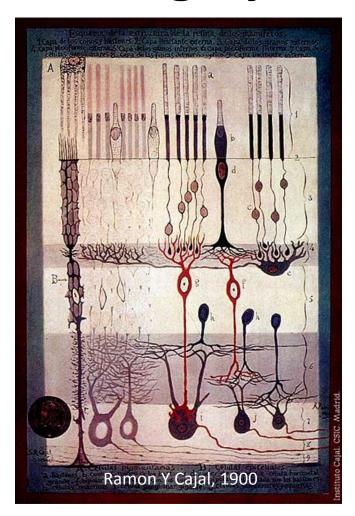
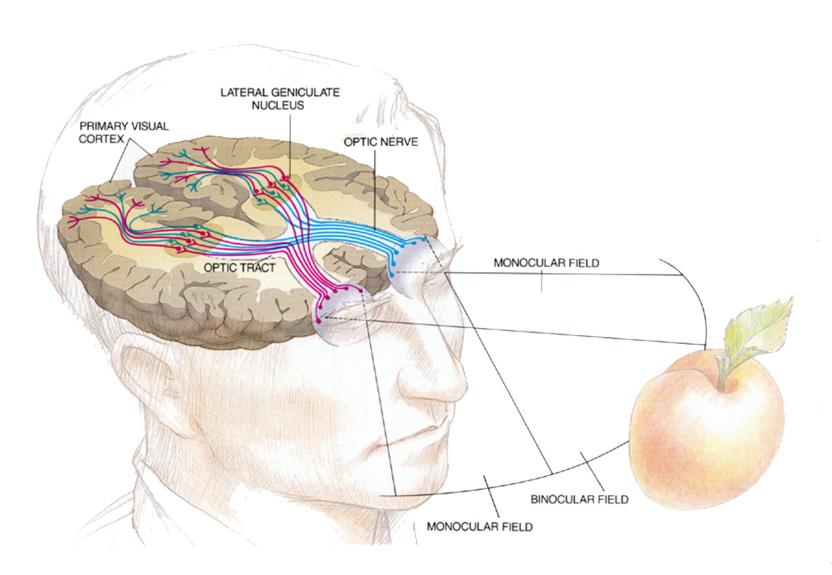
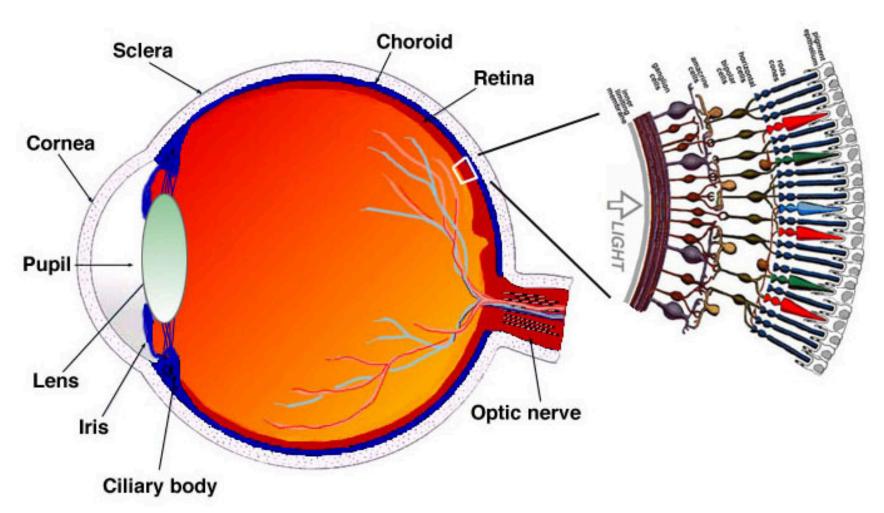
Introduction to Neuroscience: Visual Processing by the Retina



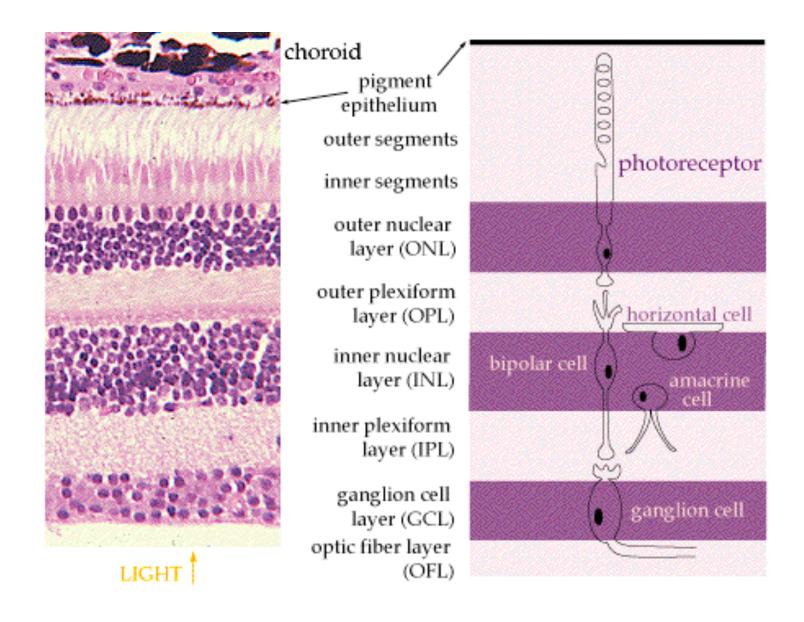
Visual processing starts in the retina, and travels through the optic nerve to the brain

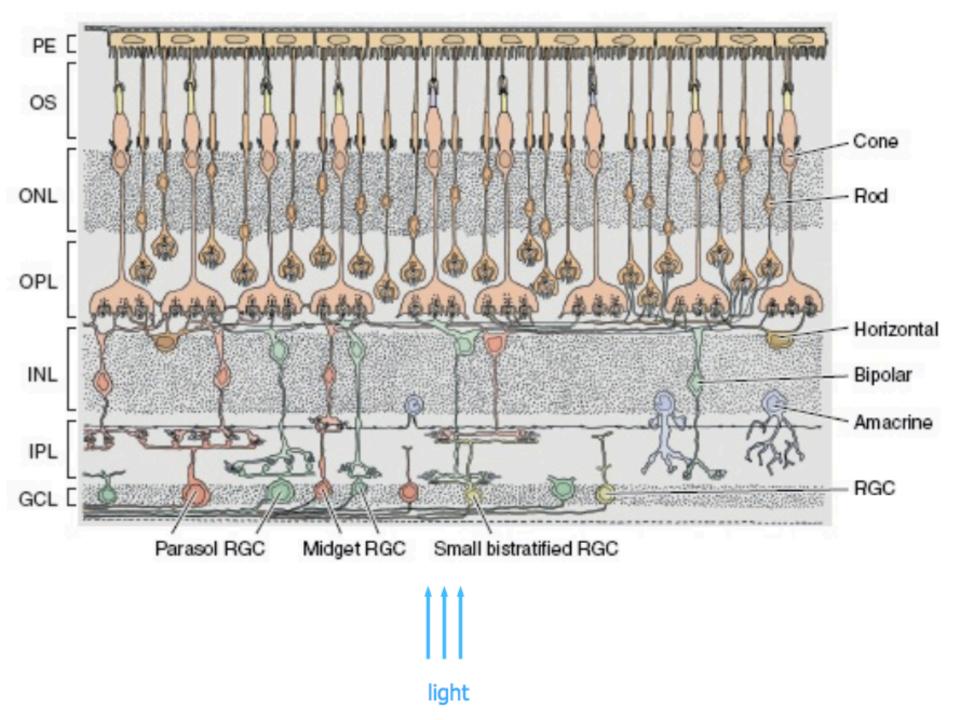


The human Eye

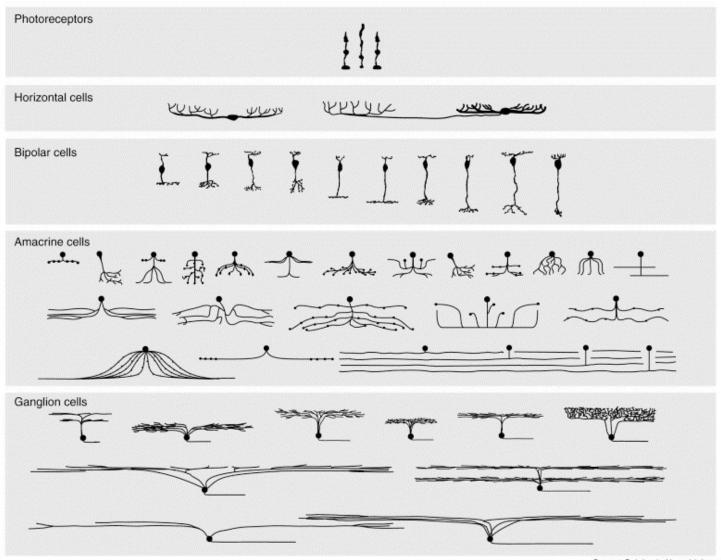


An introduction to the cell types of the retina

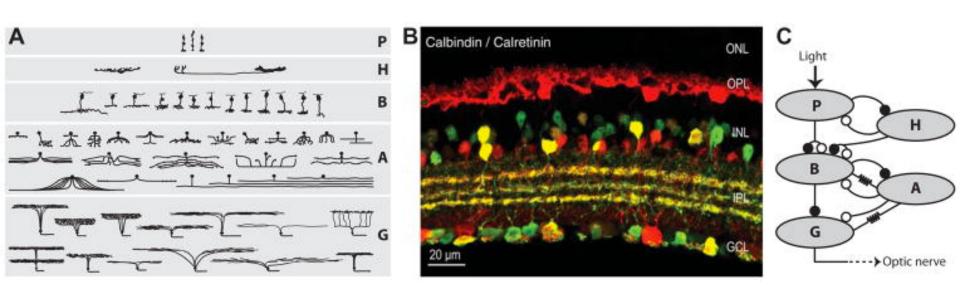




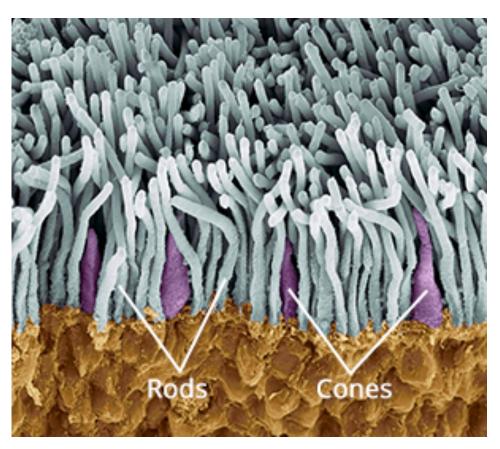
Each class is composed of multiple subtypes



Retinal neurons stratify in specific laminas of the plexiform layers

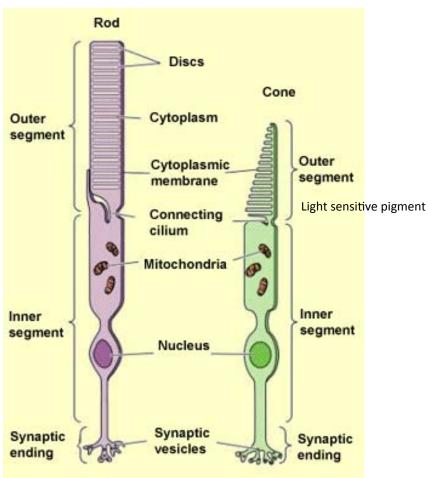


Photoreceptors: Rods and Cones

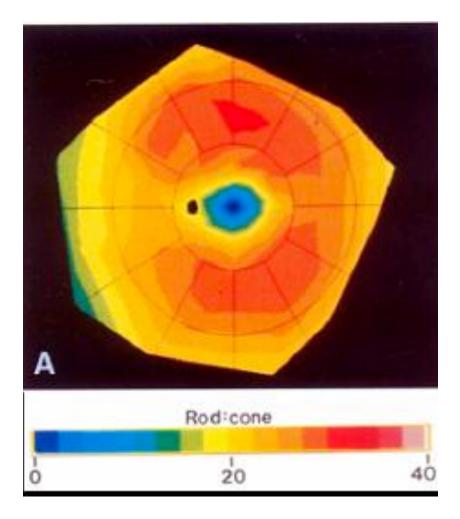


Rods: dim night vision.

Cones: bright day light vision & color vision



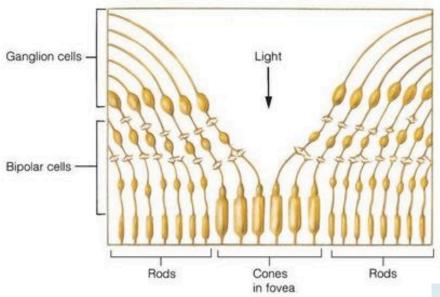
Distribution of Rods and Cones in human retina

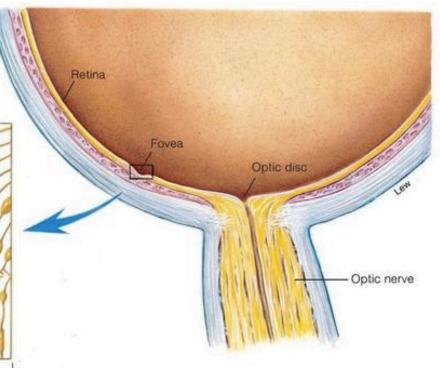


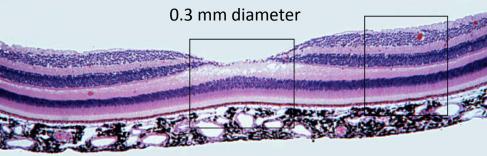
Fovea

Note: not all retinas have a fovea! Primates do.

Some species have area centralis.

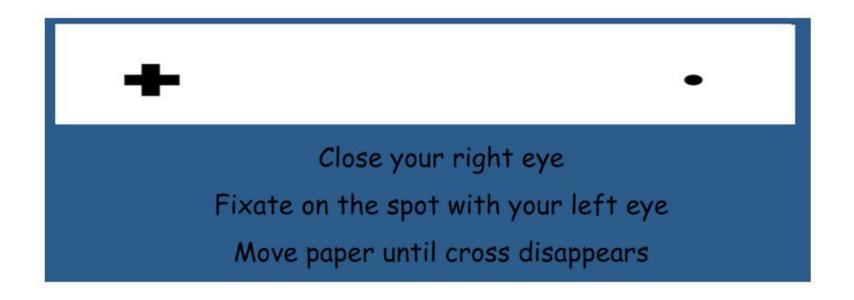




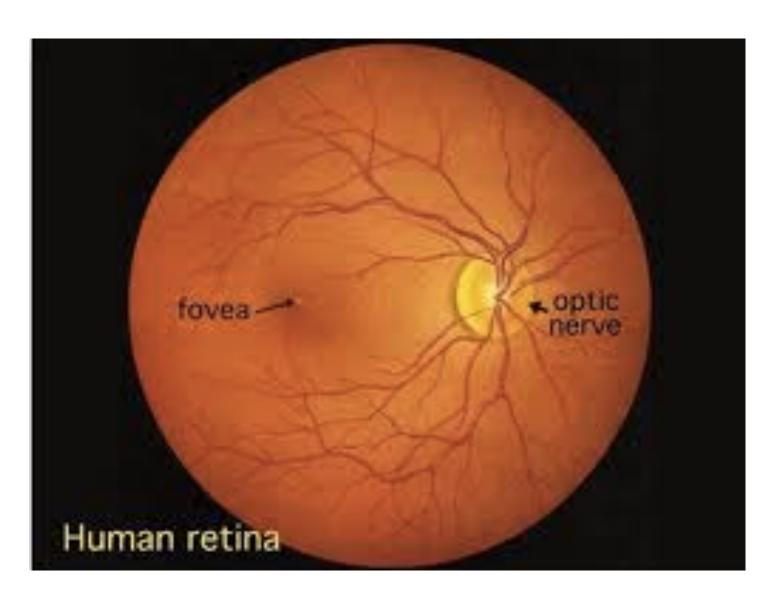


The Blind Spot

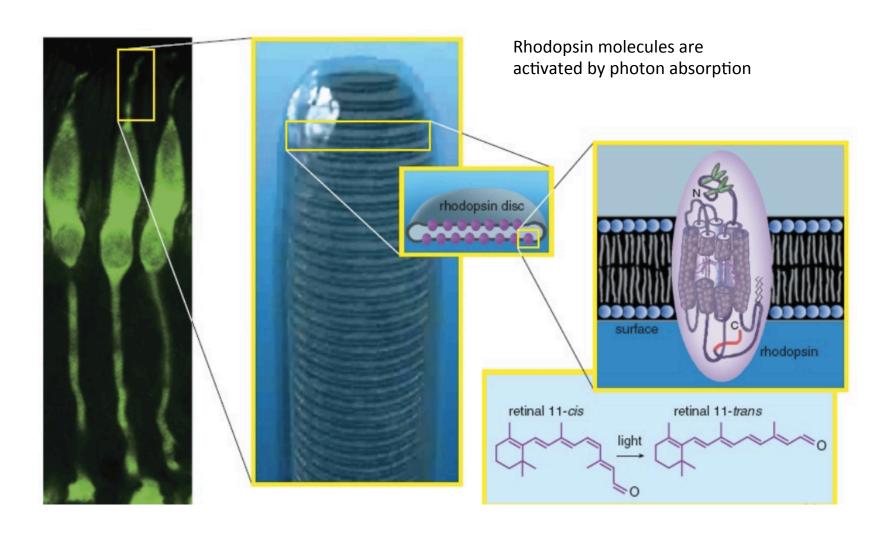
(located at the optic disc)



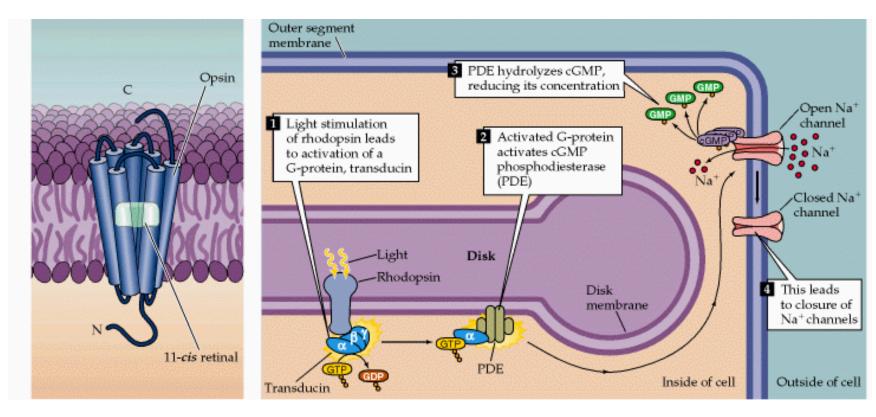
Ophthalmic view of the retina



How photoreceptors work?



Phototransduction



Rhodopsin+light -> activates the G protein transducin.

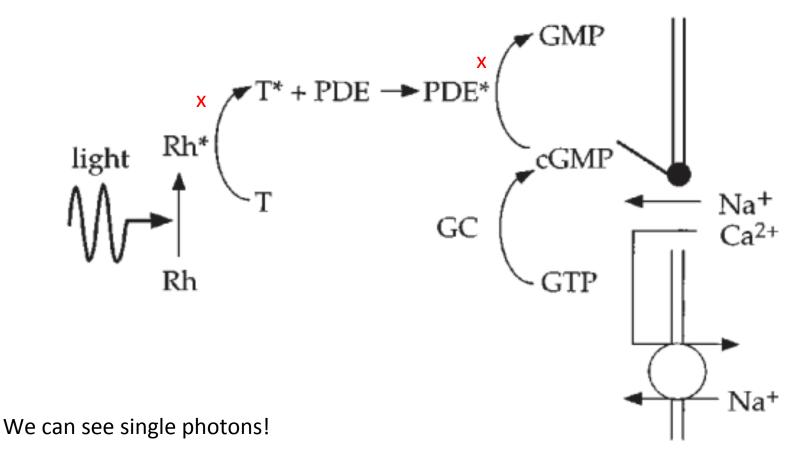
The alpha subunit of transducin releases GDP, binds GTP, and diffuses to activate PDE.

The activated PDE (phosphodiesterase) reduces cGMP levels.

Sodium channels close

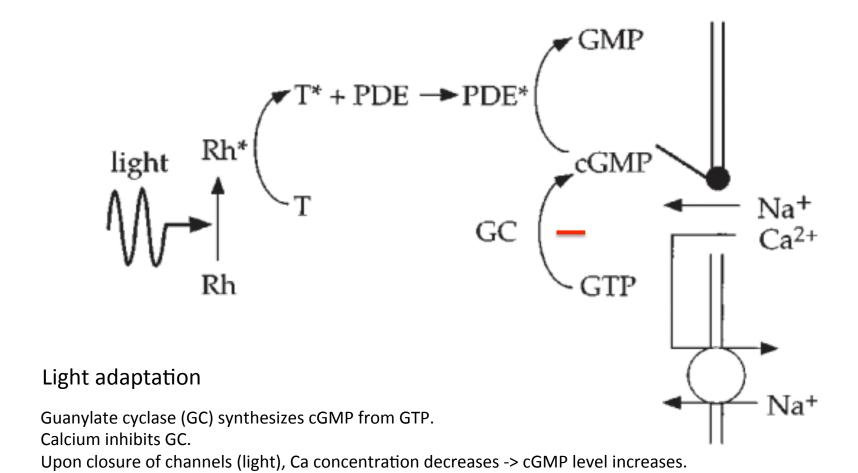
Photoreceptor hyperpolarizes

Amplifying the visual signal



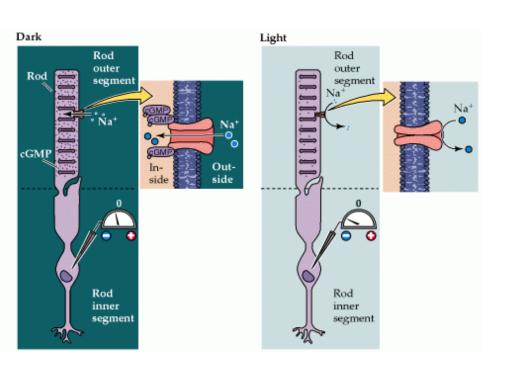
A single photoactivated rhodopsin activates ~30 transducin & PDE molecules, each PDE can break down thousands of cGMP molecules.

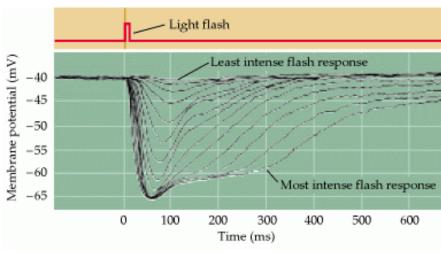
Recovery



Rieke & Baylor, 1998

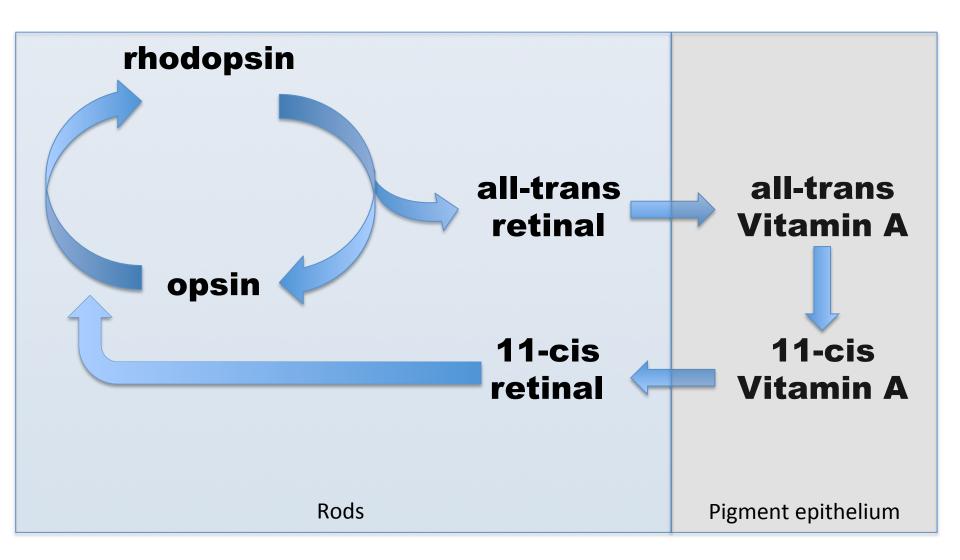
The dark current





Purves et al., 2001

The rhodopsin cycle



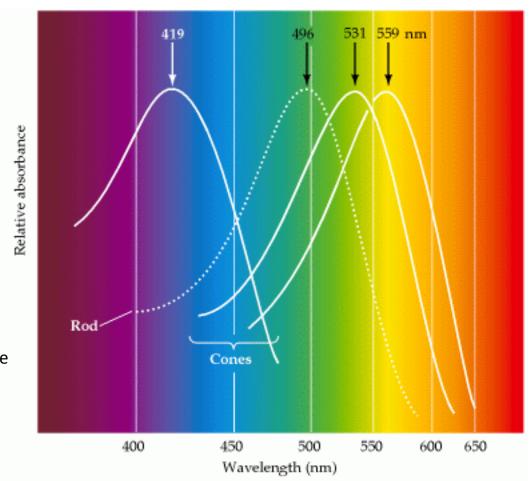
Cones mediate color vision

Color vision is based on two or more photoreceptors that bare different photopigments

- Red (L)
- Green (M)
- Blue (S)

"As it is almost impossible to conceive each sensitive point of the retina to contain an infinite number of particles..., it becomes necessary to suppose the number limited, for instance to the three principal colors."

Thomas Young, 1802



Monochromatic vision at night

Cone vision (day)

Rod vision (night)

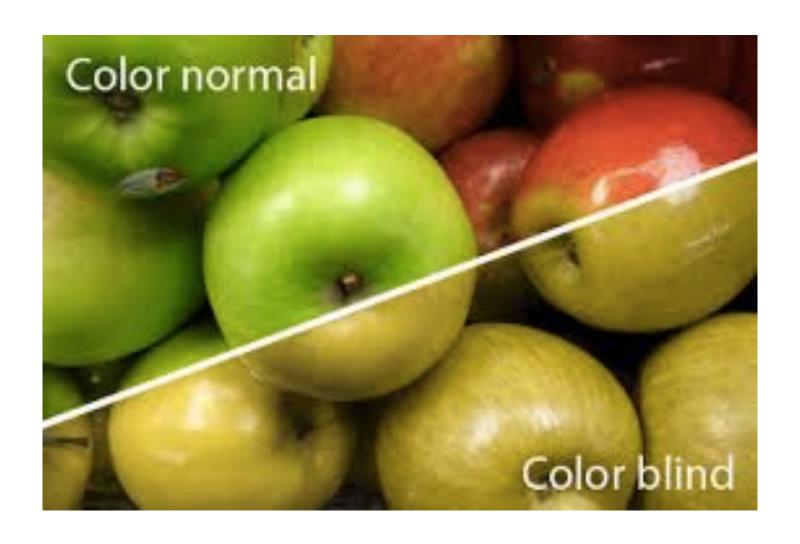




Some animals normally have monochromatic vision:



Color blindness



Visual adaptation

- Light intensities range across ~9 orders of magnitude.
 - A piece of white paper can be 1,000,000,000 times brighter in outdoor sunlight than in a moonless night.
 - If we were sensitive to this whole range all the time, we wouldn't be able to discriminate lightness levels in a typical scene.



The visual system solves this problem by restricting the 'dynamic range' of its response to match the current overall or 'ambient' light level.

Dark Indoor lighting

Dark Indoor lighting

Seattle day

Say

Seattle day

Sunny day

Seattle day

Sunny day

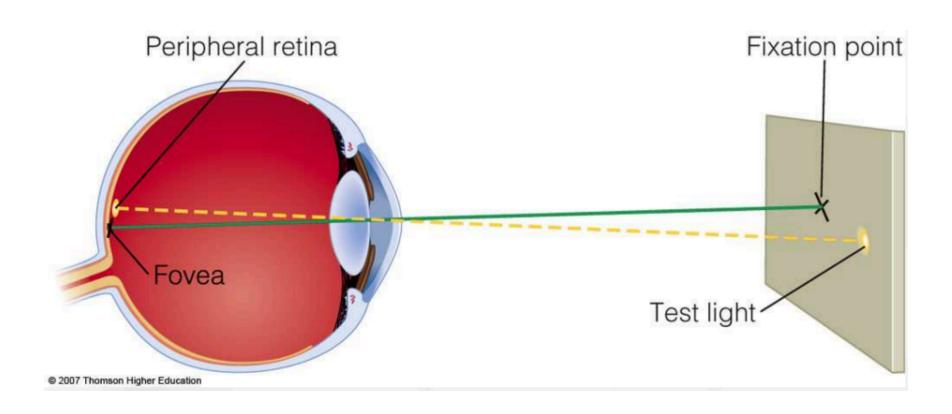
Visual adaptation

- Already at the retina!
 - Pupil's diameter: 2-8 mm.
 - Rods and cones two visual systems.
 - Both rods and cones adapt becomes less sensitive as light levels increase.

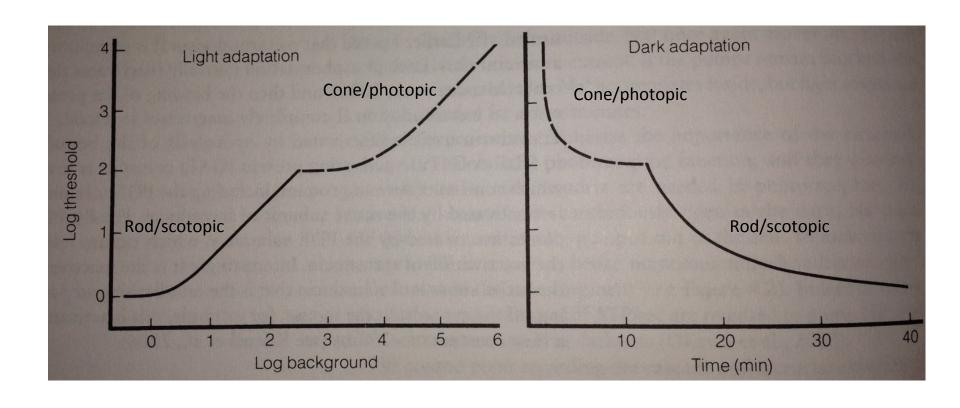


Craik & Vernon, 1941: Pressure blind experiments.

Psychophysical Measurement of Light/Dark Adaptation

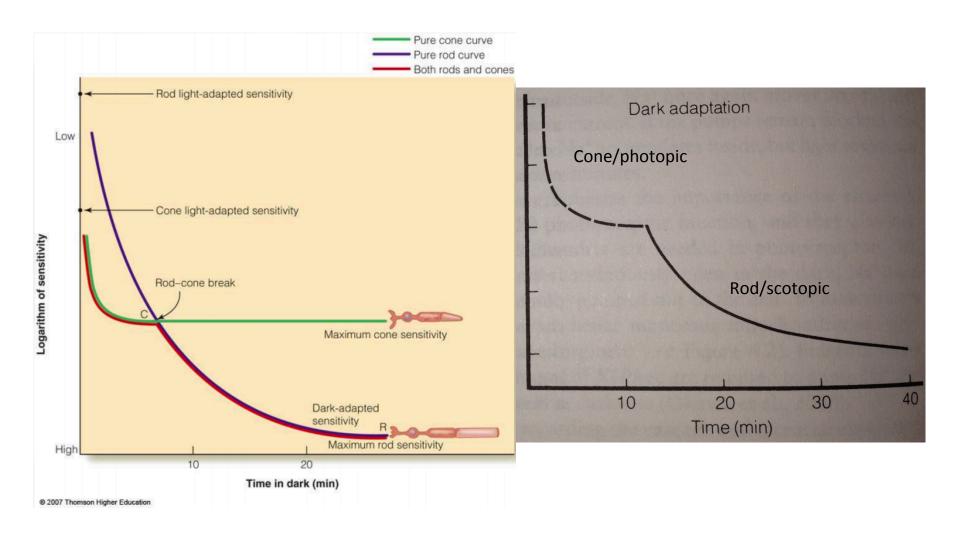


Light and dark adaptation

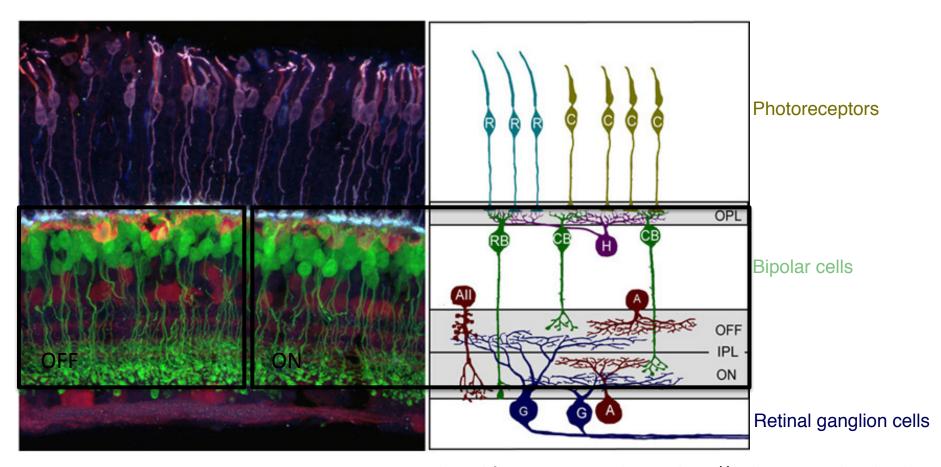


Weber law:
$$C = \frac{\Delta I}{I}$$
 $\frac{11-10}{10} = \frac{110-100}{100}$

Light and dark adaptation

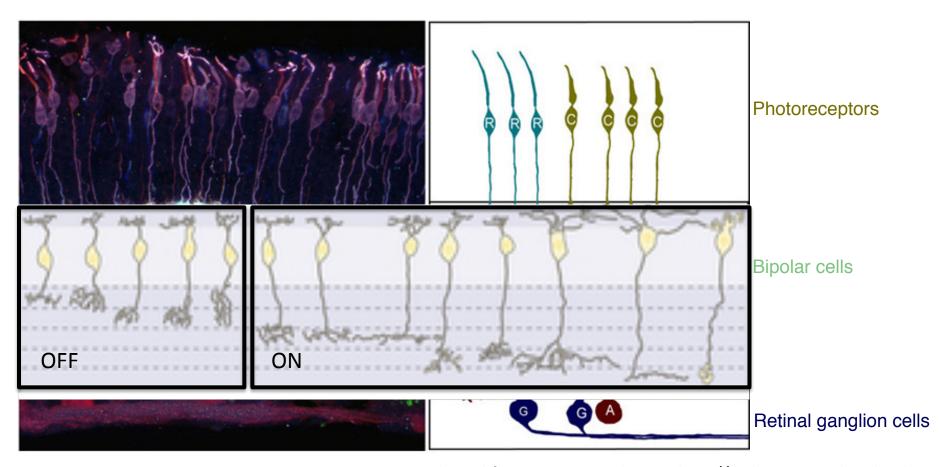


Bipolar Cells



Adapted from Morgan and Wong http://webvision.med.utah.edu And from Euler et al. 2014 Nature reviews Neuroscience

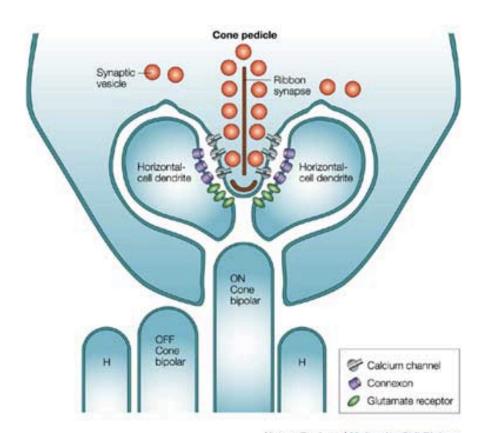
Bipolar Cells



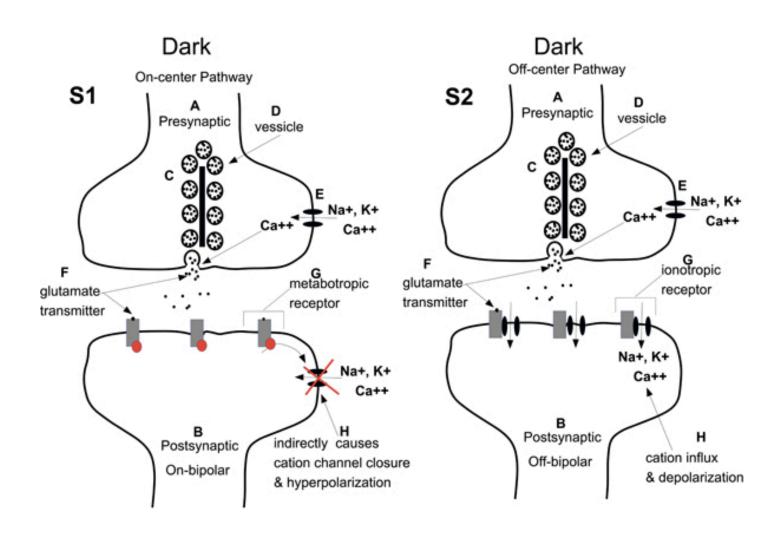
Adapted from Morgan and Wong http://webvision.med.utah.edu And from Euler et al. 2014 Nature reviews Neuroscience

Ribbon synapse

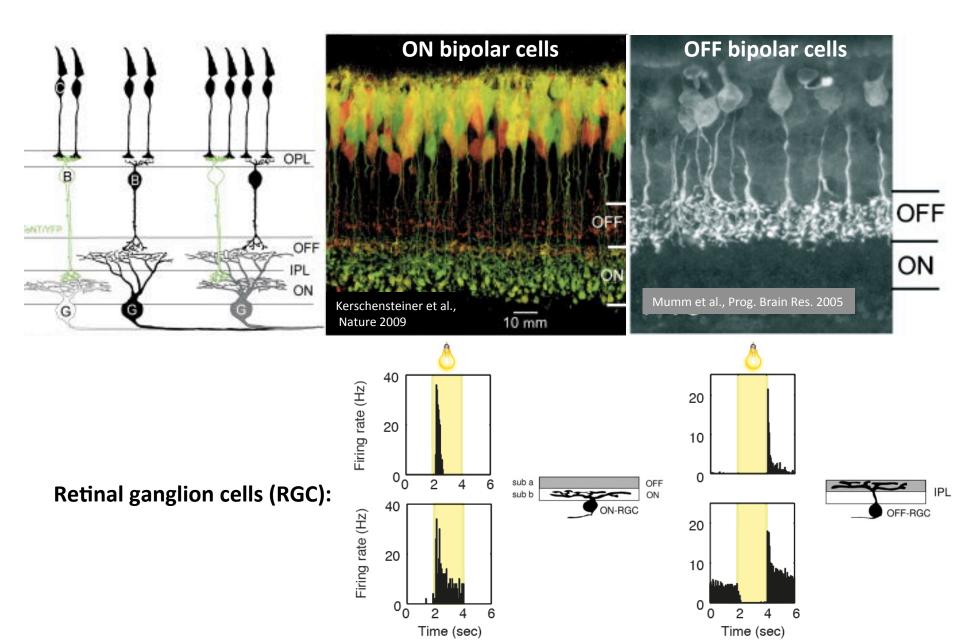
- Graded neurotransmitter release.
 Precise, sustained, and rapid.
- The synaptic ribbon releases
 100s-1000s vesicles per second.
- Each pre-synaptic cell has 10-100 ribbons.
- Requires a large pool of readily releasable vesicles.



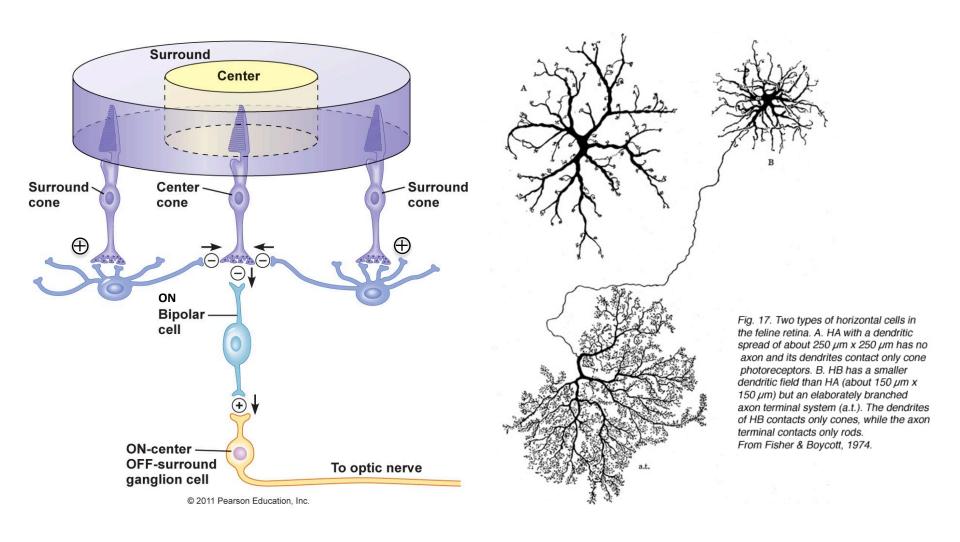
On- and Off-bipolar cells



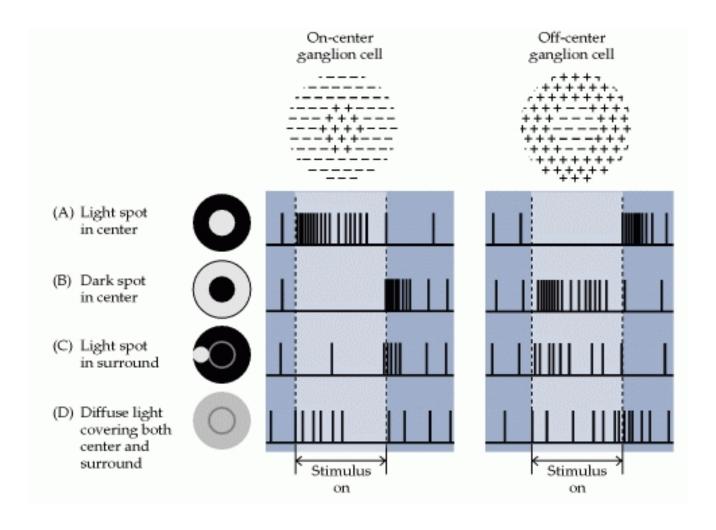
On and Off retinal pathways



Horizontal Cells



Center-surround organization of receptive fields

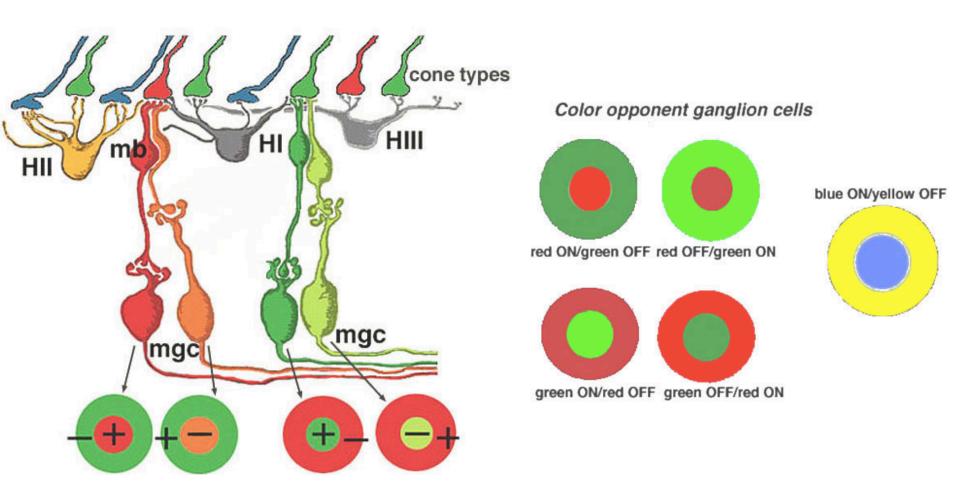


Hartline (1938) & Granit (1947) – On, Off; Barlow & Kuffler (1950) – antagonistic center-surround.

Retinal neurons signal relative intensity of stimulation



Midget ganglion cells

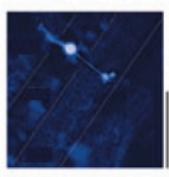


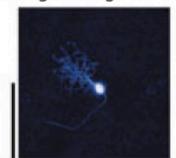




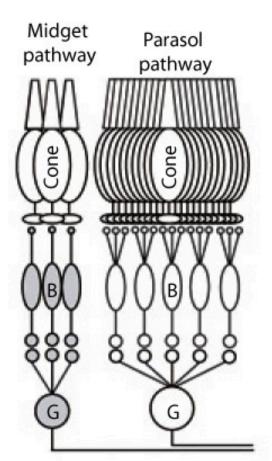
Convergence and acuity







Midget Ganglion Cell



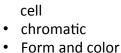
Diffuse Bipolar Cell = not selective for color



Parasol Ganglion Cell



- Large dendritic arbors
- Large receptive fields
- ~10% of ganglion cells
- Non-chromatic
- Gross features of stimulus
- movement



Small dendritic

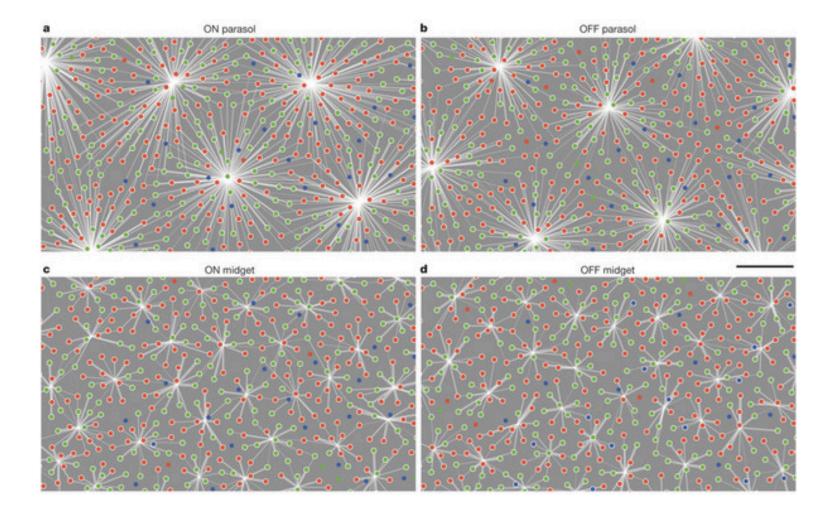
Small receptive

• ~80% of ganglion

arbors

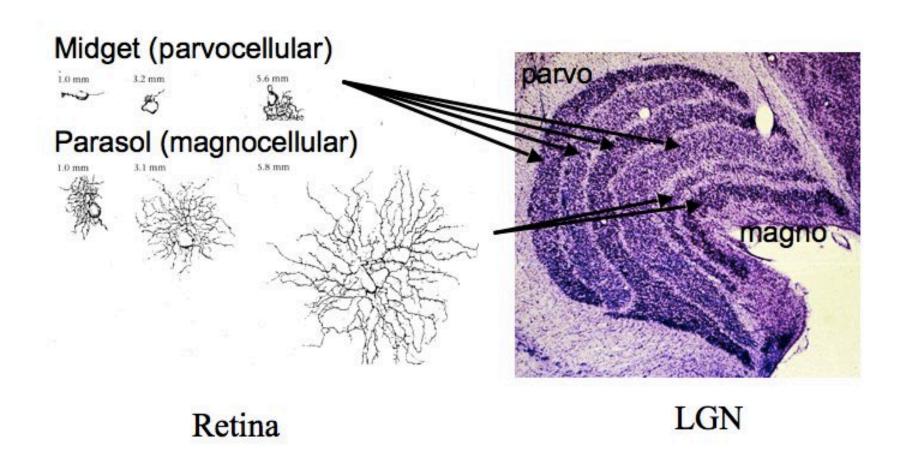
fields

Fine details

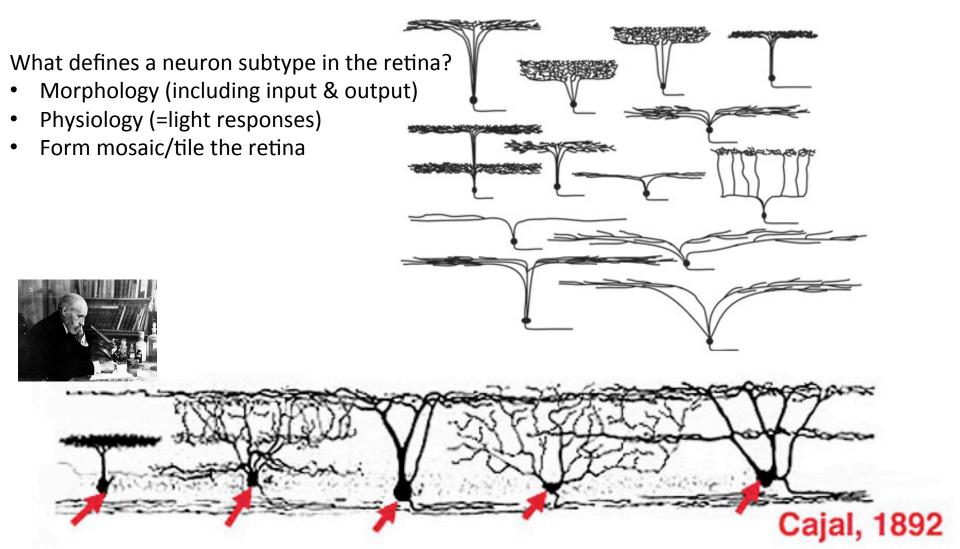


Field et al., 2010

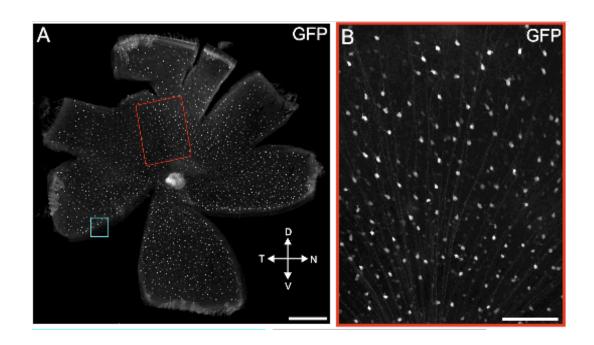
Magno and Parvo pathways



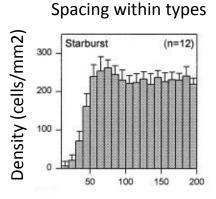
Retinal ganglion cells



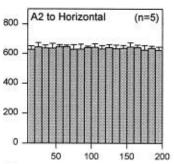
Mosaic organization



- Cover the entire visual field
- Exclusion zone for same cell type

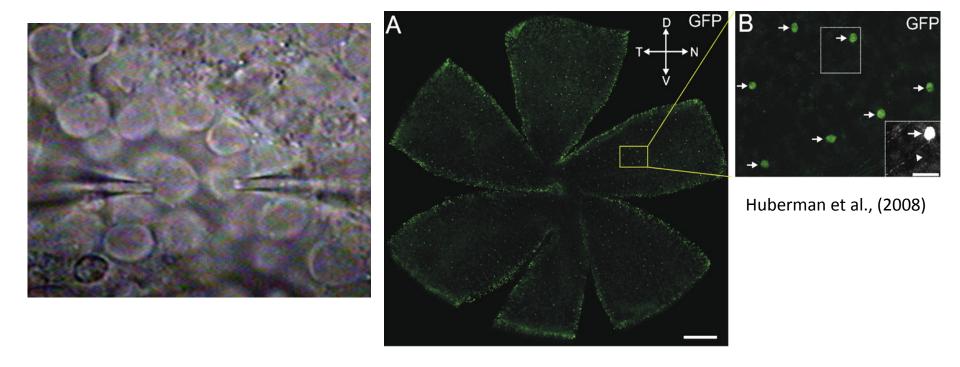


Spacing between types

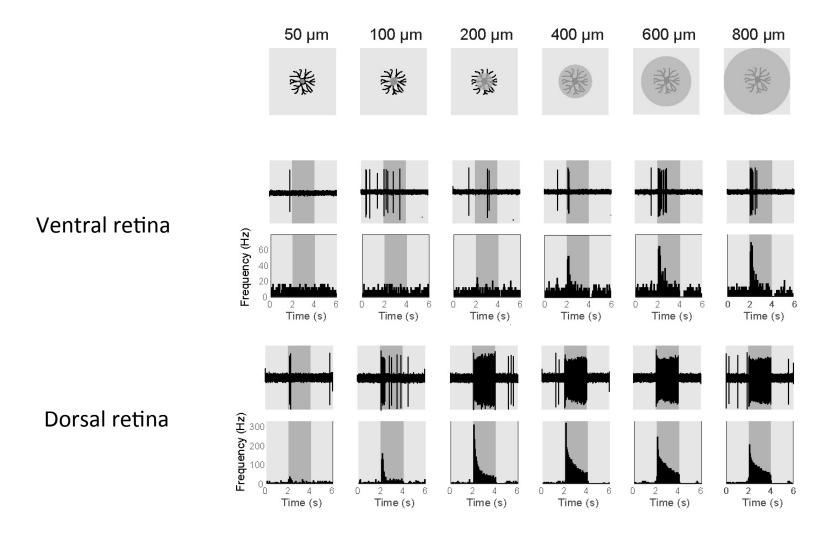


Distance from reference cell

Two-photon targeted recordings from transgenic mouse lines

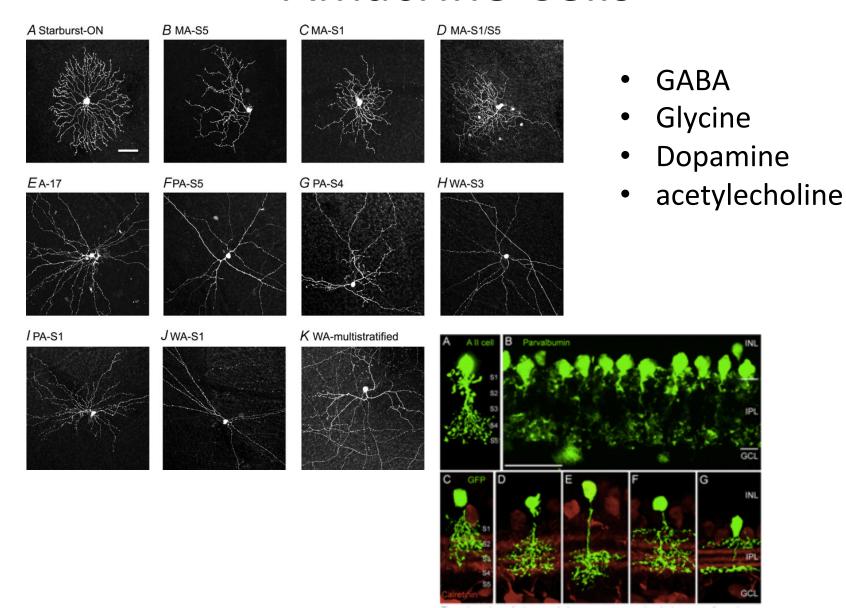


Transient-OFF-α-retinal ganglion cells display different response properties at different retinal locations

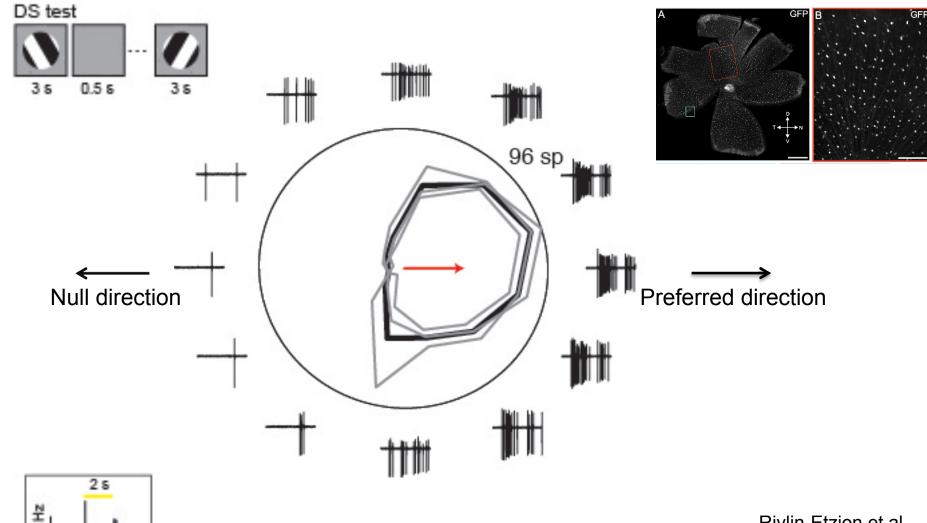


Recent evidence from mouse retina reveal some of the accepted concepts may be overruled. For example, transient-Off-alpha RGCs show different response properties in different retinal locations

Amacrine Cells

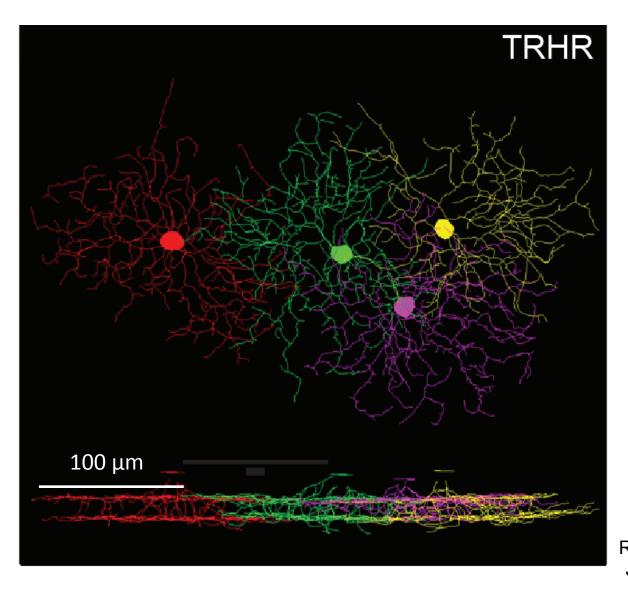


Direction selective retinal ganglion cell has preferred and null directions



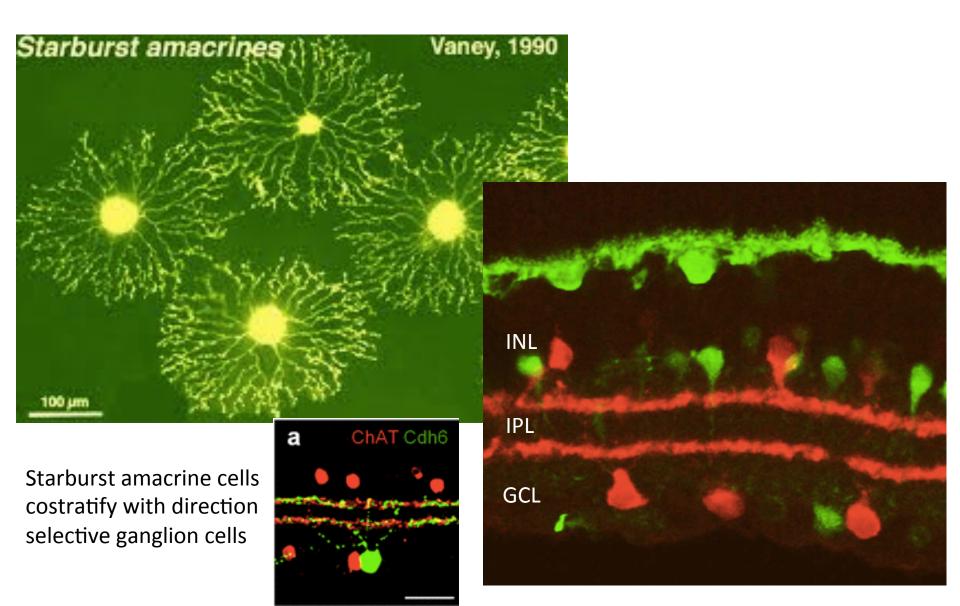
Rivlin-Etzion et al., J Neurosci, 2011

On-Off Direction Selective Ganglion Cells

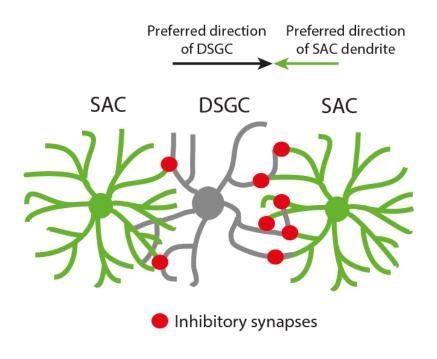


Rivlin-Etzion et al., J Neurosci, 2011

Starburst amacrine cells

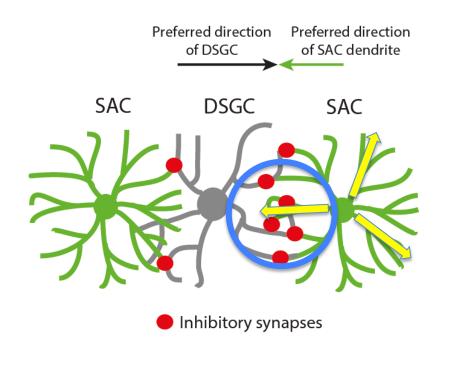


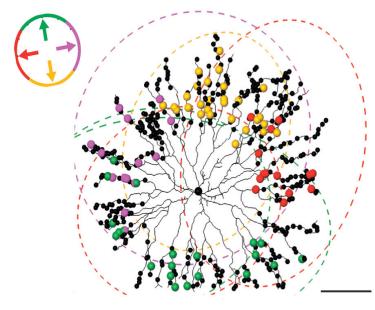
Directional responses are mediated by asymmetric inhibition from starburst amacrine cells





Directional responses are mediated by asymmetric inhibition from starburst amacrine cells



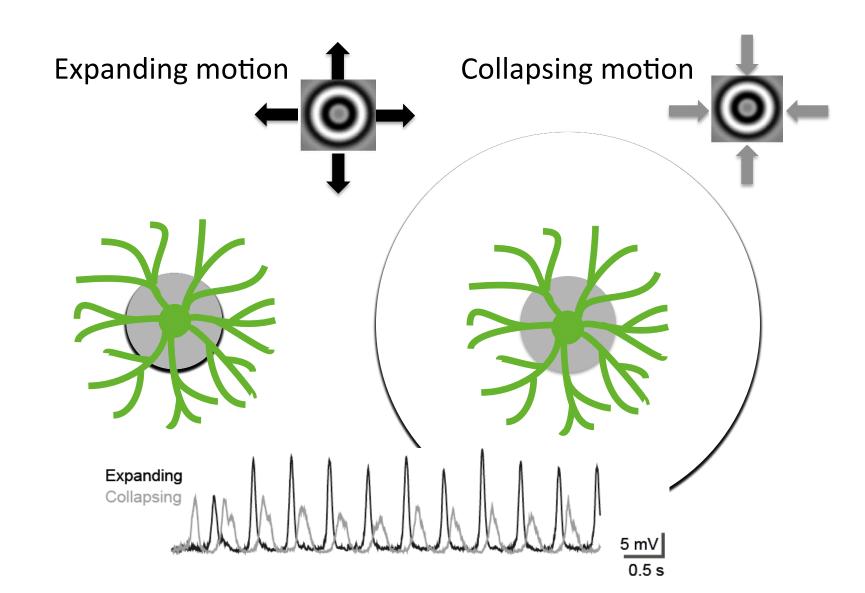


Brigmann & Denk 2011



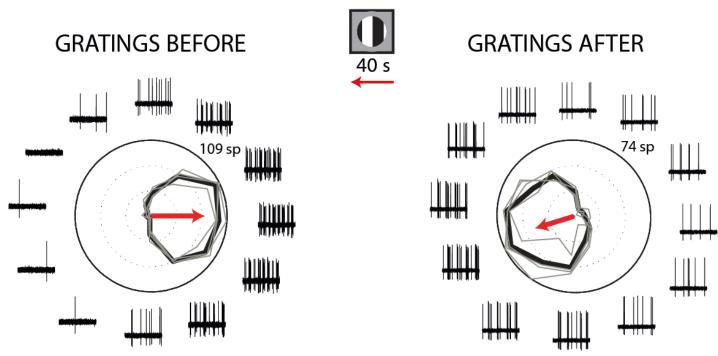
Direction selective Starburst amacrine

SAC processes prefer centrifugal (=expanding) motion



On-Off DSGCs reverse their directional preference upon visual stimulation

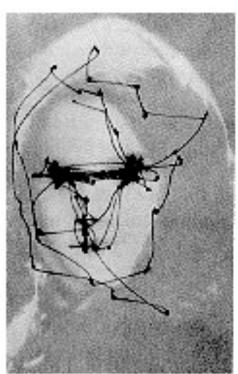




Recent evidence from mouse retina reveal some of the accepted concepts may be overruled. For example, following a specific adaptation protocol, DSGC overcome the anatomy of the circuit and reverse their directional preference.

The visual image is not stable on the retina





saccades

Some computations performed by RGCs

- Texture motion: RGCs that respond to high spatial frequencies moving in all directions.
- Object motion-sensitive (OMS) ganglion cells fire only when a local patch on the receptive field center moves with a trajectory different from the background.
- Approaching motion.
- Prediction of location of a moving object.
- Omitted stimulus response: after exposure to a periodic stimulus, if one stimulus is omitted, some neurons generate a pulse of activity at the time corresponding to the missing stimulus.
- Saccadic suppression.

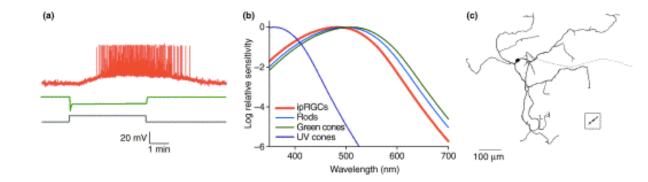
Review: Gollisch & Meister 2010

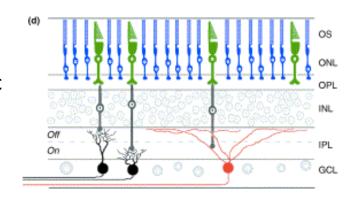
Retina: advantages

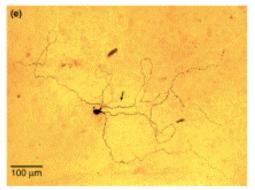
- Simple system, but performs complex computations.
- Organized structure.
- Full control on its input.
- Approachable for recordings/imaging.
- Similar anatomical arrangement in all mammals.

Intrinsic photosensitive RGCs (ipRGCs)

- Express the visual pigment melanopsin
- Mice, rats, cats, monkeys, human
- Light responses develop slowly
- Sustained depolarization to light
- Slow recovery
- Project to the suprachiasmatic nucleus – controls the circadian clock
- Projects to the pretectal nucleus, which controls the pupillary light reflex







TRENOS in Neurosciences

Visual acuity

- The human visual system is capable of acuity of 1 min of arc or 60 cycles/degree of visual angle.
- One degree of visual angle is thought to cover approximately 280-300 μm of retinal distance.

One factor that determines visual acuity is the size of the eye: The larger the eye, the bigger the area that covers a given object in the visual field.

The bigger the area, the more neurons are involved in encoding of the object...

