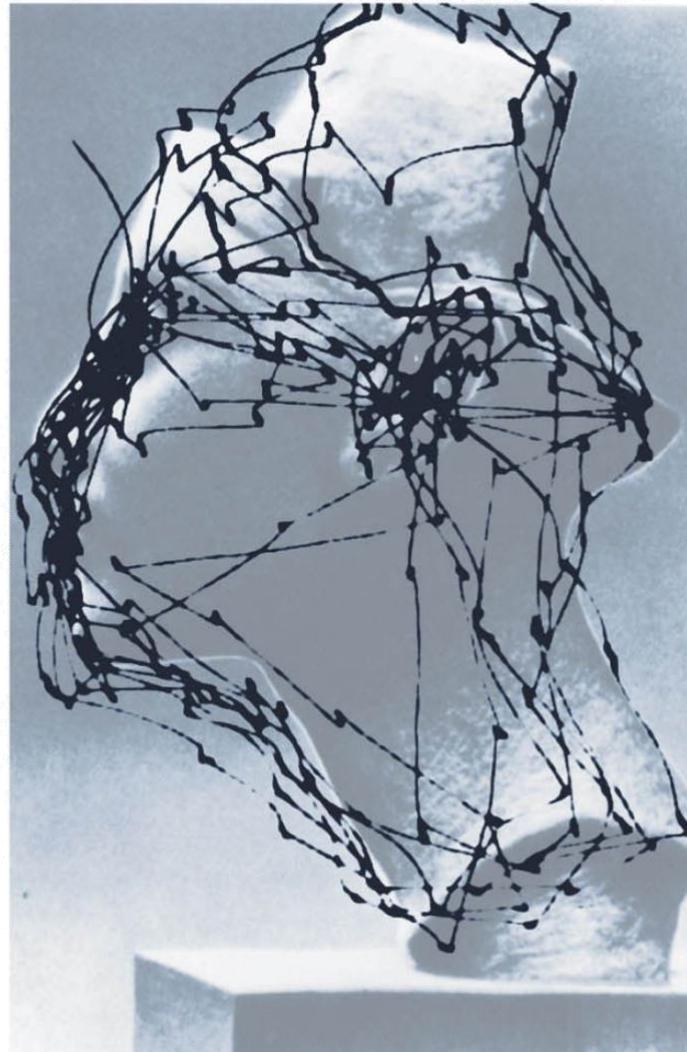
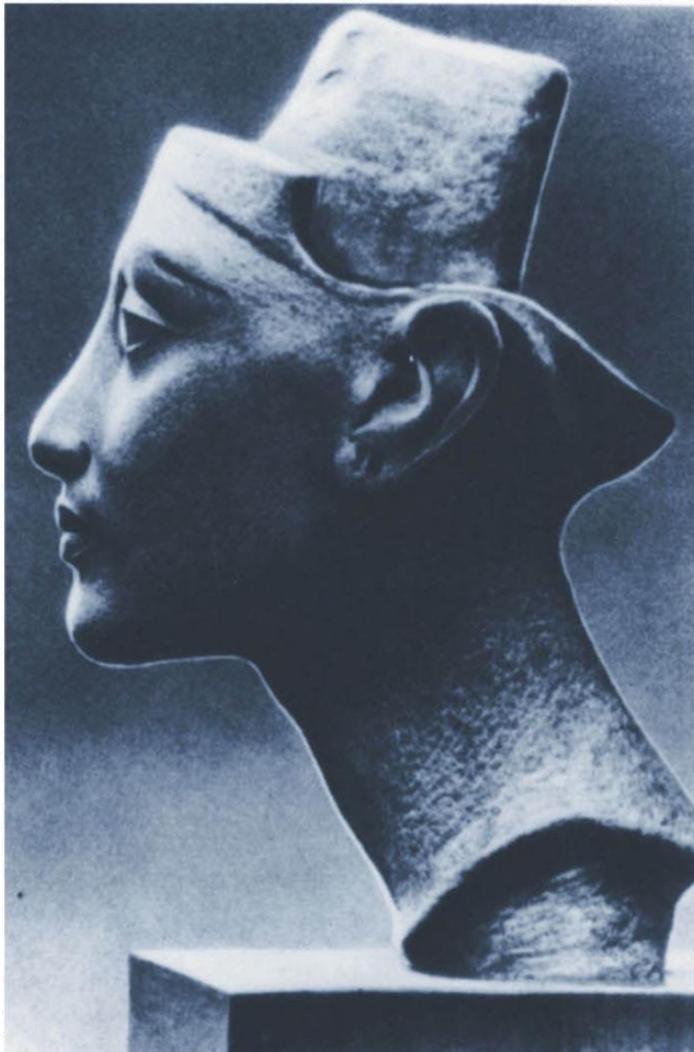
Periphery: rods > cones → worse resolution, better night vision

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:

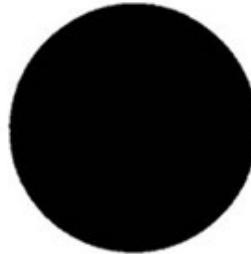
- Saccade to new point
- That part of the image can now be processed with high resolution (fovea)
- Repeat

Blind spot

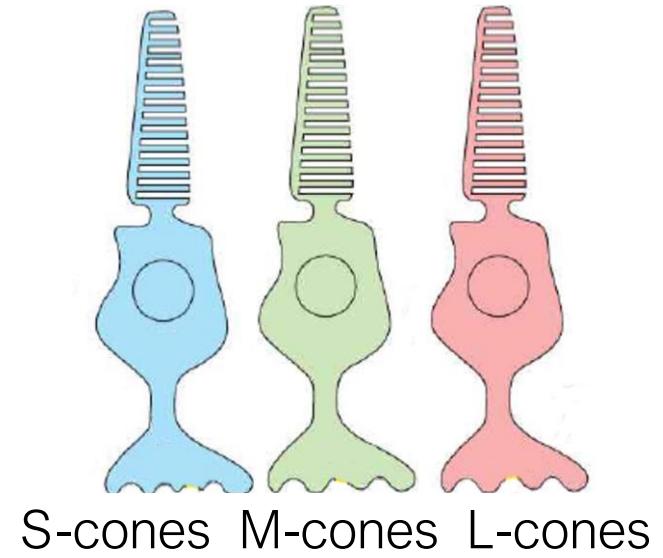
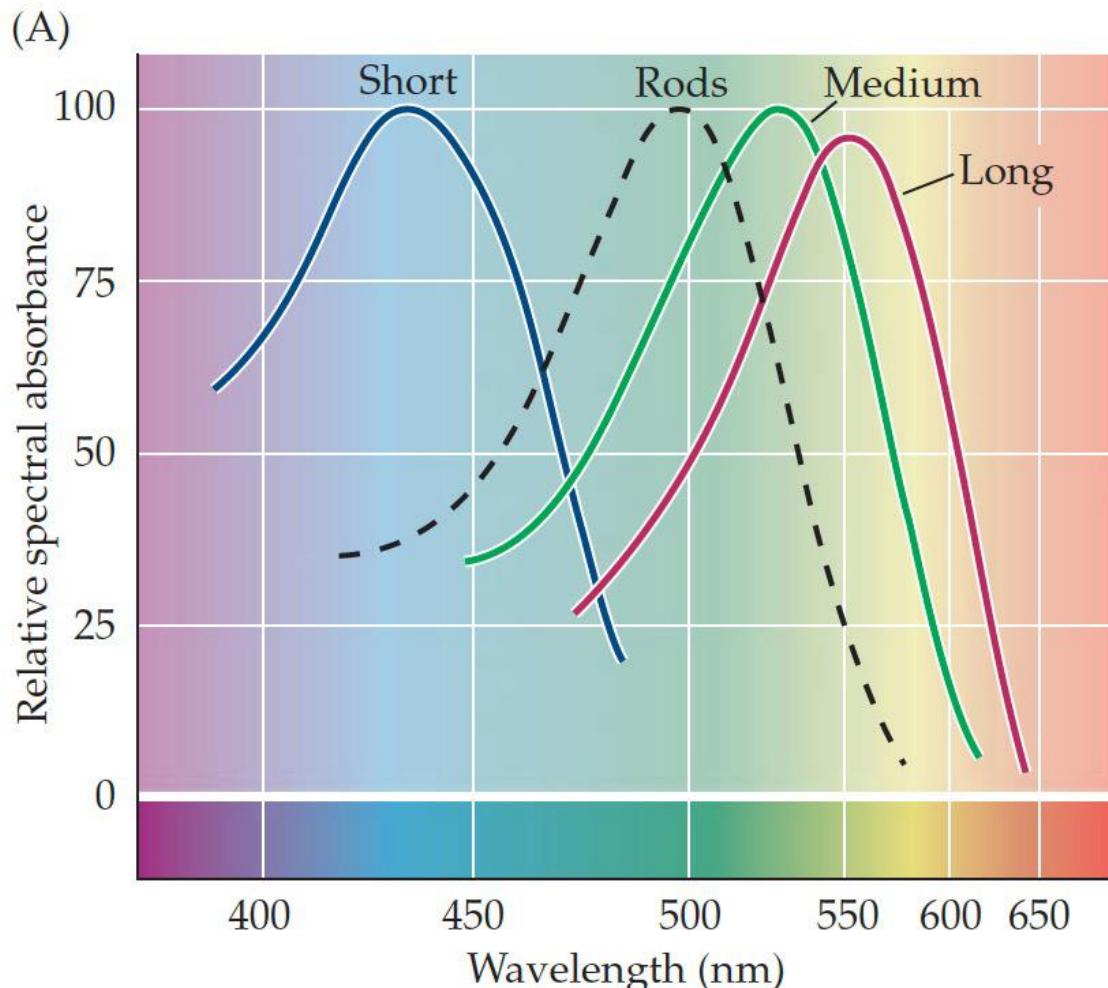
A demonstration of the blind spot:

Courtesy of Pavan Ramdy'a'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!)



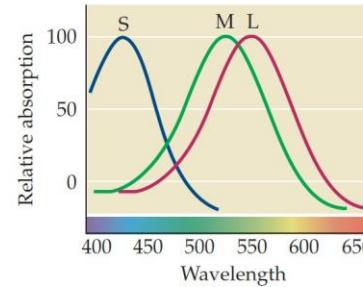
Color vision is realized by 3 types of cones with different spectral sensitivities



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

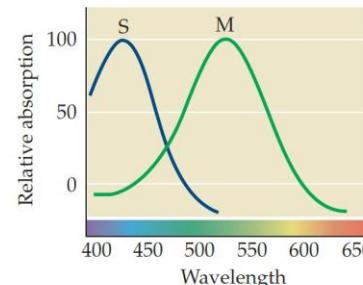
Color blindness (dichromacy)

(A) Normal (trichromat)



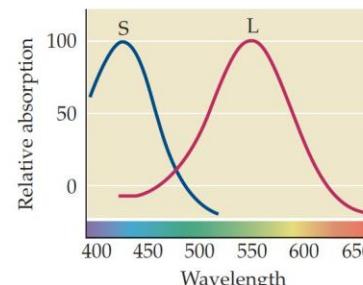
Neurotypical

(B) Protanopia



Loss of L-Cones → Red-Green color blind

(C) Deutanopia

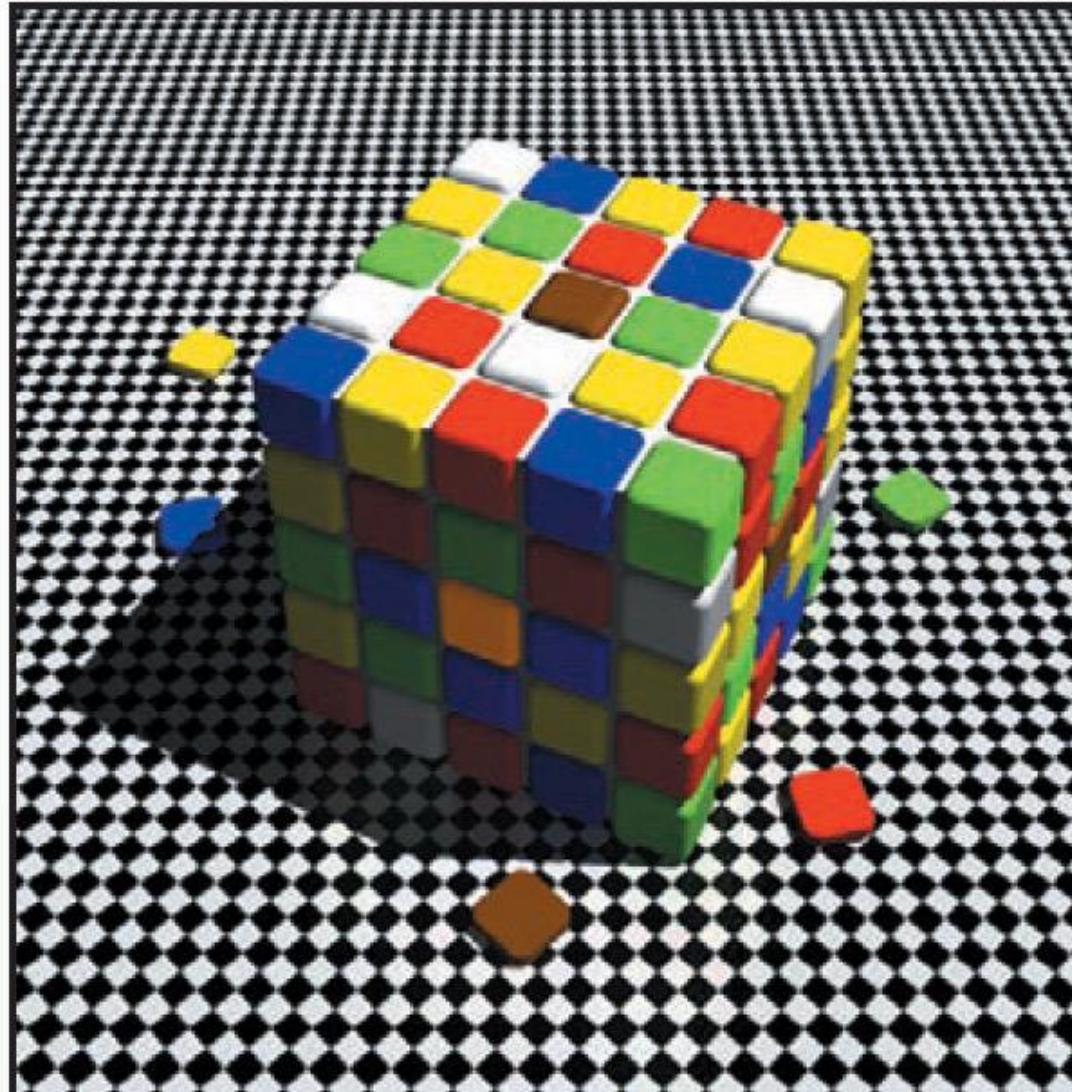


Loss of M-Cones → Red-Green color blind

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

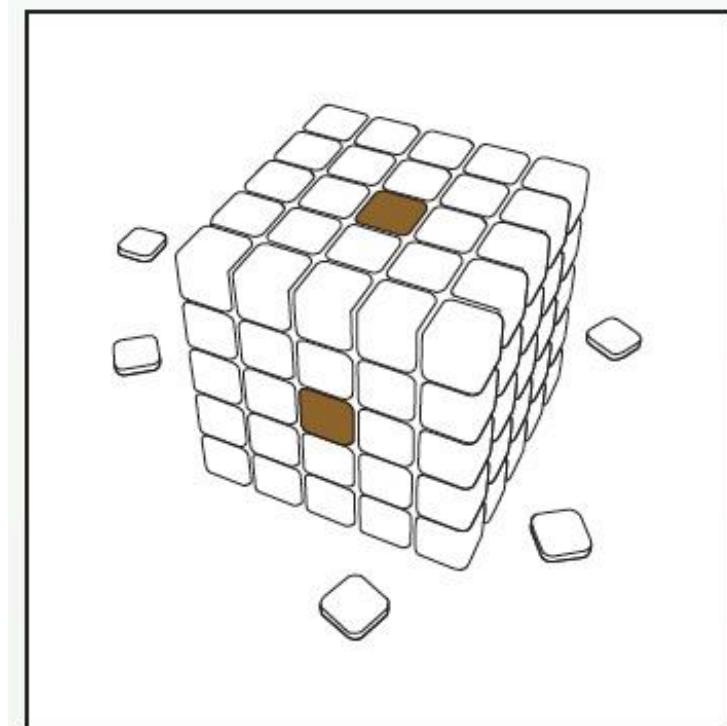
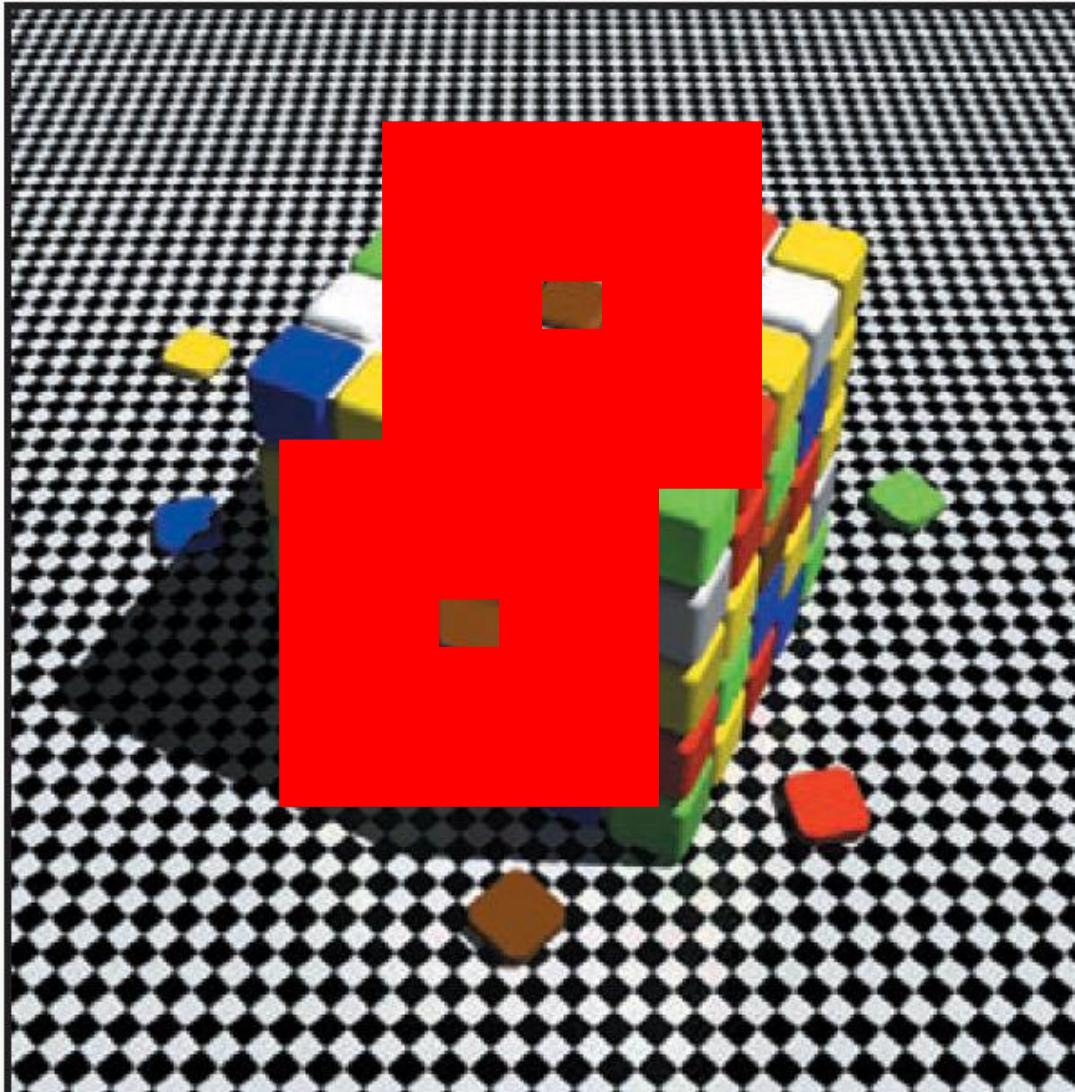
Visual Context

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?



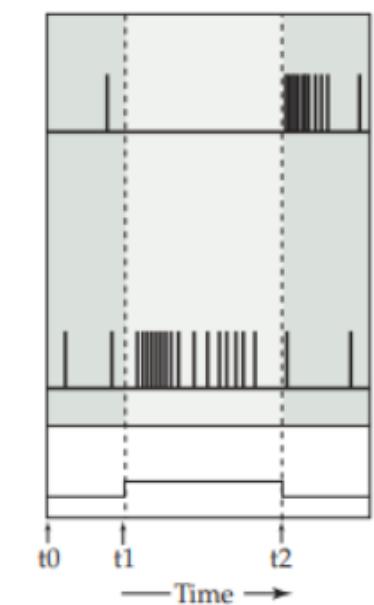
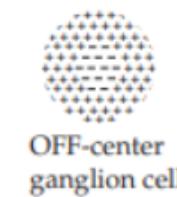
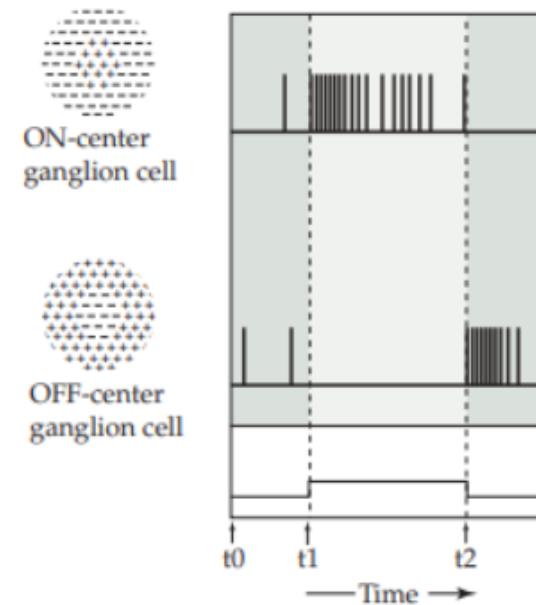
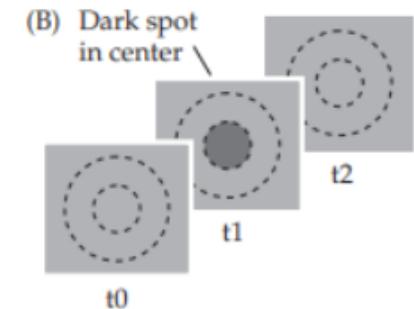
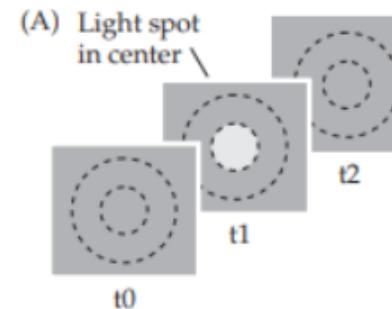
A simple form of context in retinal ganglion cells: ON- and OFF-center

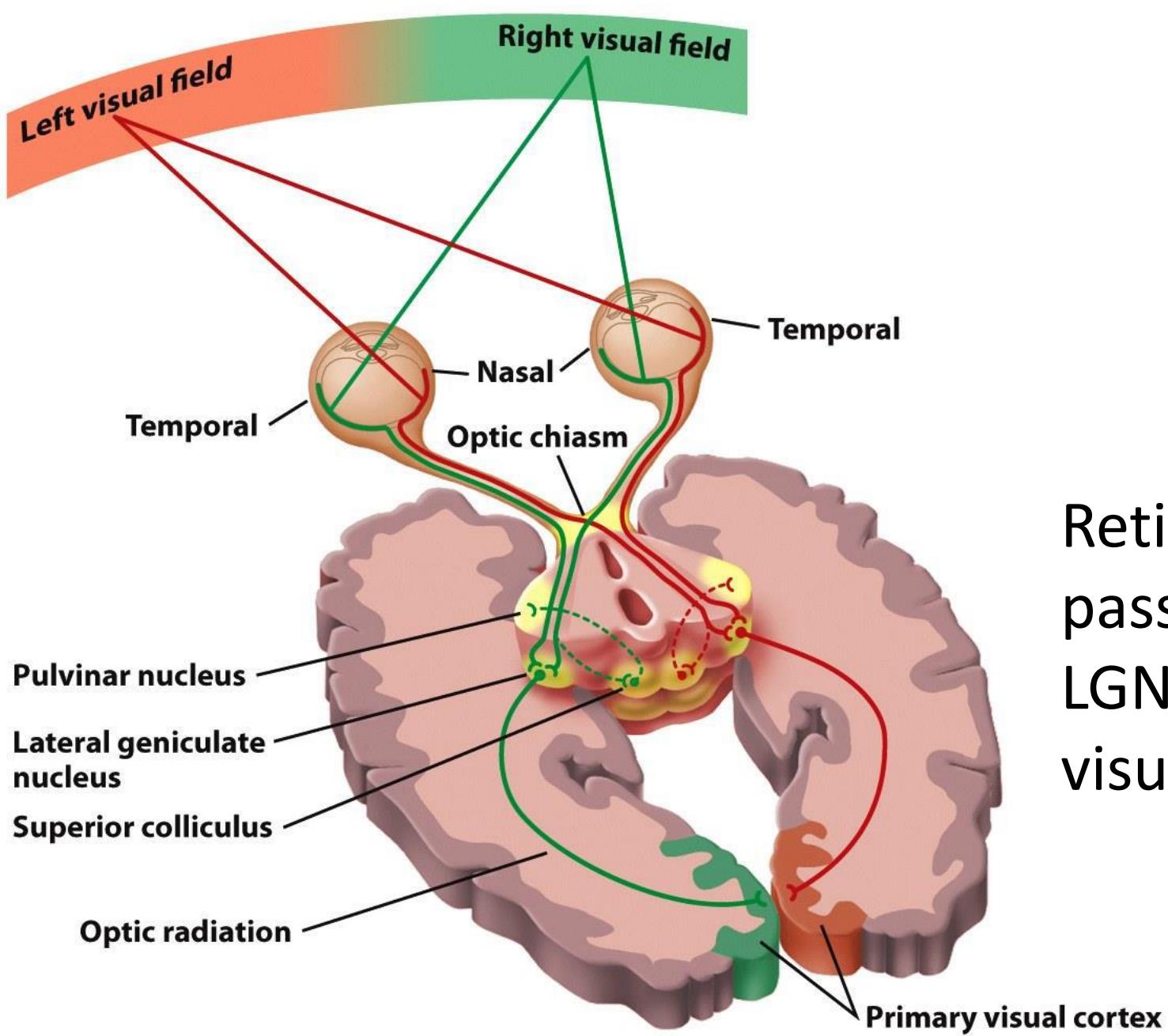


"ON-center":
↑ AP- frequency
in the center of the receptive field (RF)

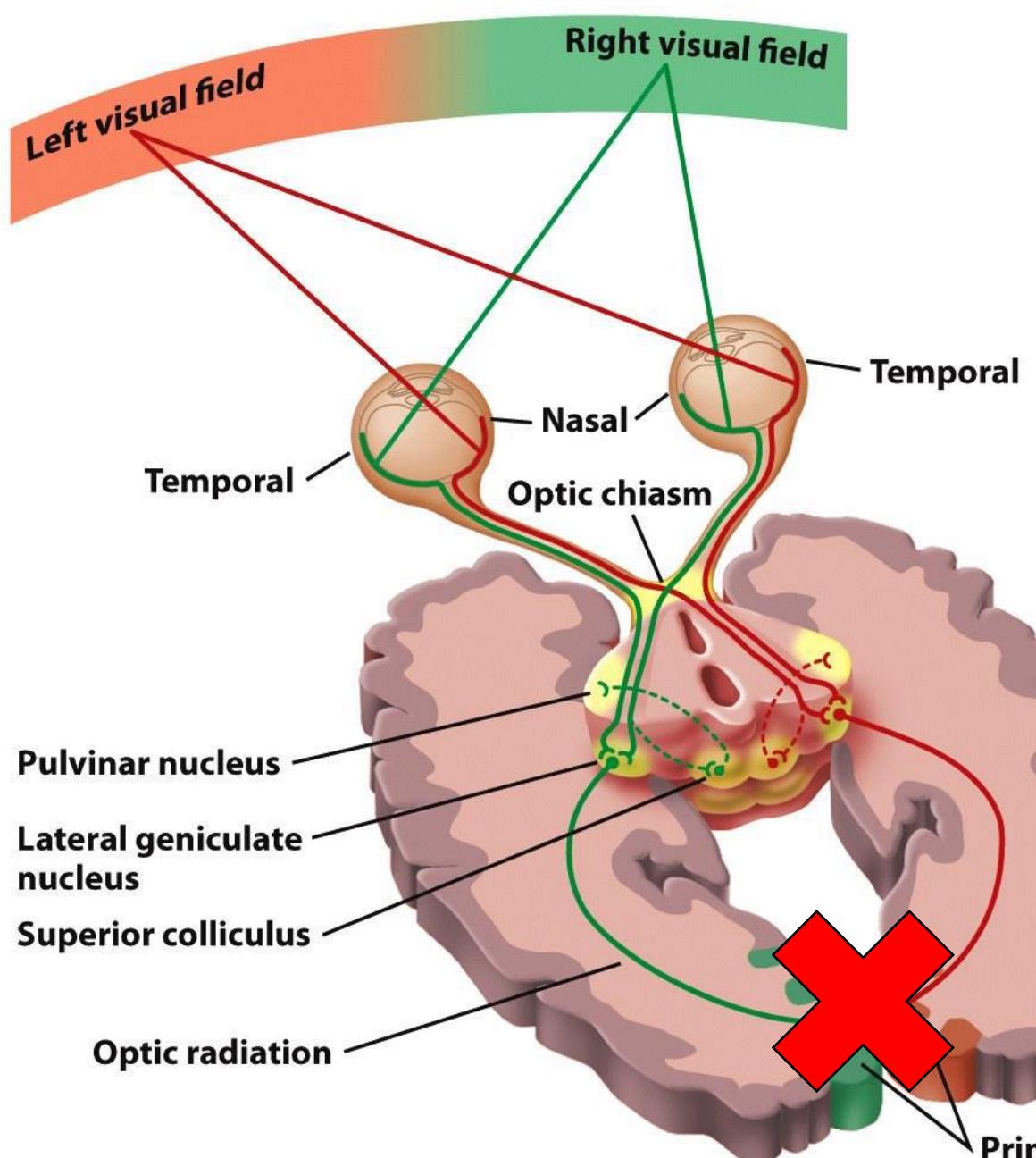


"OFF center"
↓ AP-frequency
in the center of the RF





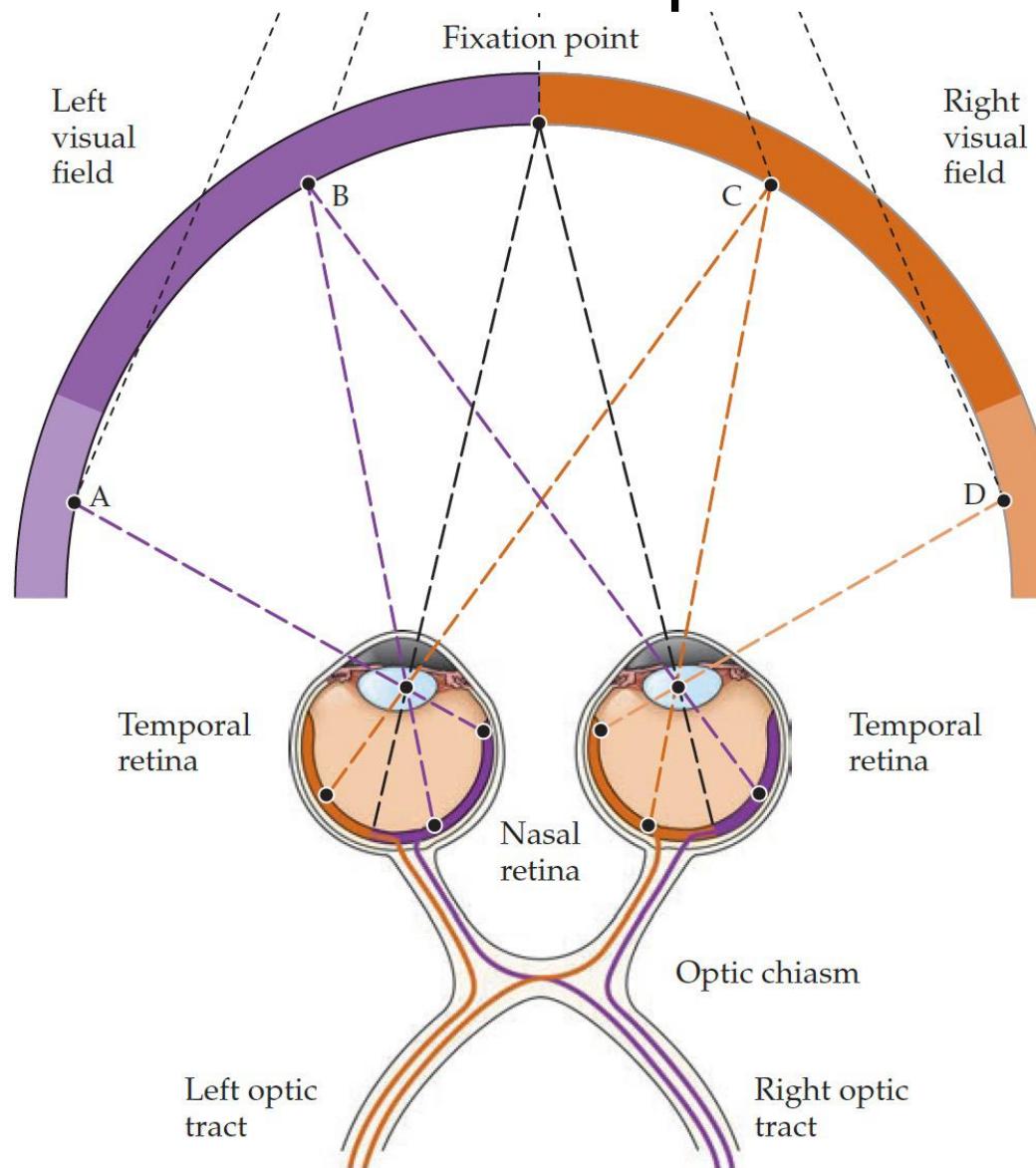
Retinal input
passes through
LGN into primary
visual cortex



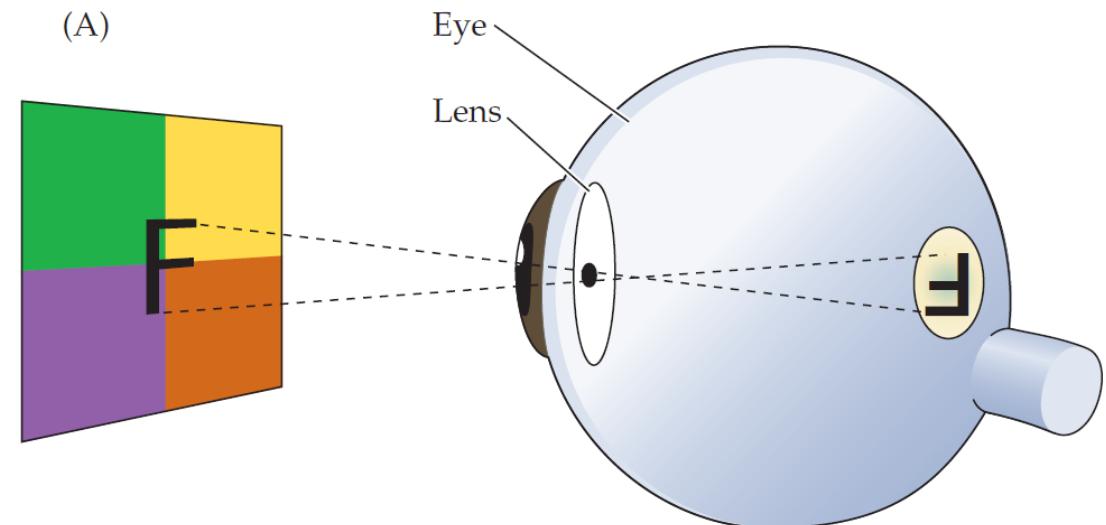
Blindsight?

Some visual function without input into primary visual cortex

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

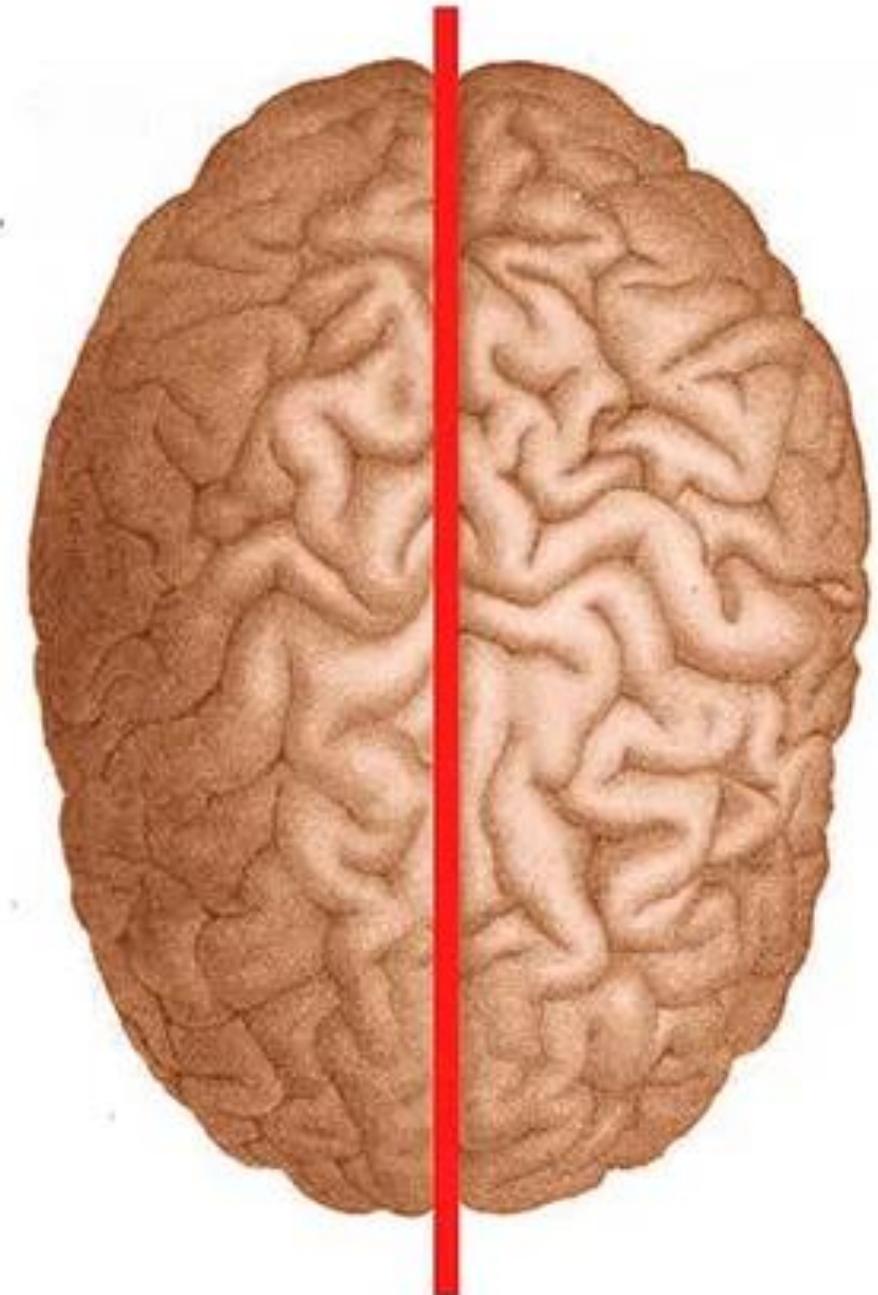
Two hemispheres are connected via corpus callosum.

Sever this connection as a last resort to treat epilepsy.

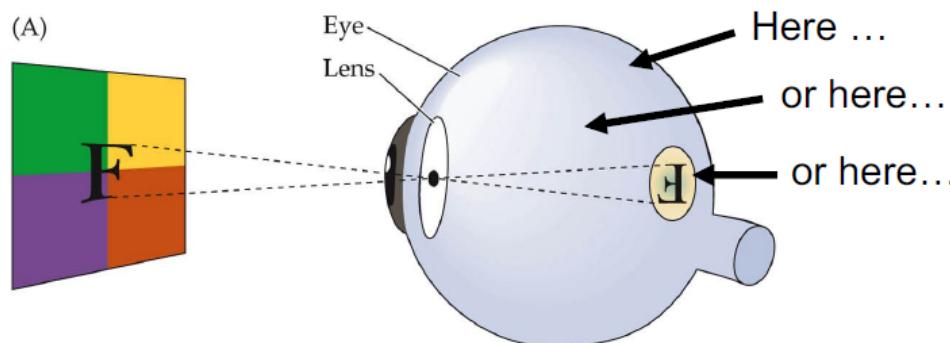
Effect in some patients: **alien-hand syndrome**, e.g.

- 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)

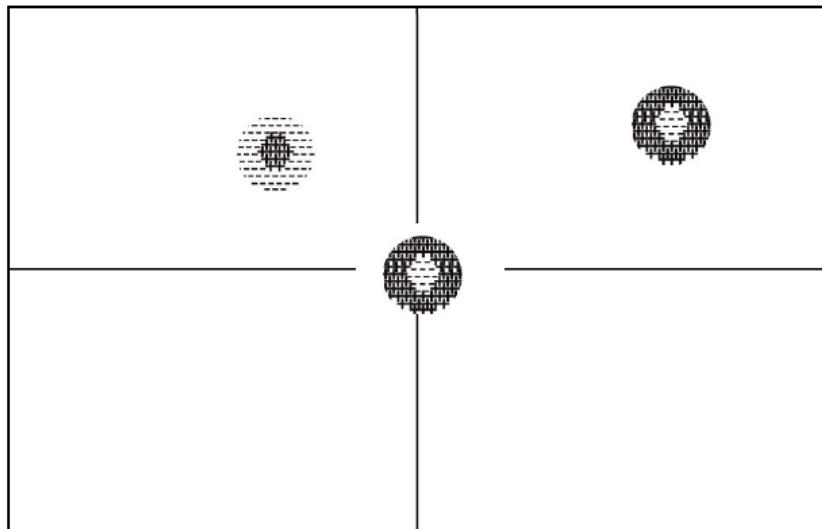
Seen as evidence for dual consciousness after corpus callosotomy.



Receptive fields



Field of view:



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



On-center
ganglion cell

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



Off-center
ganglion cell

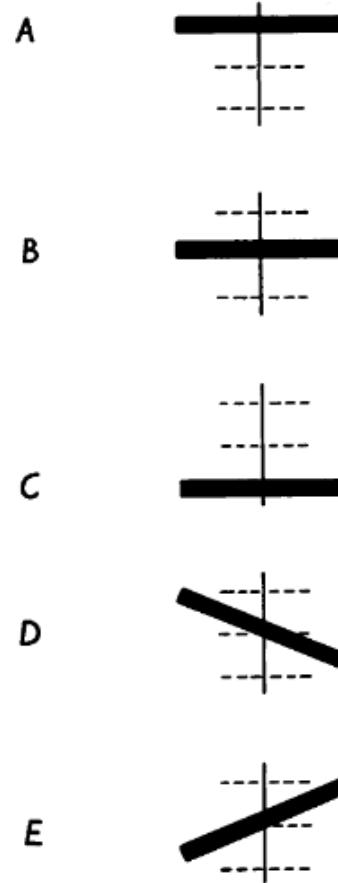
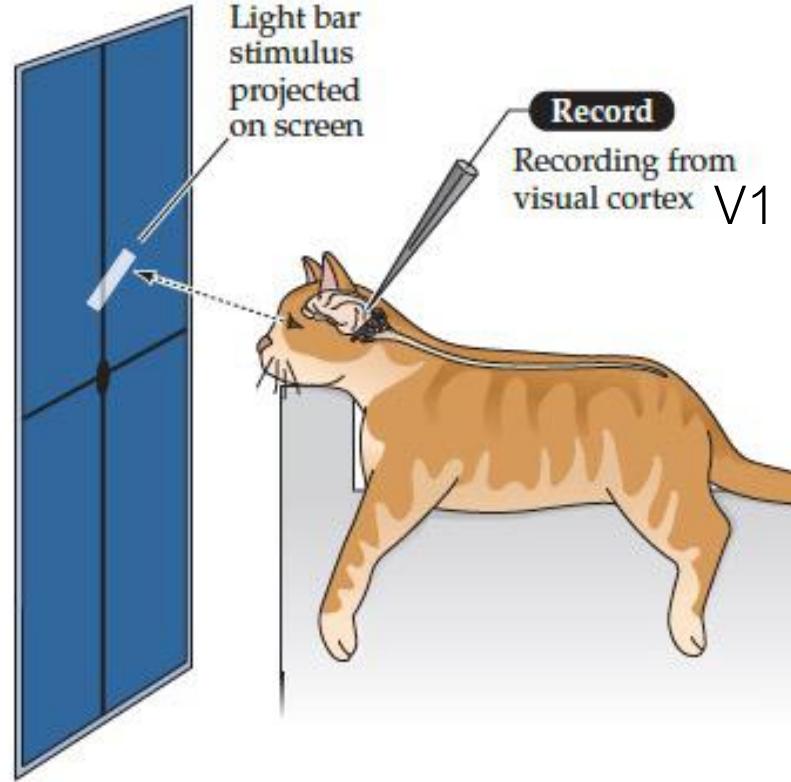
"OFF center"
↓ AP-frequency
in the center of the RF

V1

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

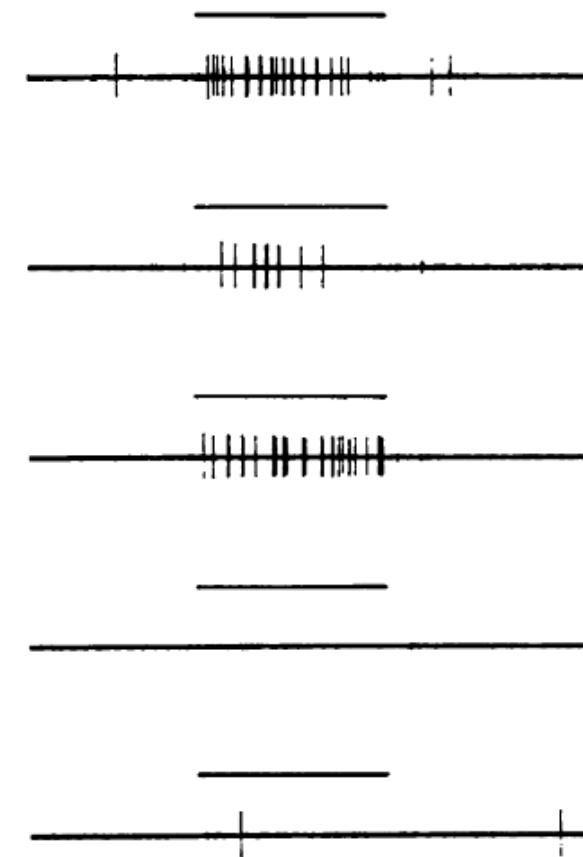
Textbook function: edge detection.

(A) Experimental setup



Hubel and Wiesel 1962

Nobel Prize 1981

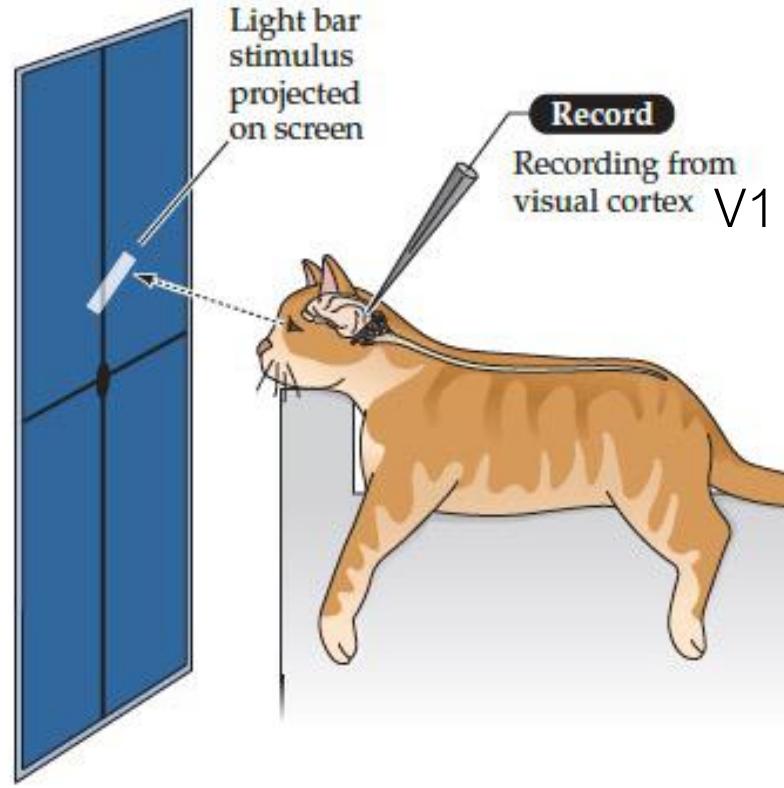


V1

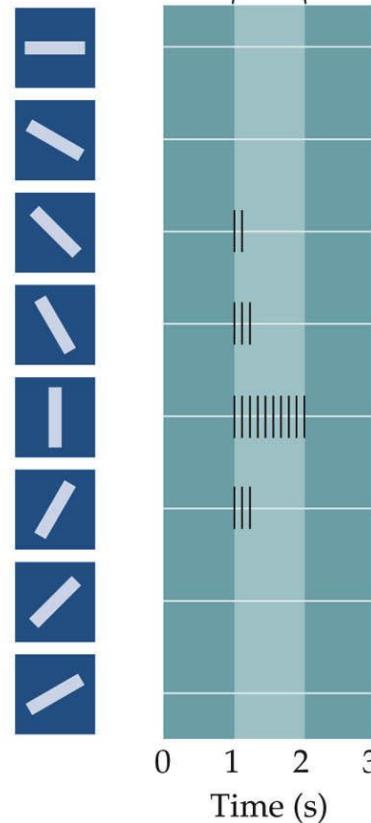
Receptive fields: $\sim 1^\circ$ visual angle.

Textbook function: edge detection.

(A) Experimental setup



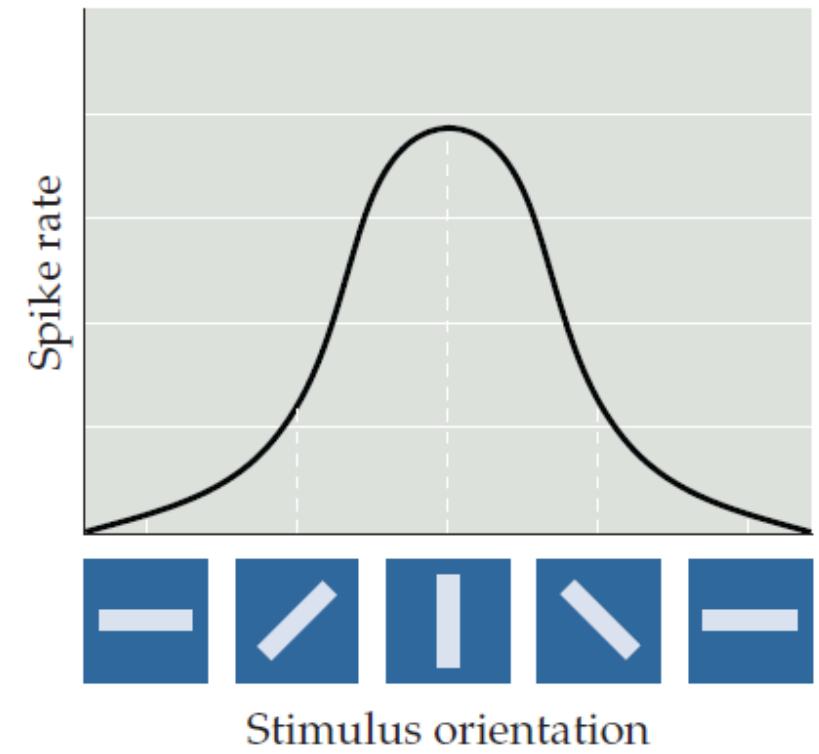
(B) Stimulus orientation Stimulus presented



Hubel and Wiesel 1962

Nobel Prize 1981

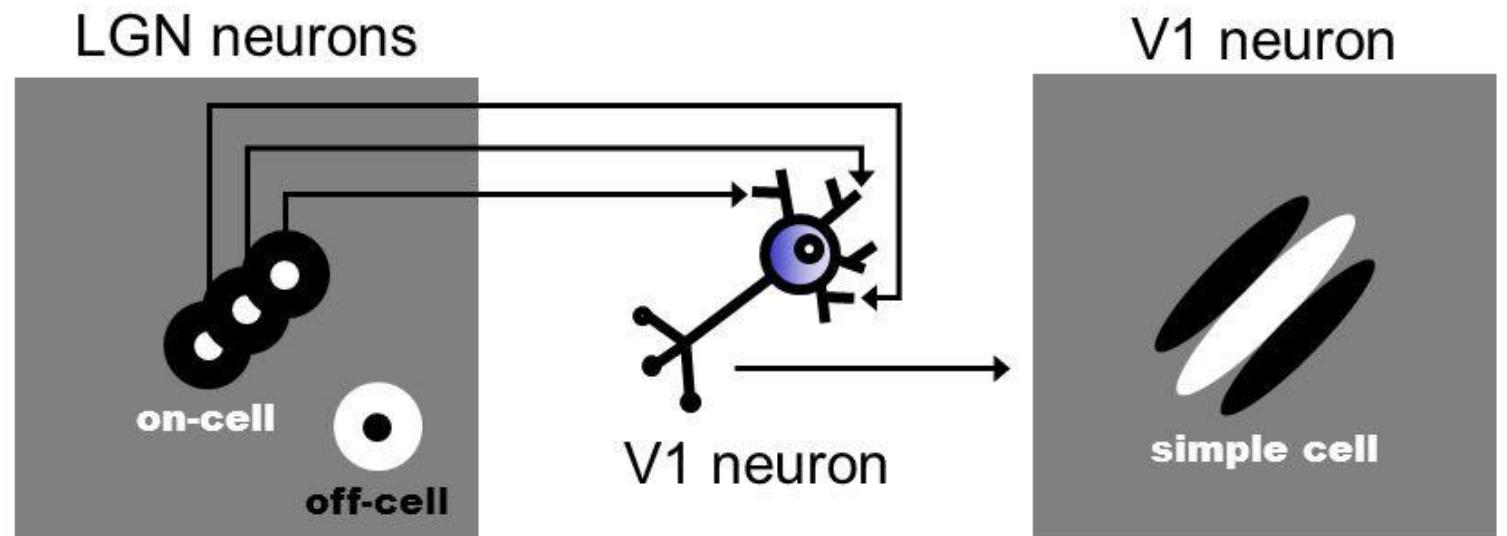
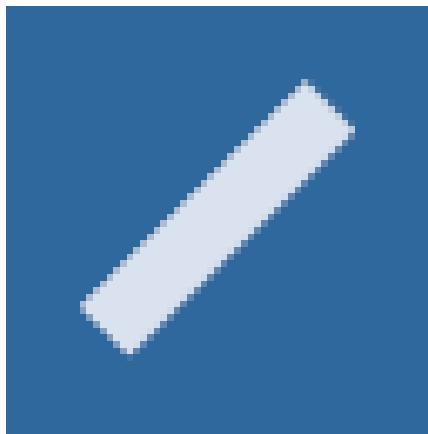
(C)



V1

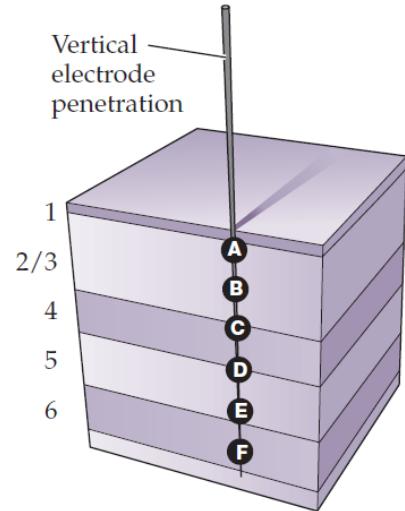
Receptive fields: $\sim 1^\circ$ visual angle.

Textbook function: edge detection.

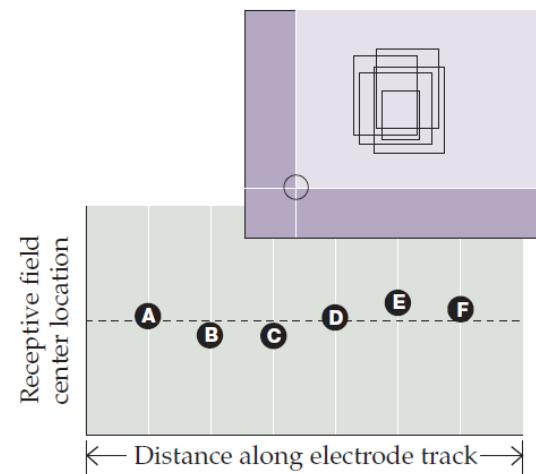


V1 columnar organization

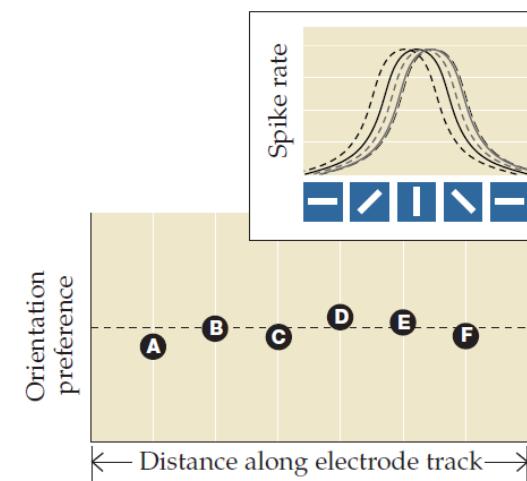
(A)



Receptive field position

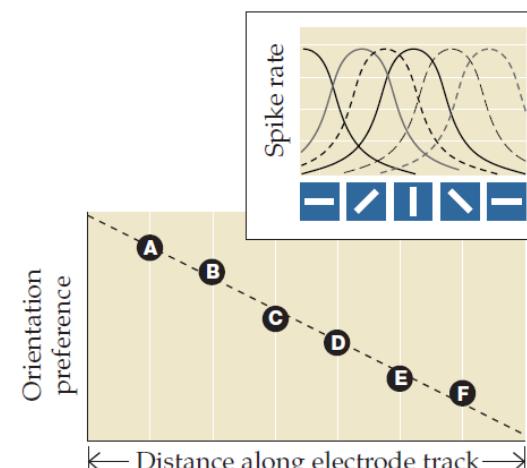
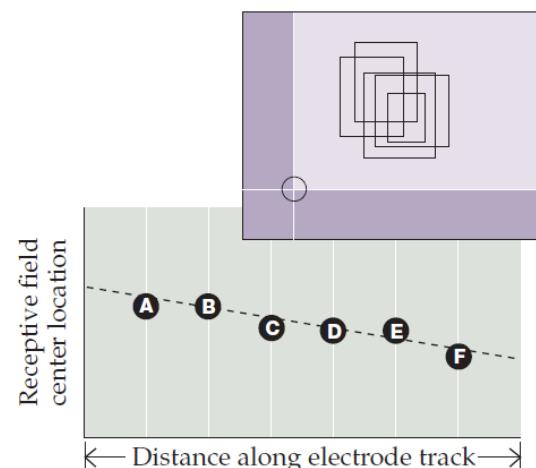
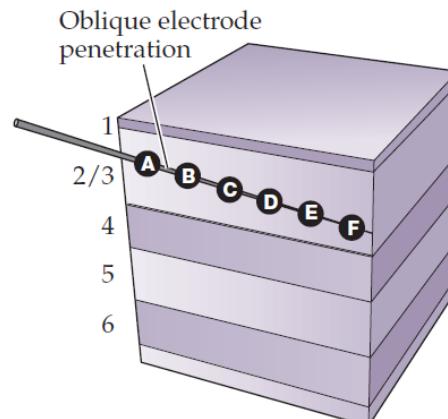


Orientation tuning curves



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

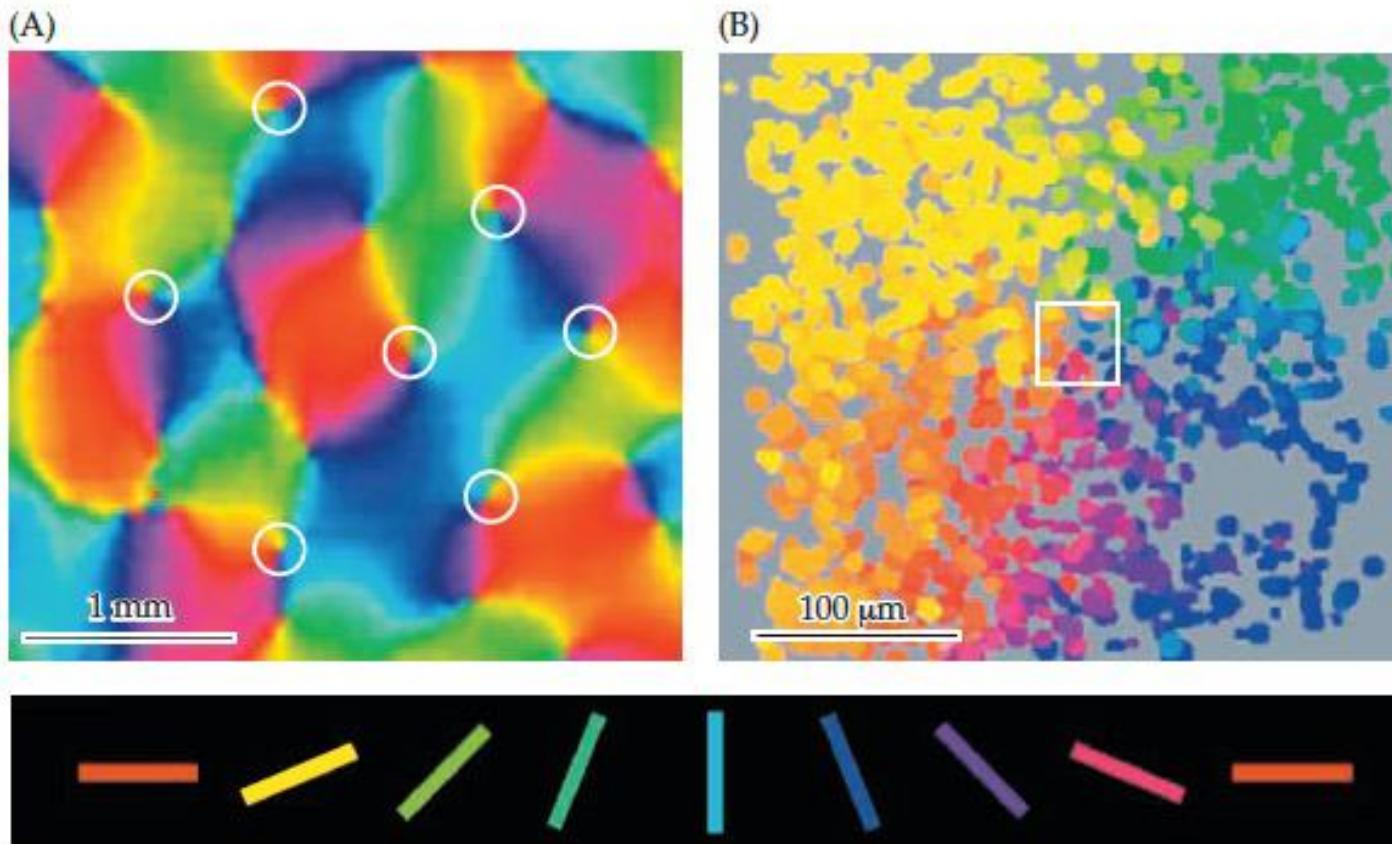
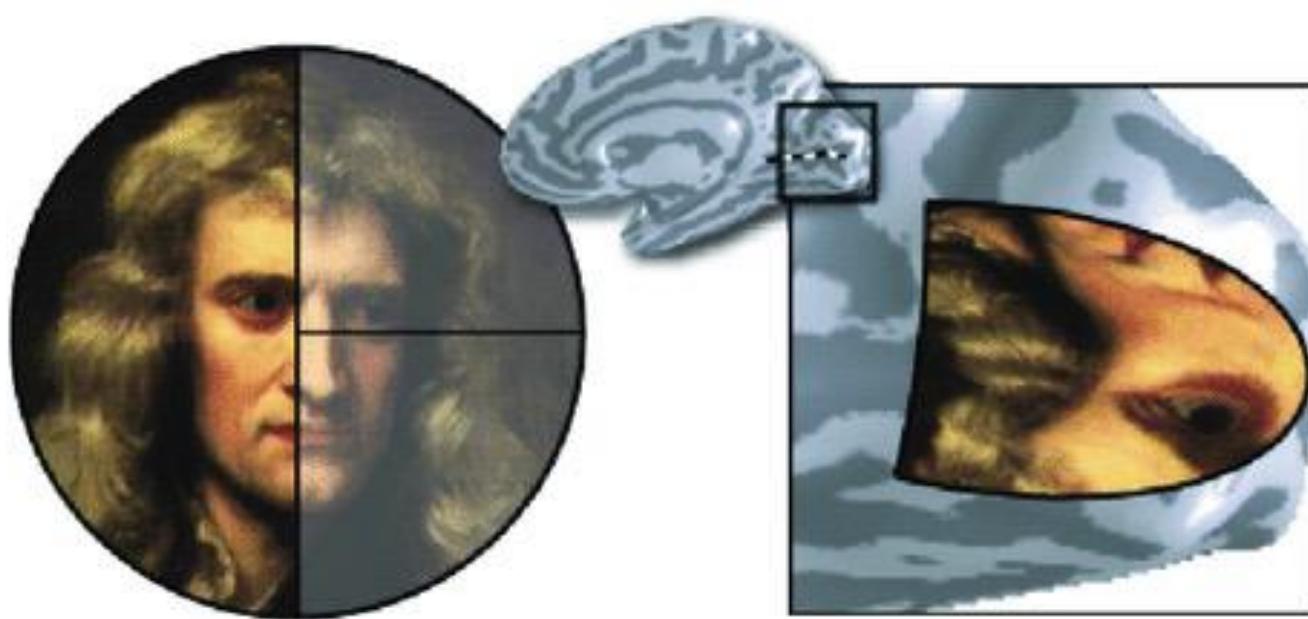


FIGURE 12.10 Functional imaging reveals orderly mapping of orientation preference in the primary visual cortex.

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

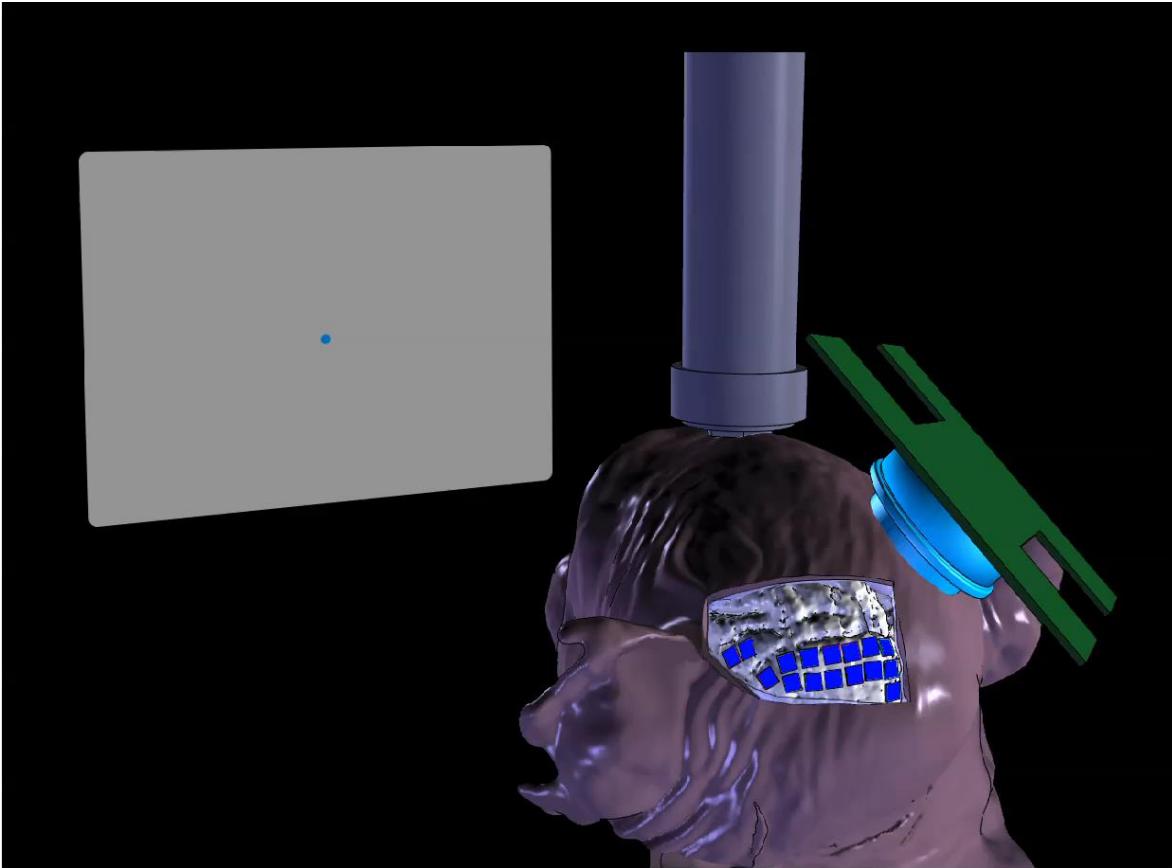
V1 retinotopy



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

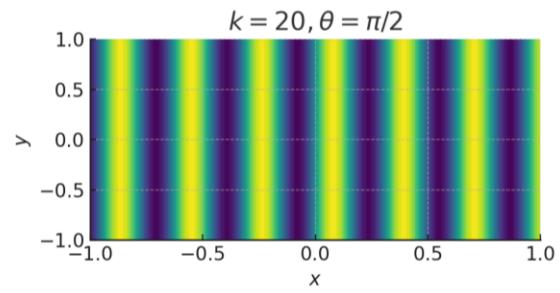
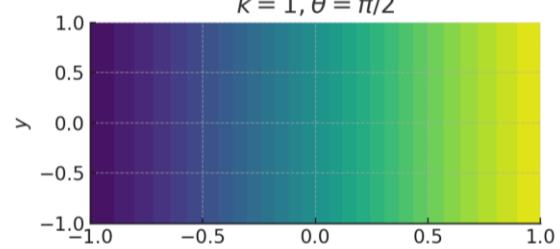
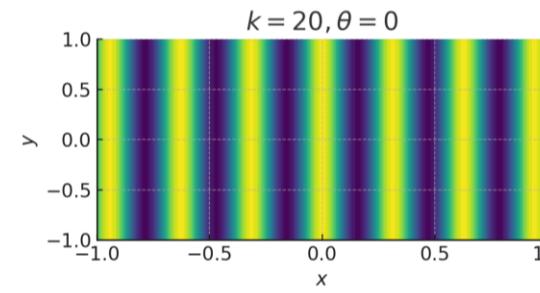
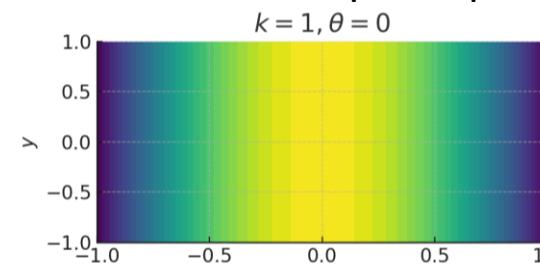
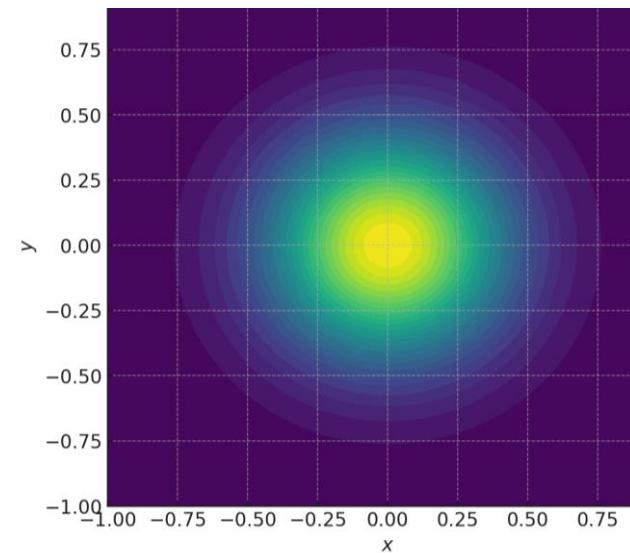
Computational model of V1: Gabor filters

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

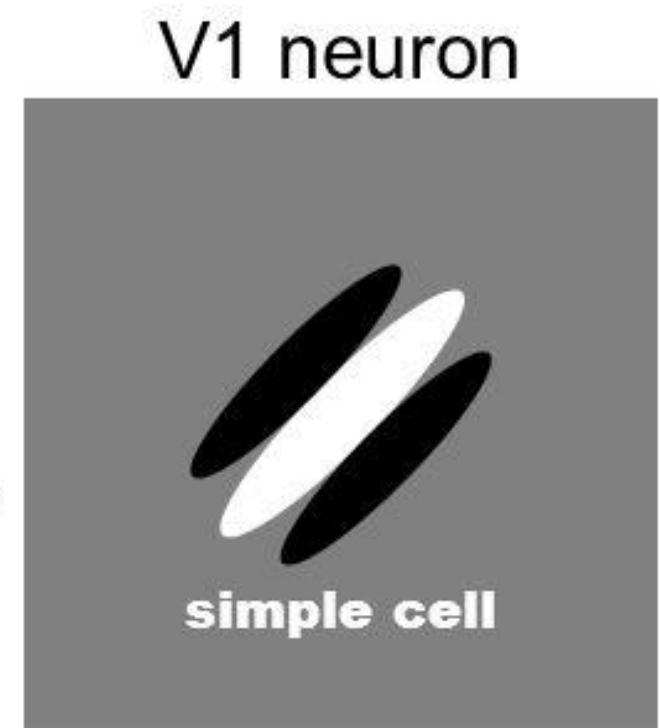
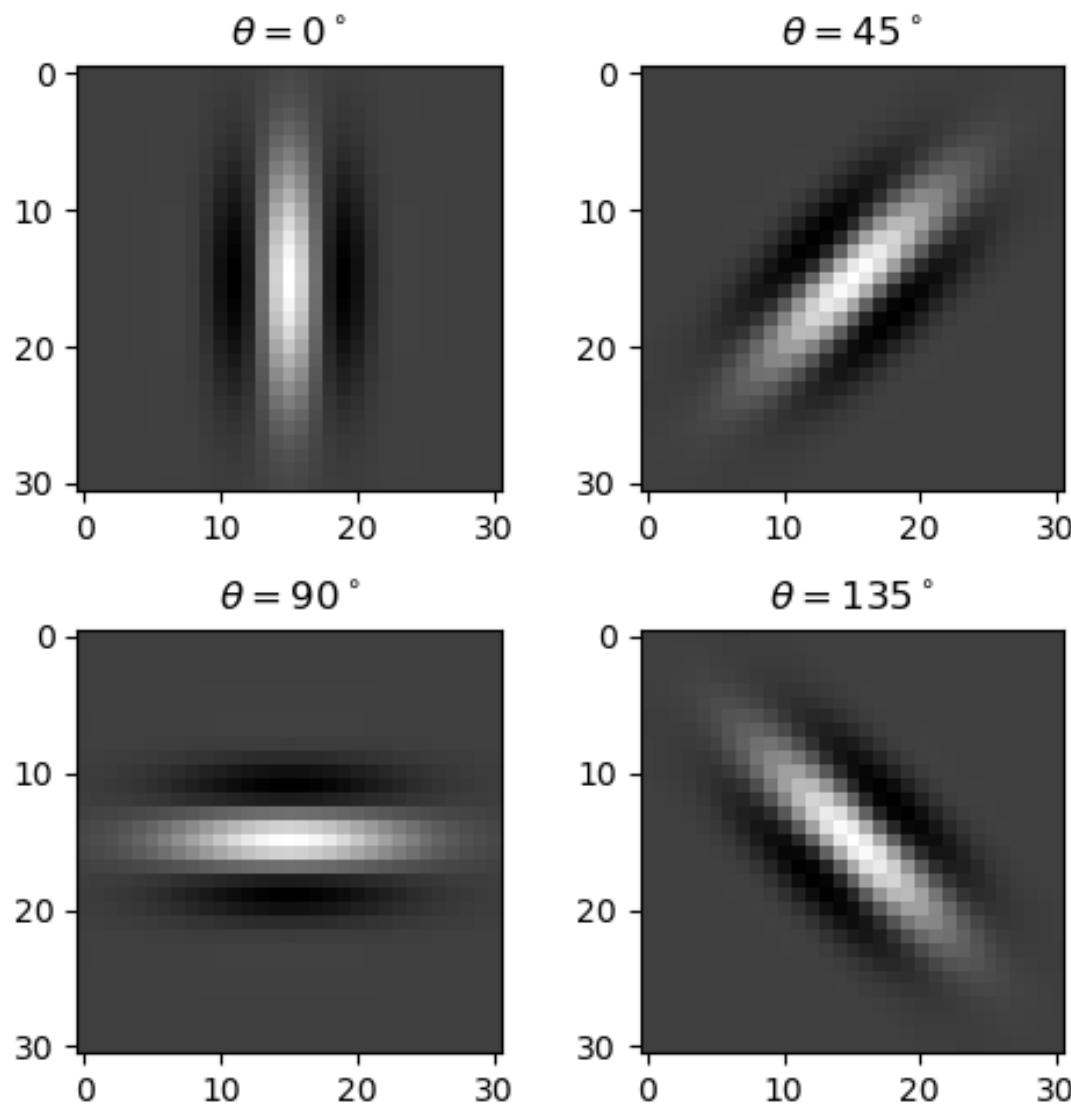
no rotation with this formulation, need x+y dependency



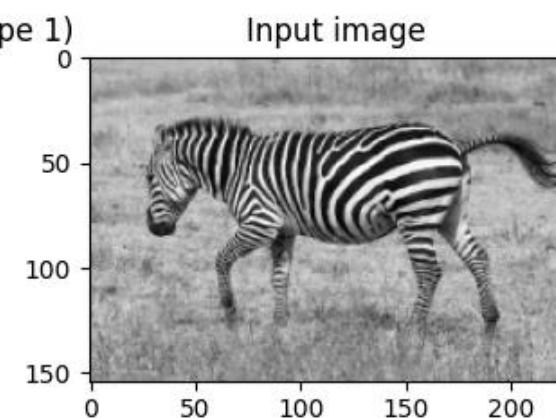
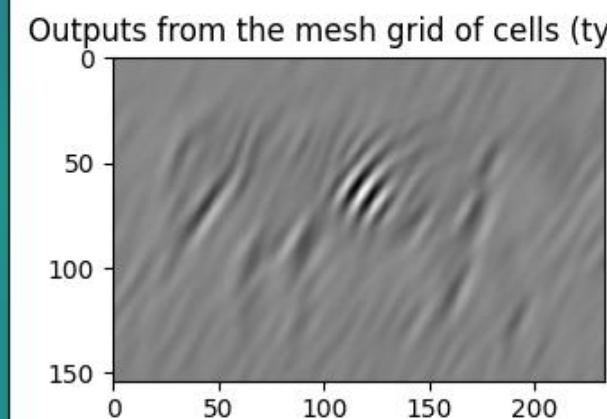
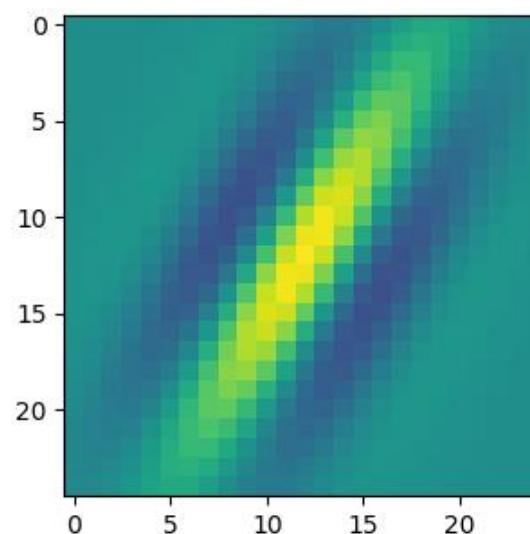
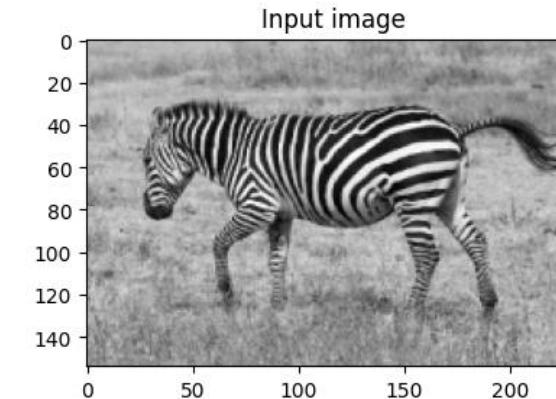
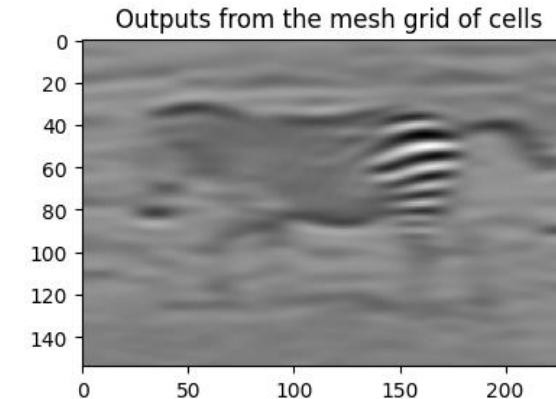
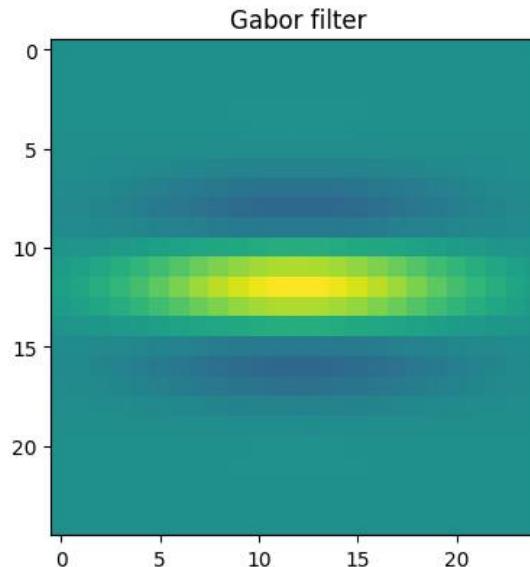
with:

- σ_x, σ_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)
- φ is the preferred spatial phase

Gabor model: example filters



Gabor model: example filters applied



Visual hierarchy

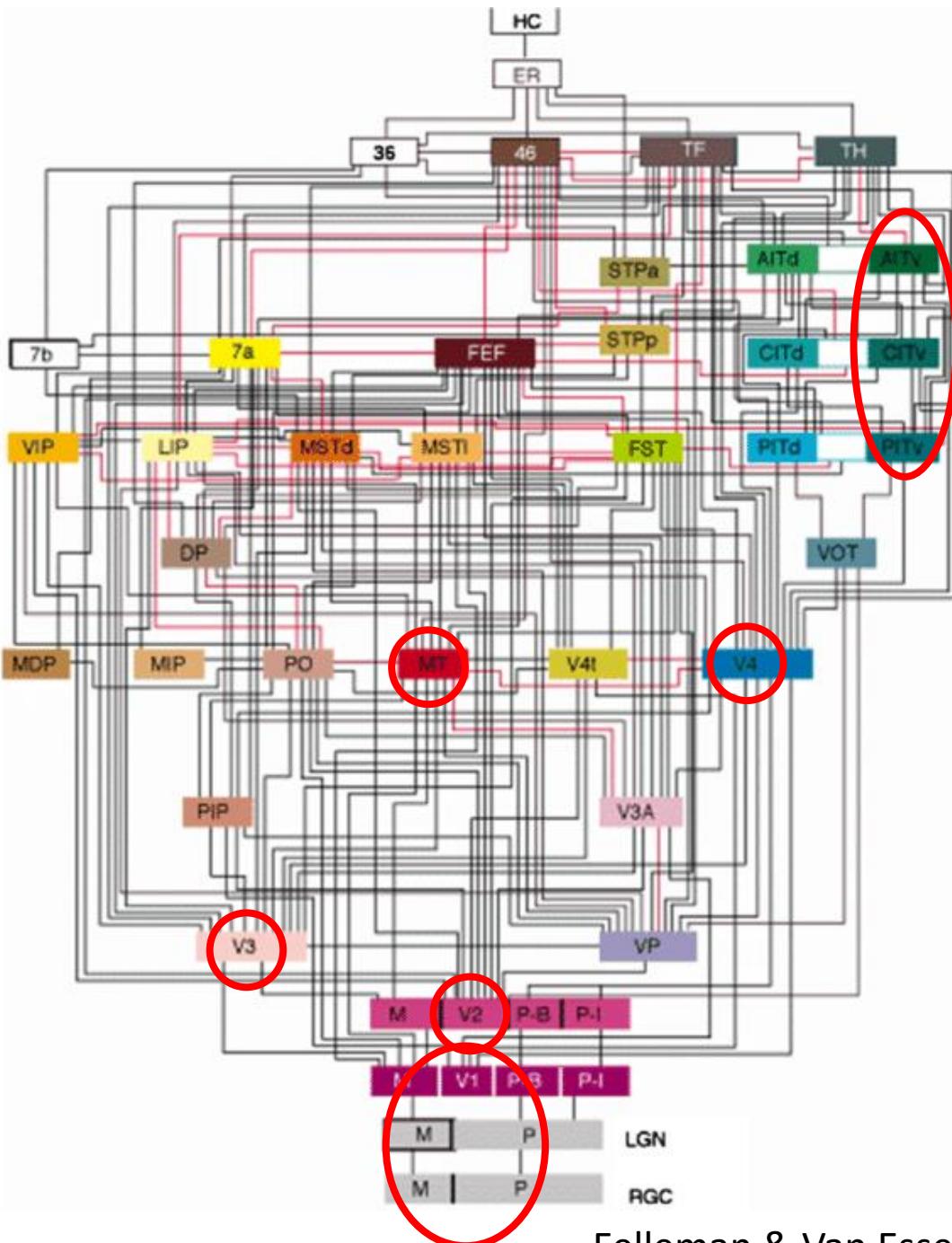
Subcortical structures

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

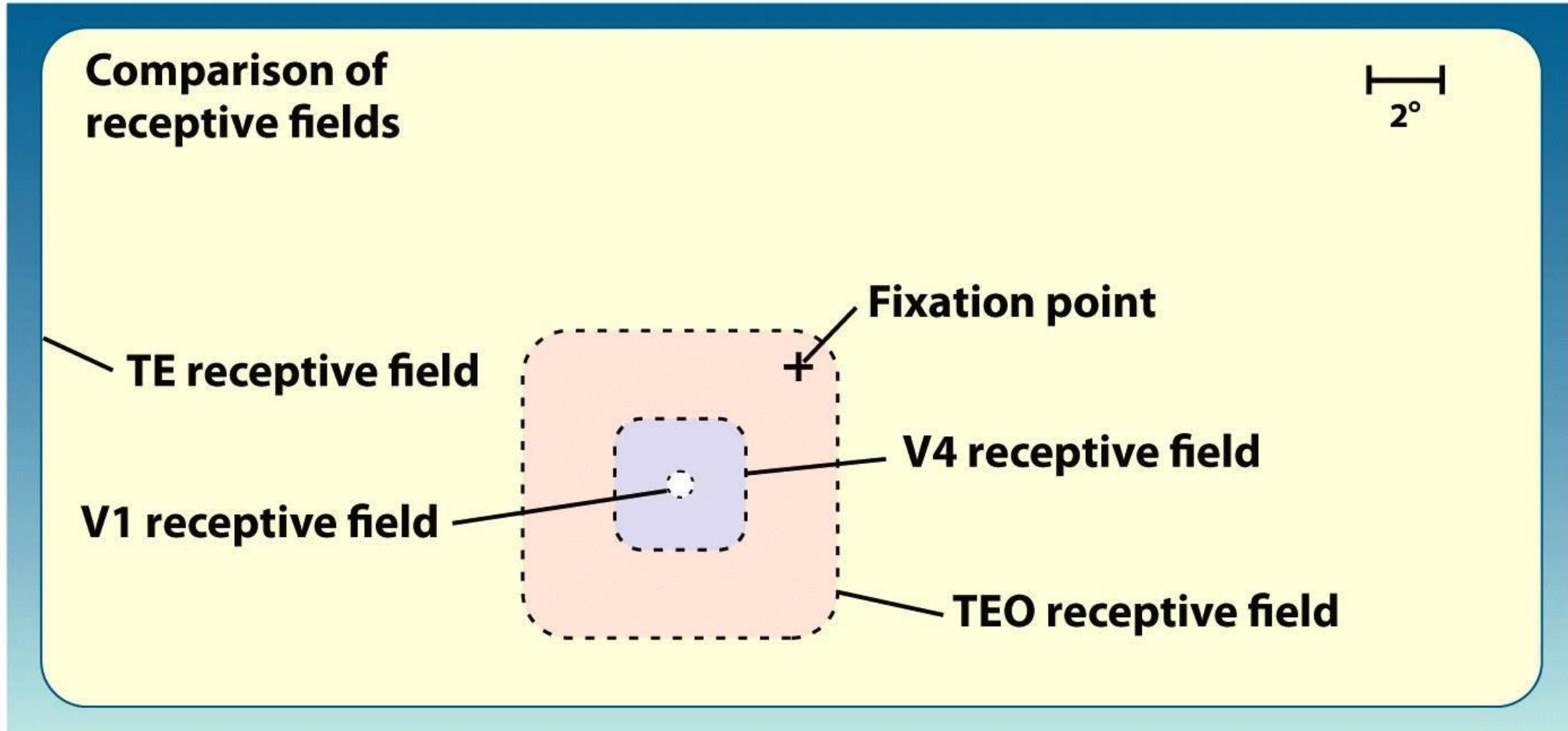
Cortex

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

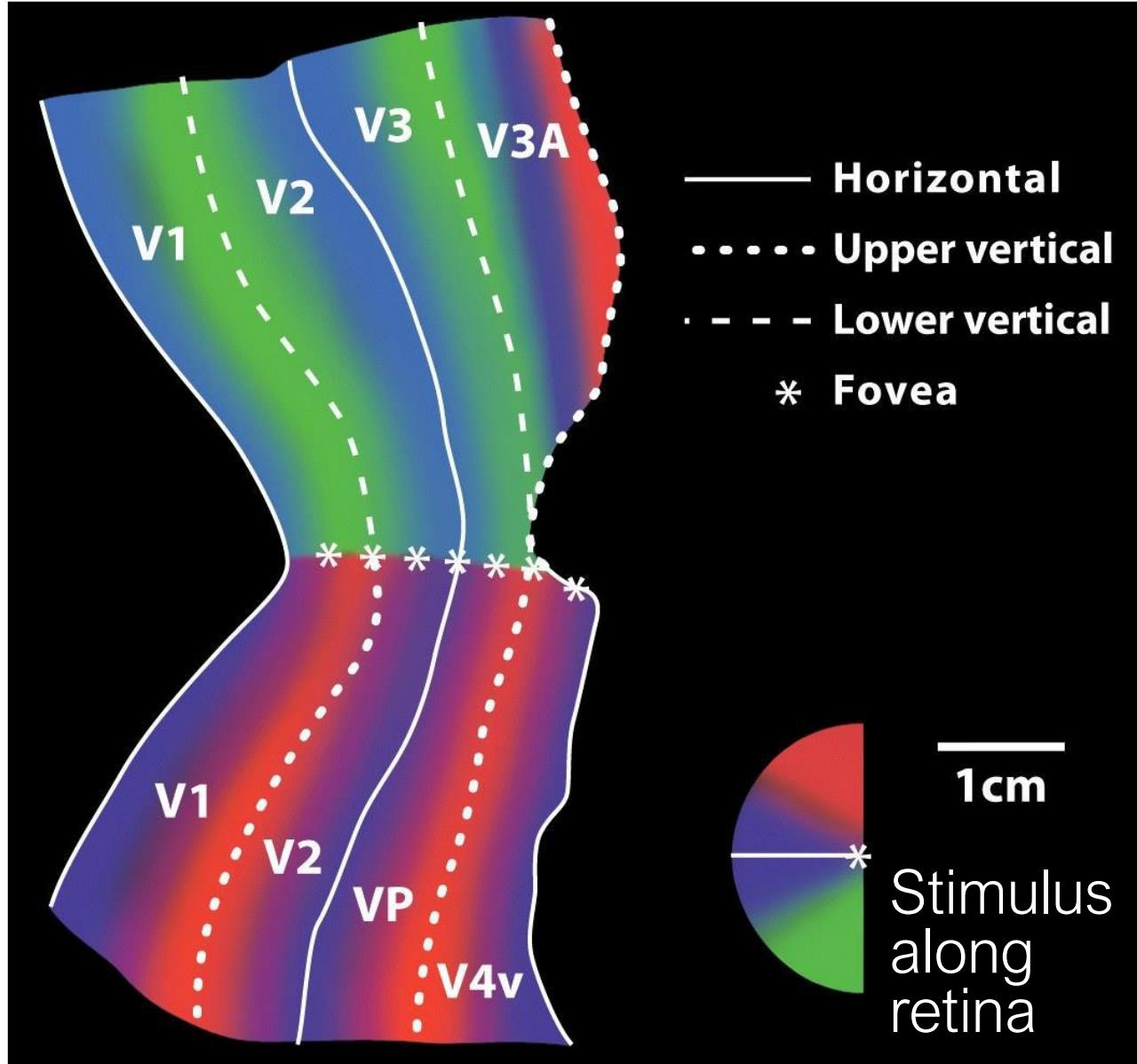
...and to a whole lot of other areas!



Beyond V1: receptive field sizes increase along the visual hierarchy



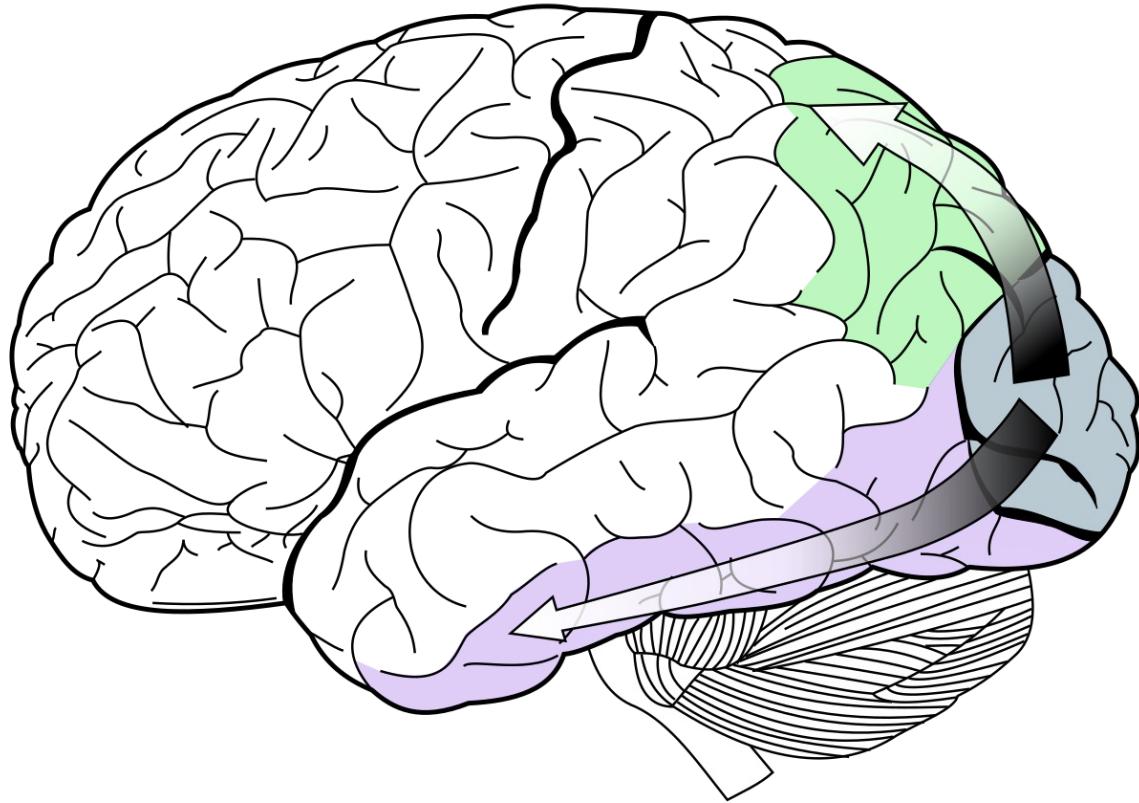
Retinotopy



Mapping of visual input from retina to neurons

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

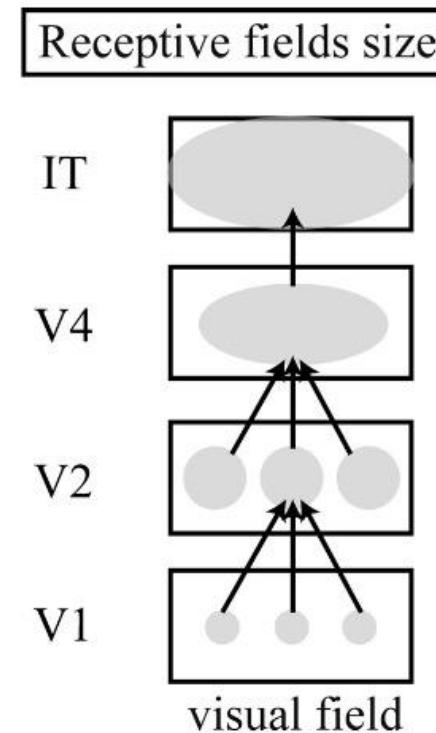
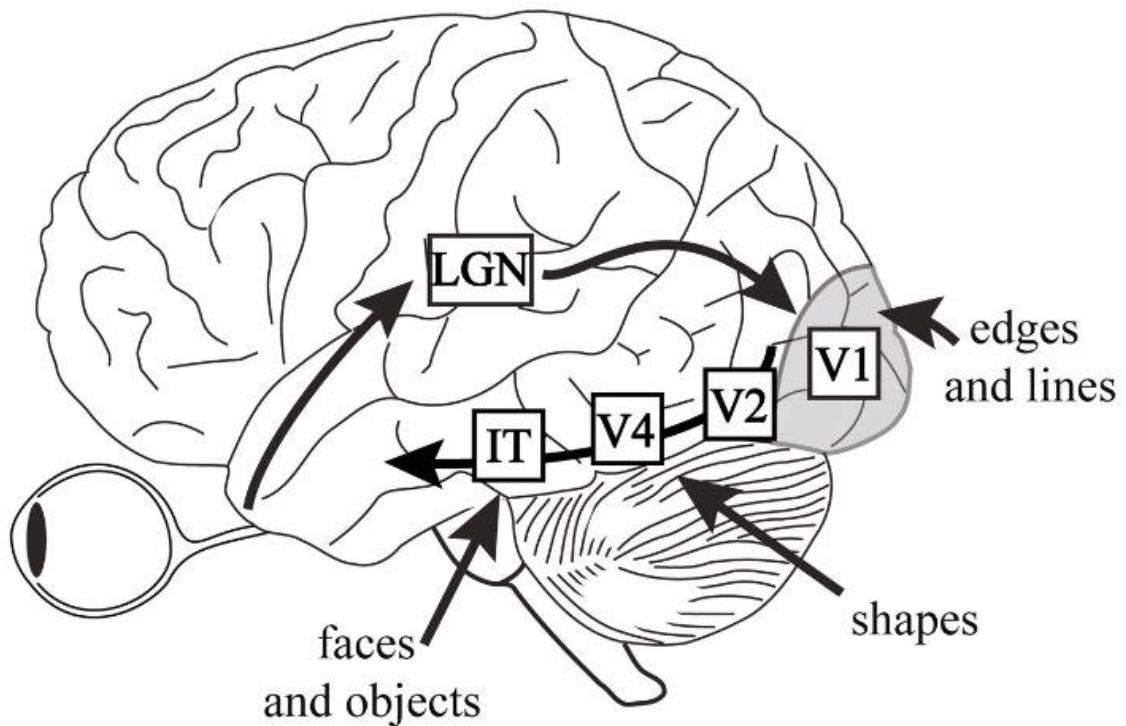
Two-streams hypothesis



Dorsal pathway:
spatial vision
“where?”

Ventral pathway:
object recognition
“what?”

Visual ventral stream

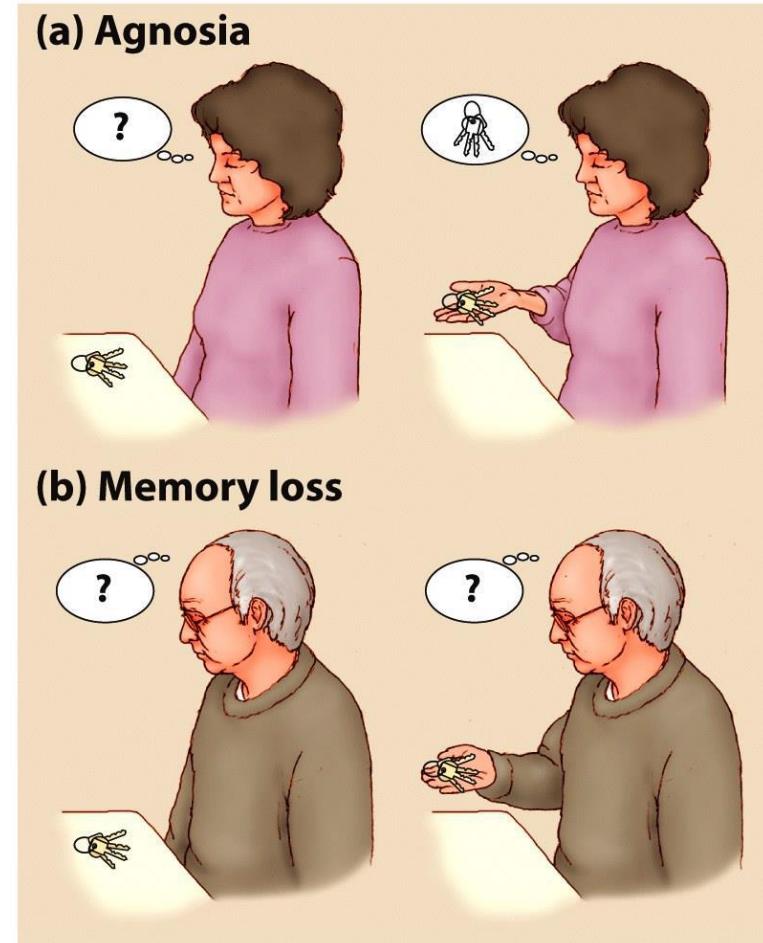


Visual agnosia: Inability to recognize objects despite normal vision

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.



Courtesy of Bob Desimone's class MIT 9.017

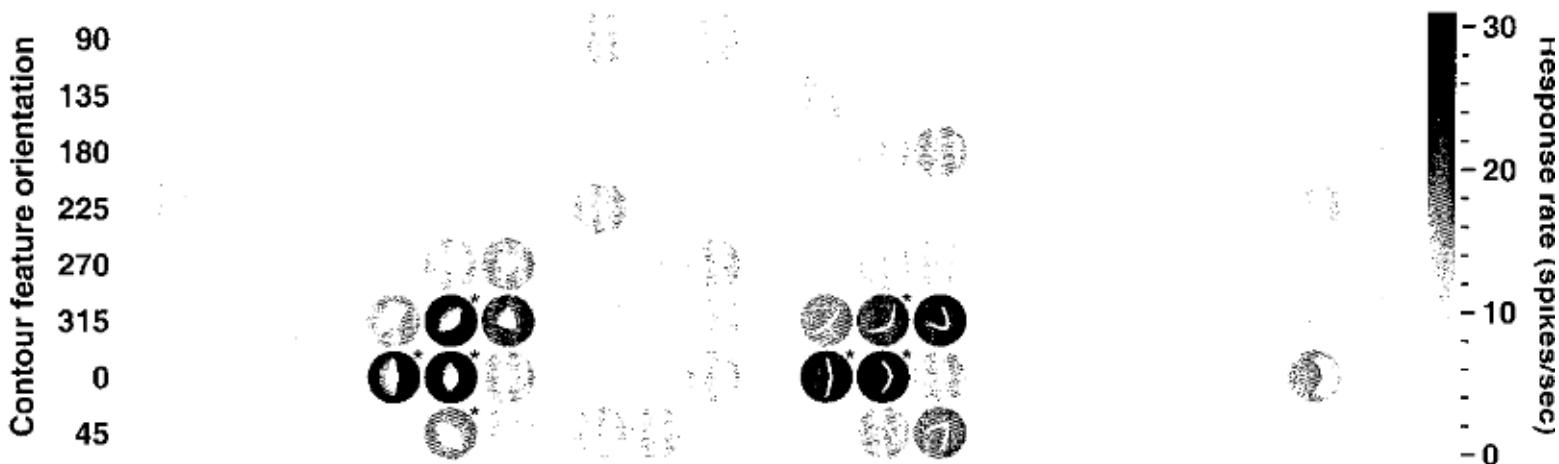
Ventral stream: V2, V4

V2

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

V4

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

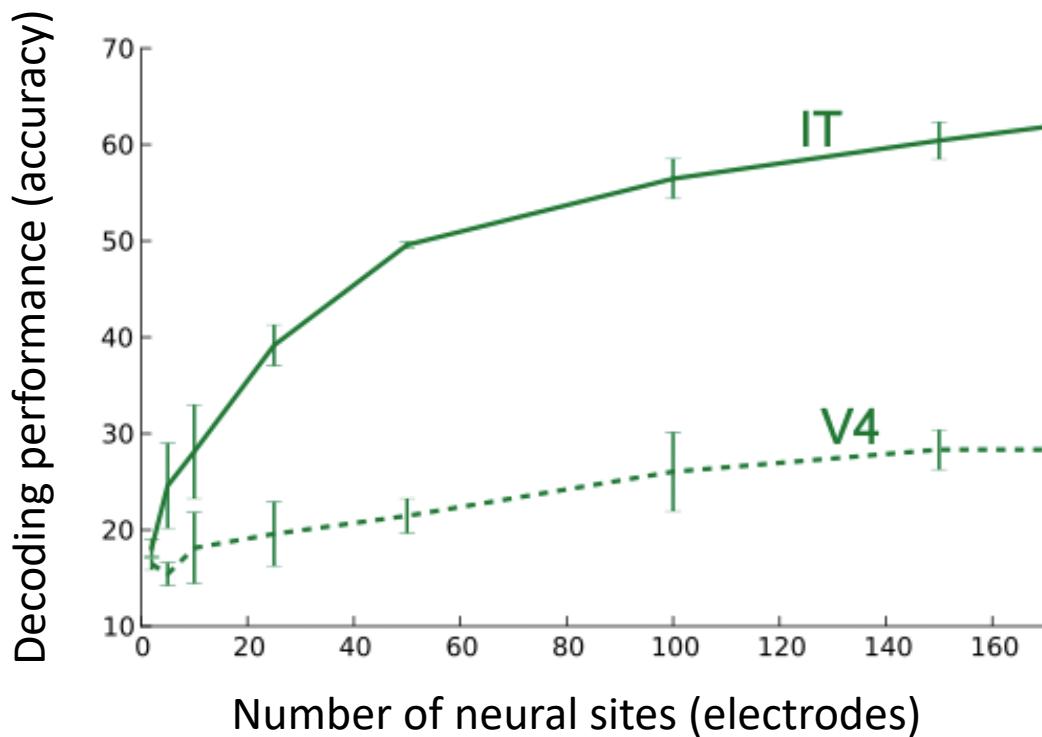
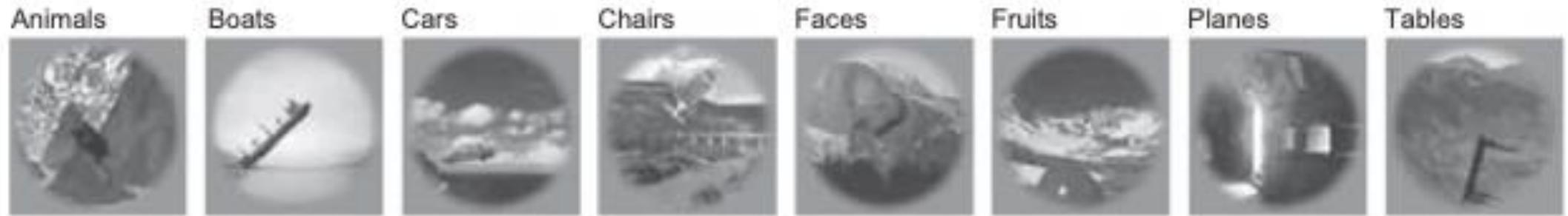


Ventral stream: IT

- RF size: ~12°
- Function: objects
- Directly supports object recognition
(correlational and causal evidence)

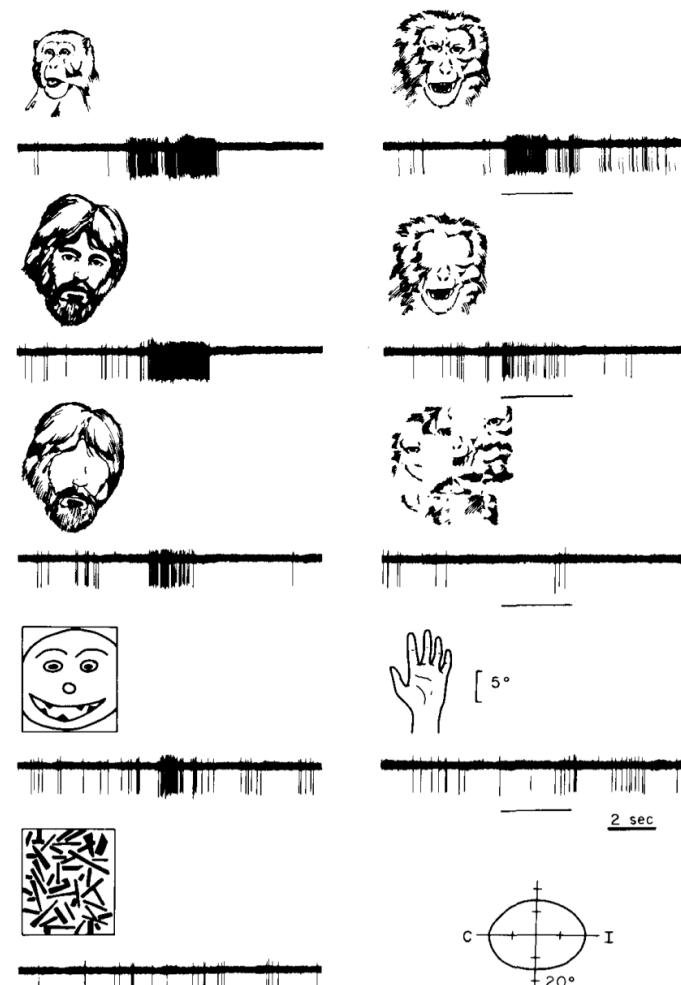
IT linearly provides object category

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



Ventral stream: IT Face cells

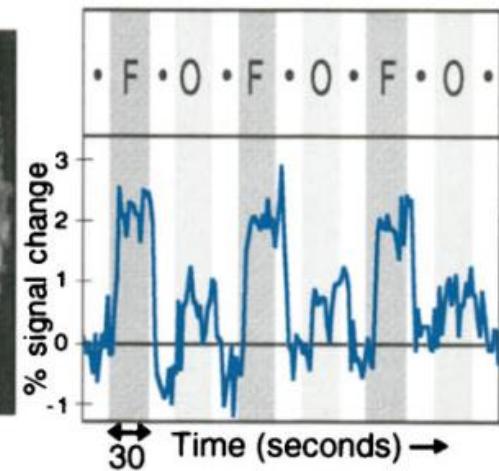
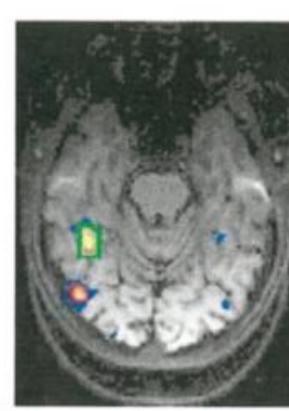
Single neuron preference for faces



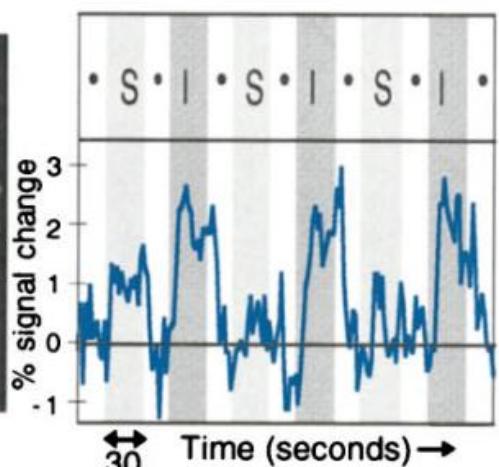
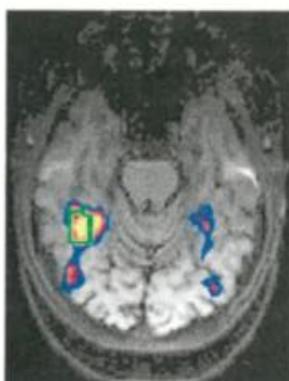
Bruce, Desimone, Gross 1981

Face patches

3a. Faces > Objects



3b. Intact Faces > Scrambled Faces



Kanwisher et al. 1997

Stimulating face-selective sites distorts face perception



"You just turned into somebody else. Your face metamorphosed."

Stimulating face-selective sites *induces* face perception

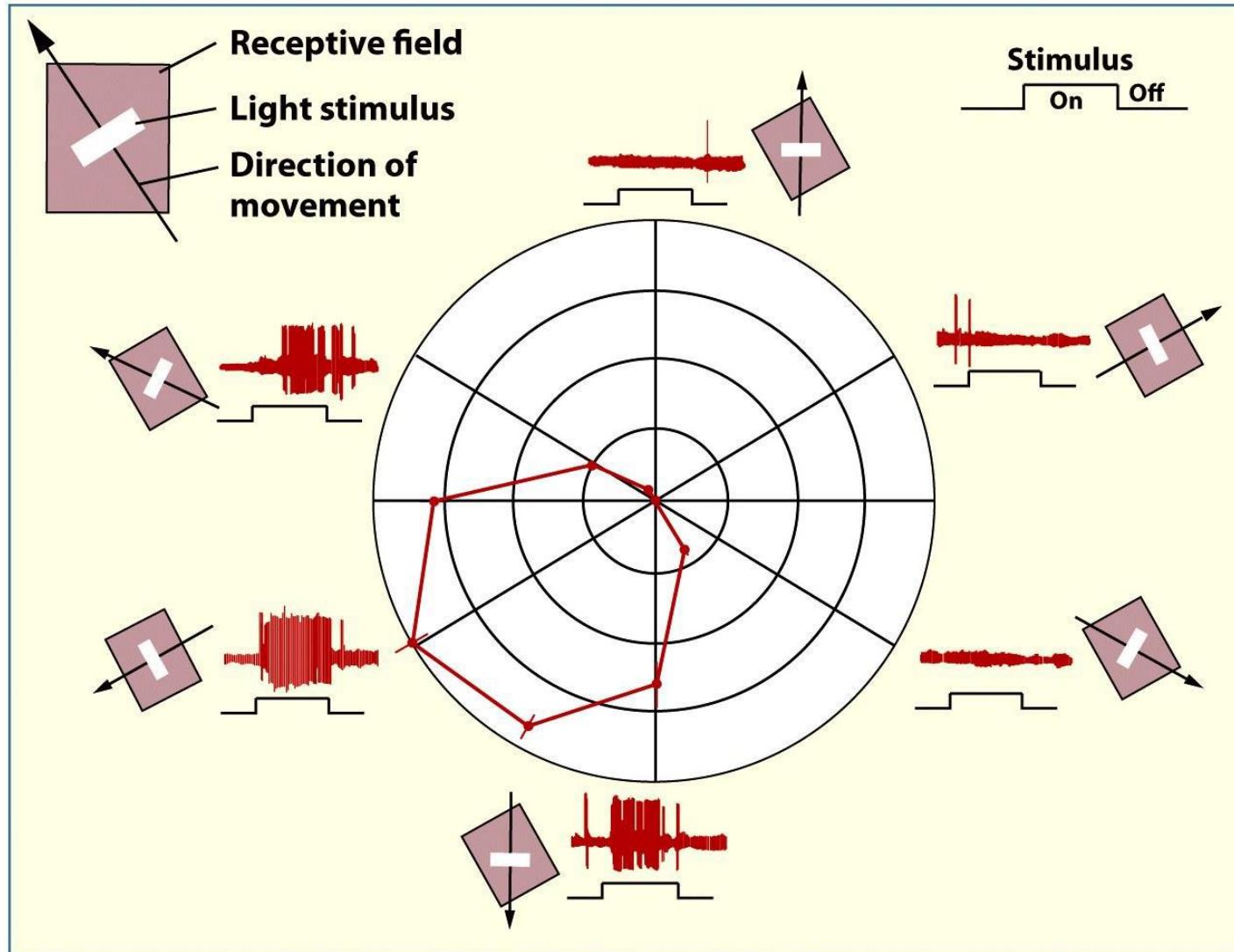
				
Stim. Elec. 181-182	<i>...just for the very first second... I saw an eye, an eye, and a mouth.</i>	<i>How do I explain this? Just like the previous one, I see an eye, an eye, and a mouth, sideways.</i>	<i>Your face completely changed...I don't know what's going on. Your eyes change.</i>	<i>Hm. Am I just 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

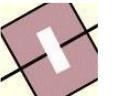
- Highly selective for motion direction and speed
- Invariant to object form or texture
- Thought to underlie motion recognition

Dorsal stream: MT motion preference



single-neuron
direction tuning

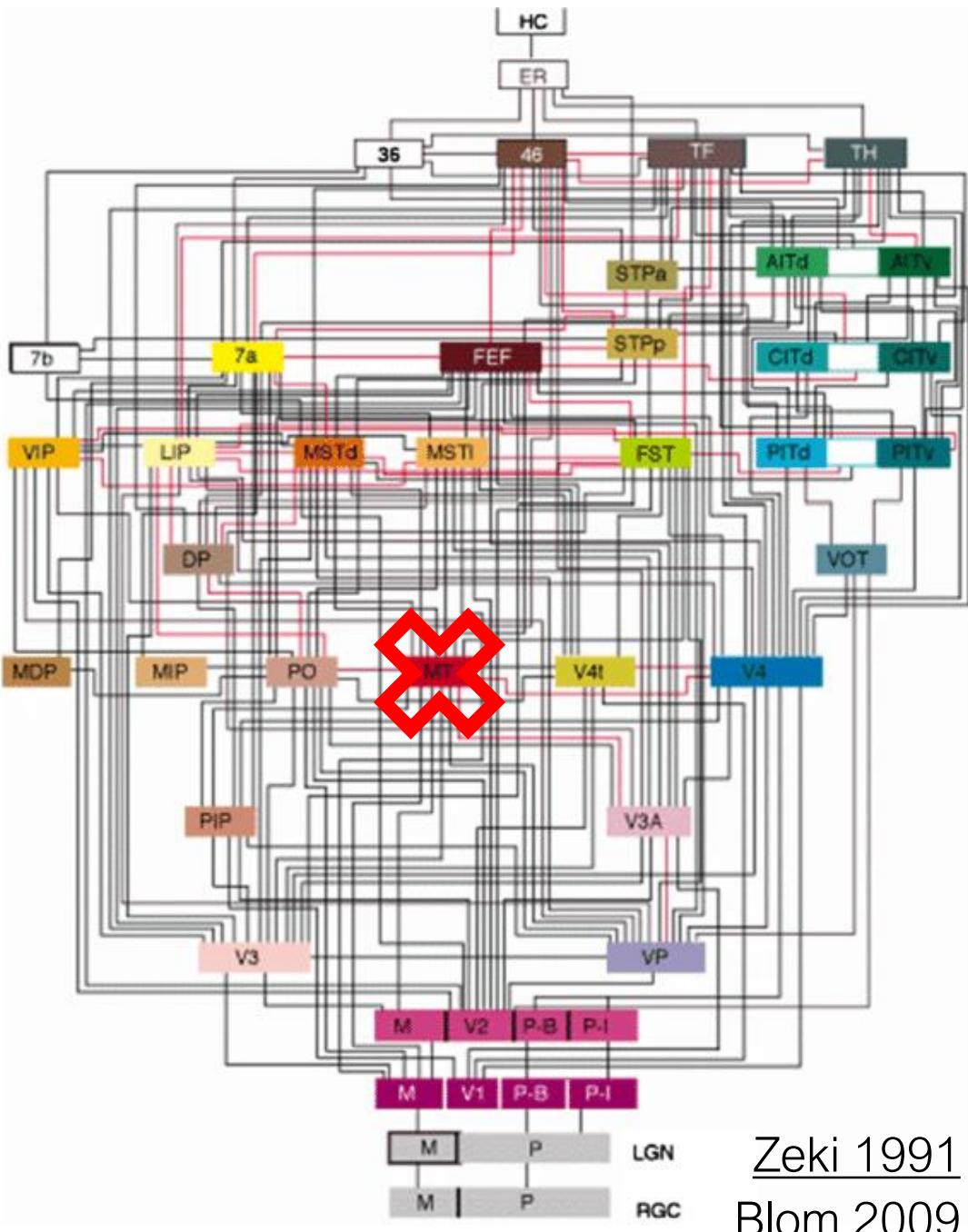
Same bar orientation
evokes different
responses depending
on movement



Akinetopsia

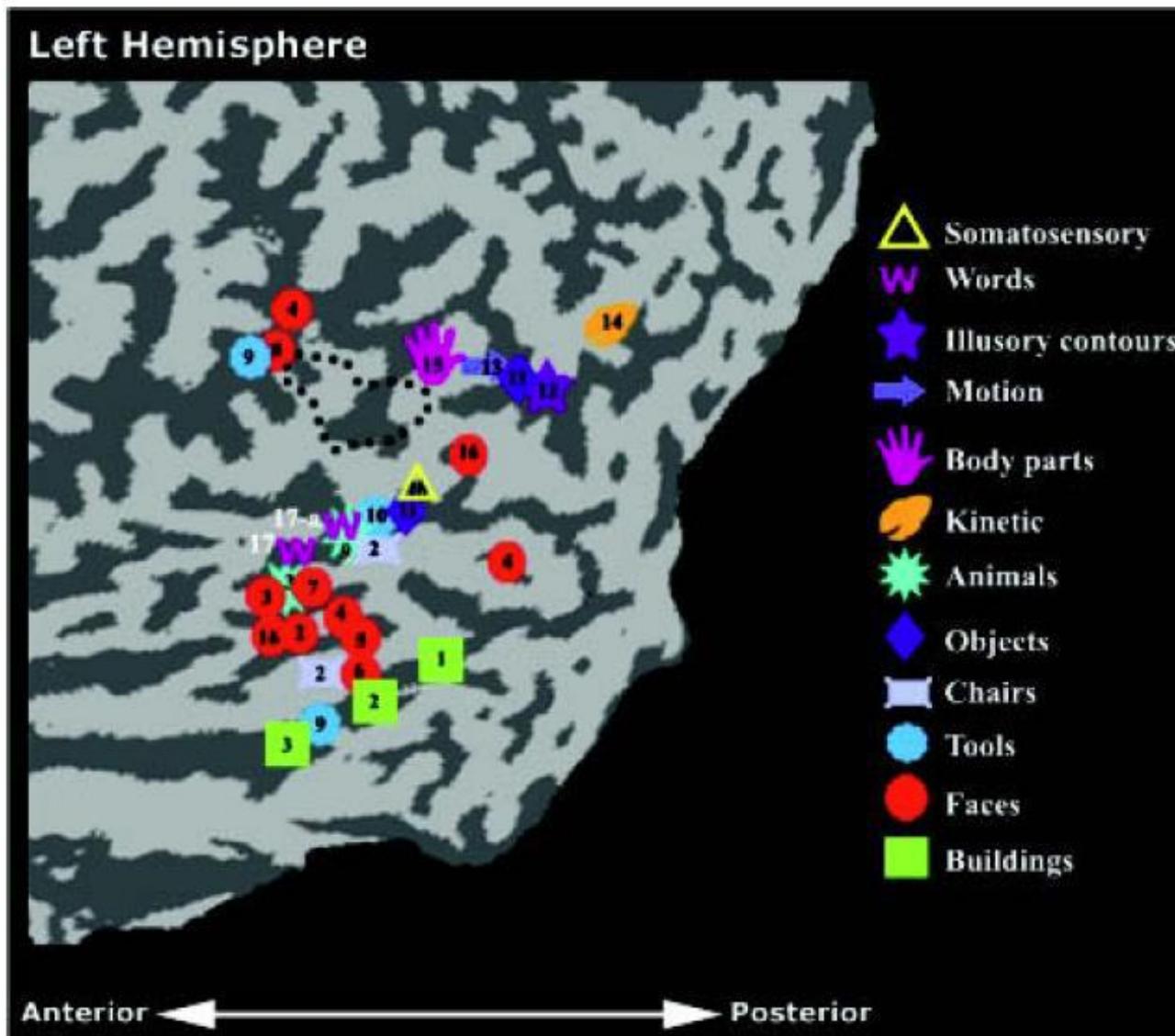
Damage to MT (middle temporal visual area, also called V5)

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

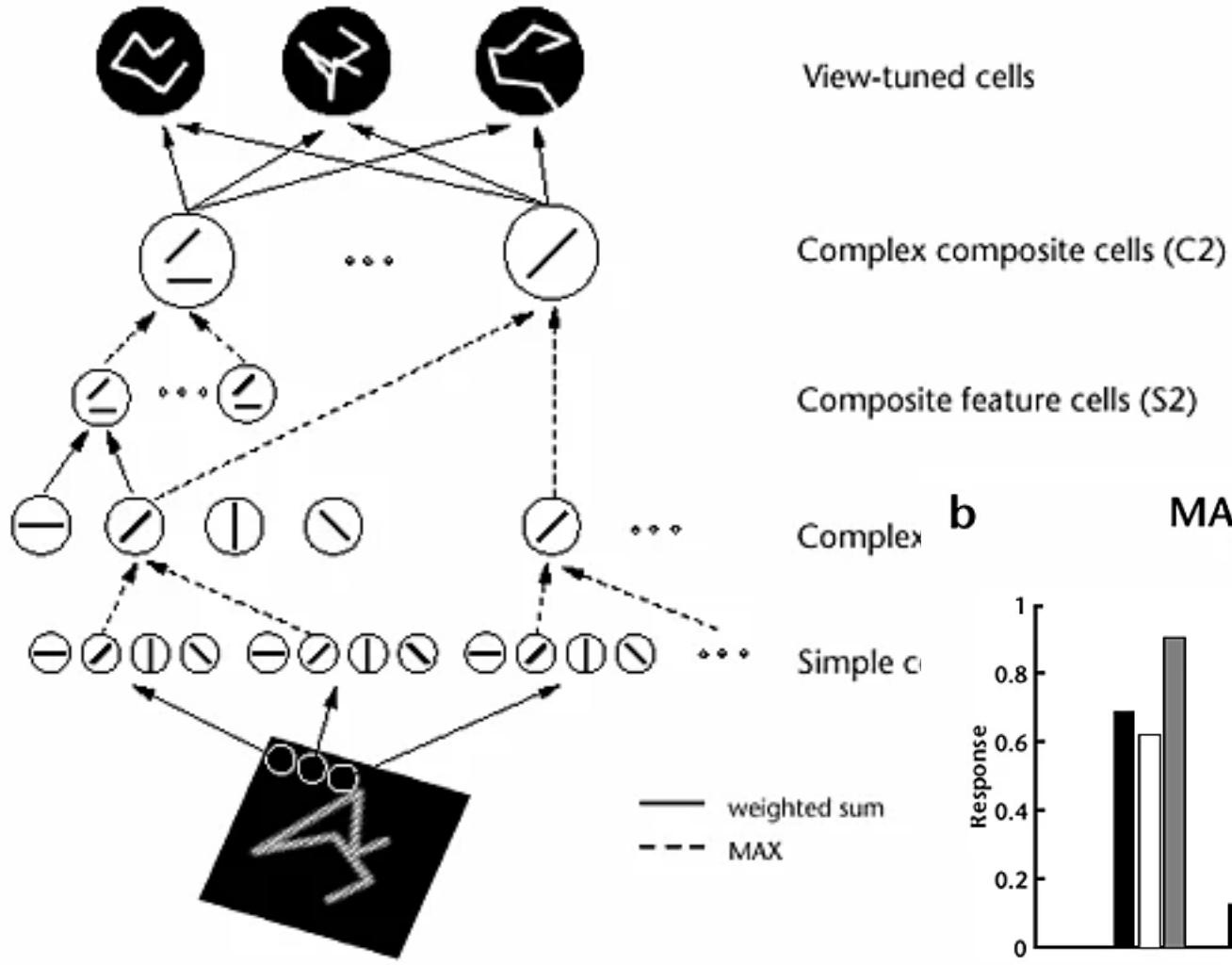


Zeki 1991
Blom 2009

Swiss army knife of visual areas

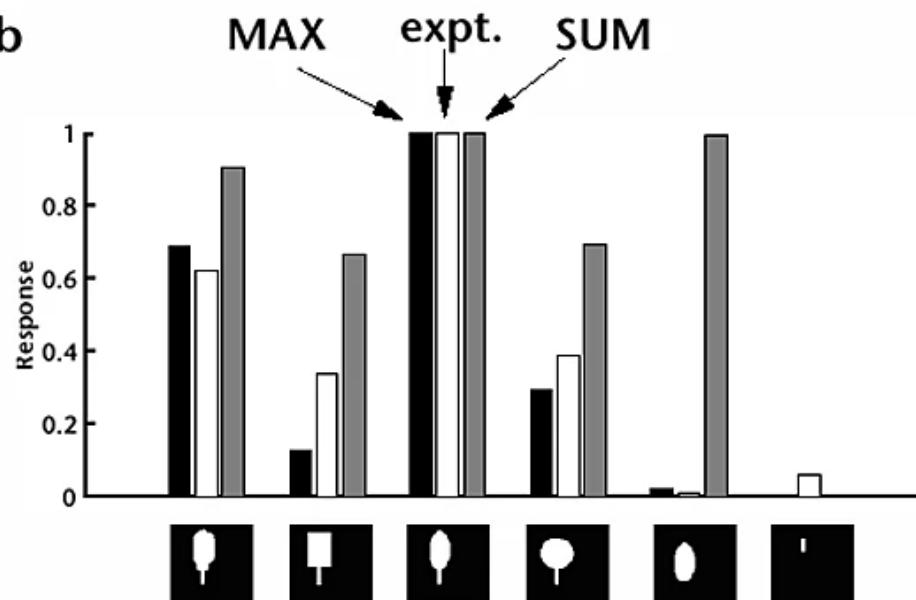


Computational Models of Vision: HMAX



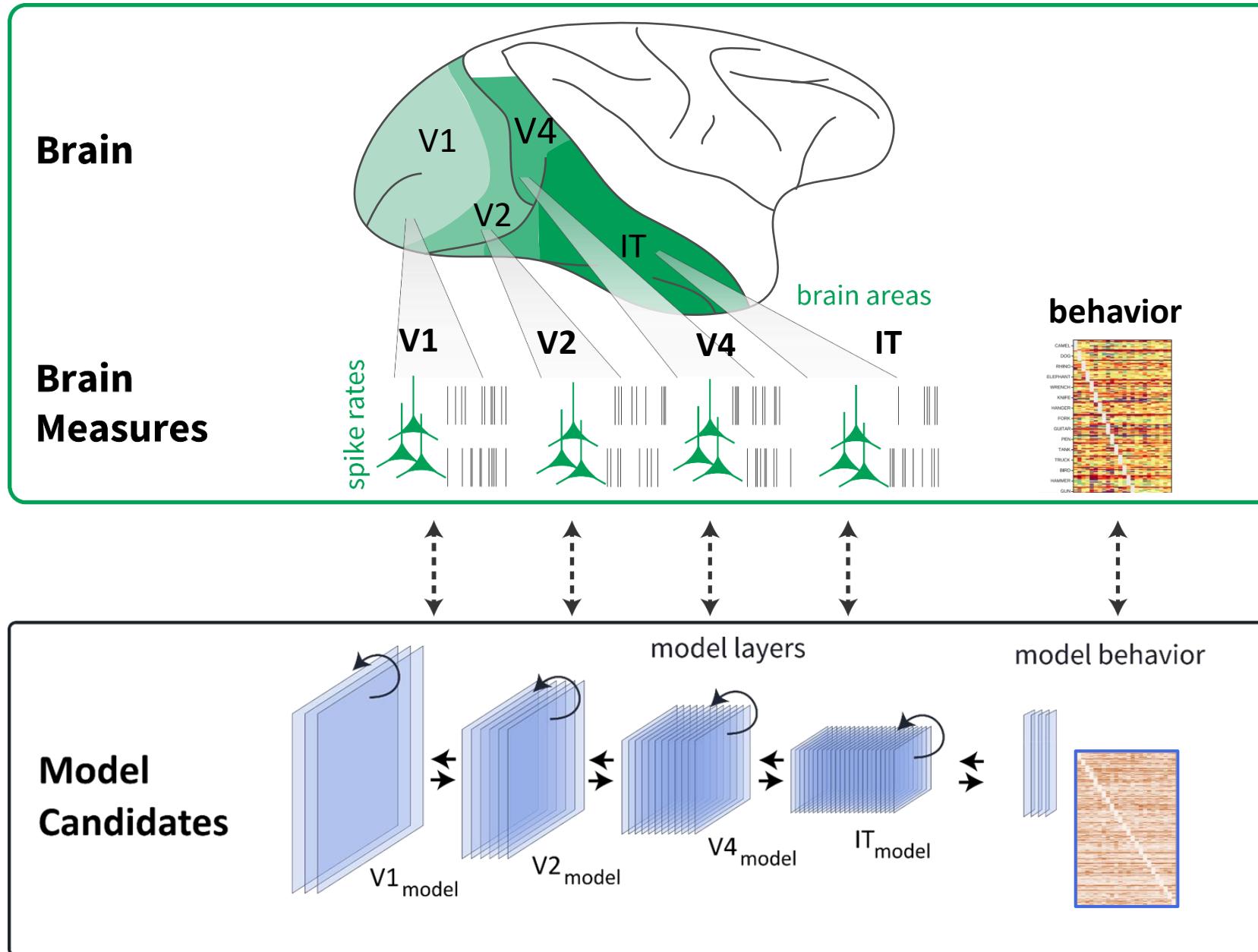
Hierarchical feedforward architecture
Simple cells: linear operations
Complex cells: nonlinear pooling (max-pool)

Replicates higher-order effects in IT



lenhuber & Poggio 1999

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?



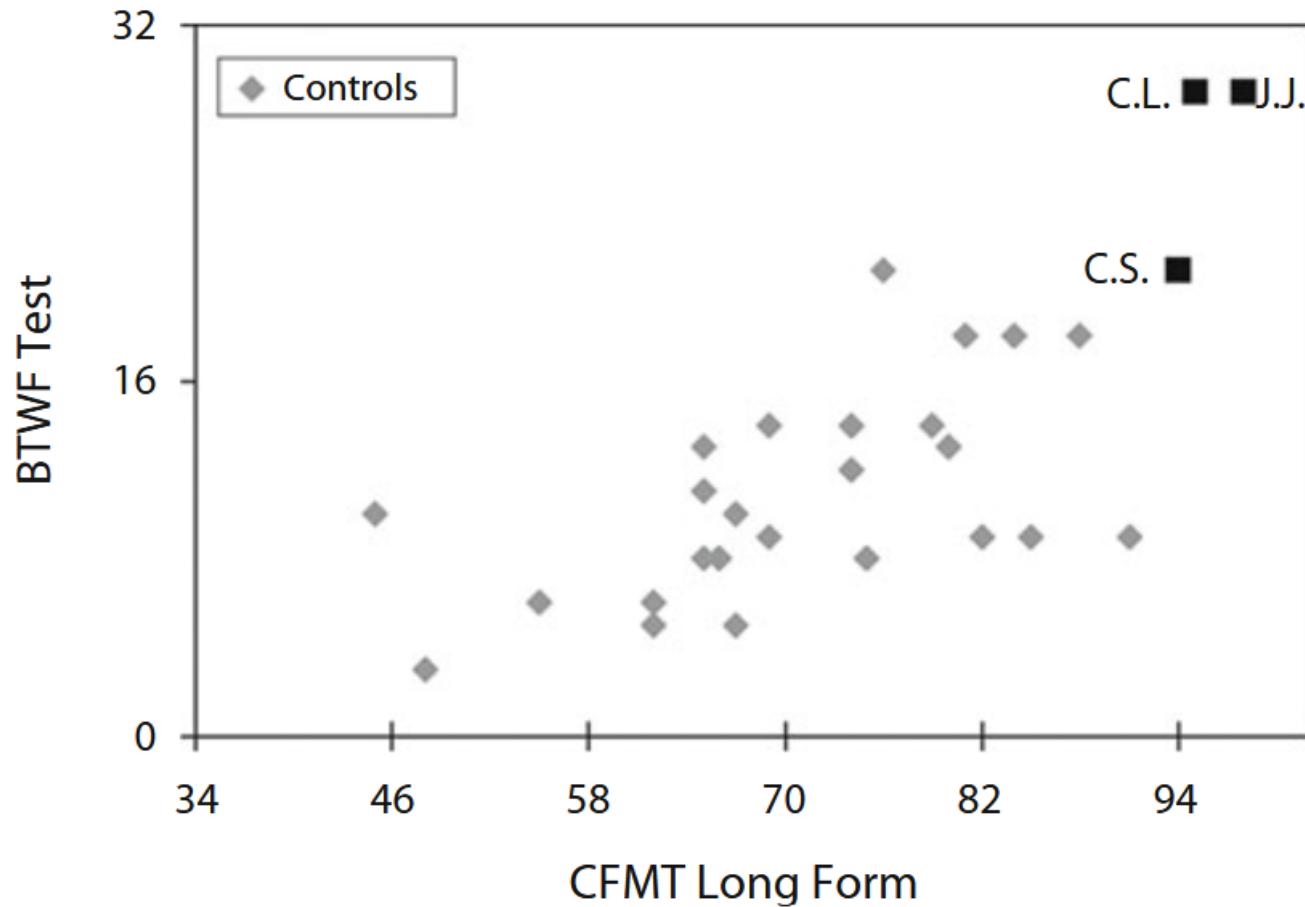
Which is easier?



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?