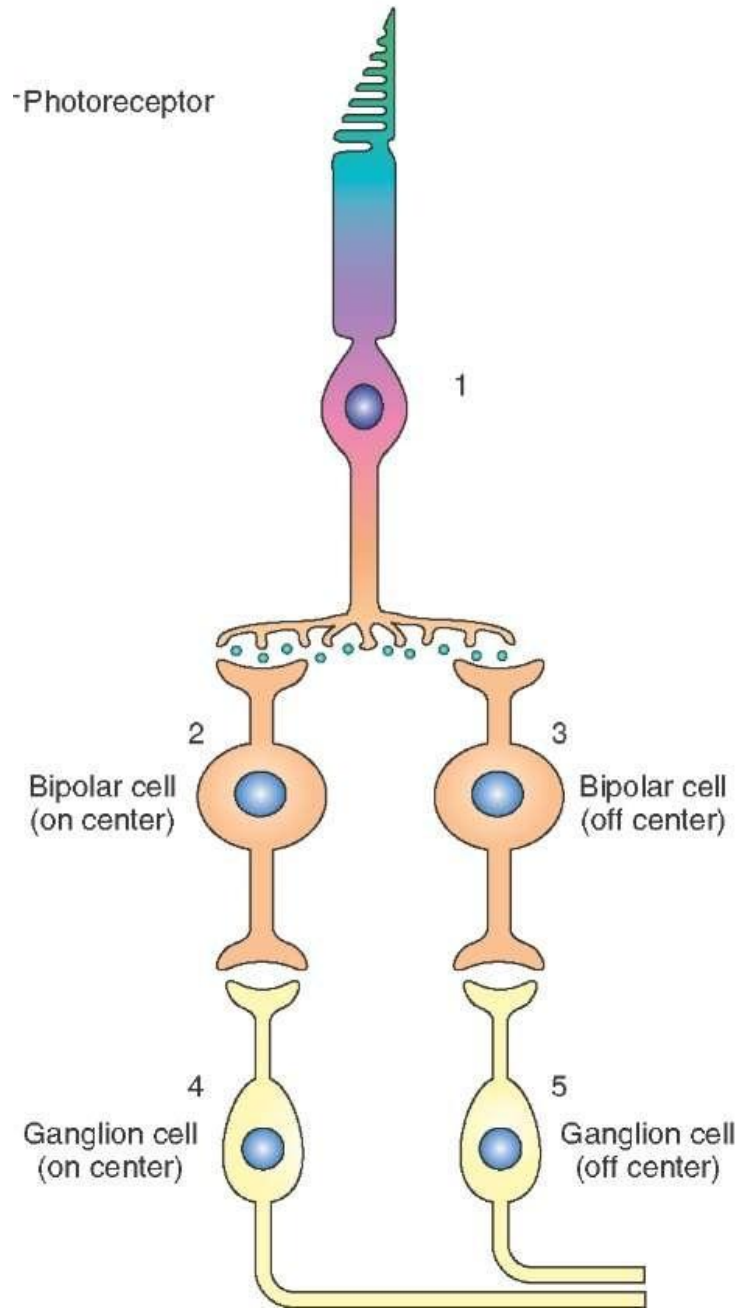


(more) vision

COGS17 - WEEK 4
7/22/19

“it doesn’t matter what one cell does, it matters HOW they are connected”

Review of the potential connectivity patterns of photoreceptors, bipolar cells, and retinal ganglion cells

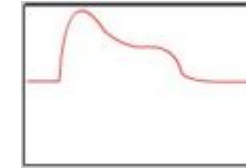


ON center vs OFF center bipolar cells

What happens when the photoreceptor is...

In the light:

ON → depolarizes

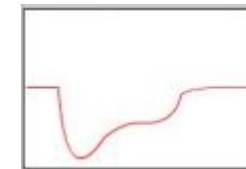


OFF → hyperpolarizes

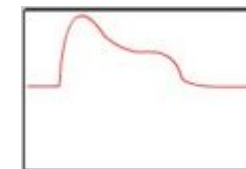


In the dark:

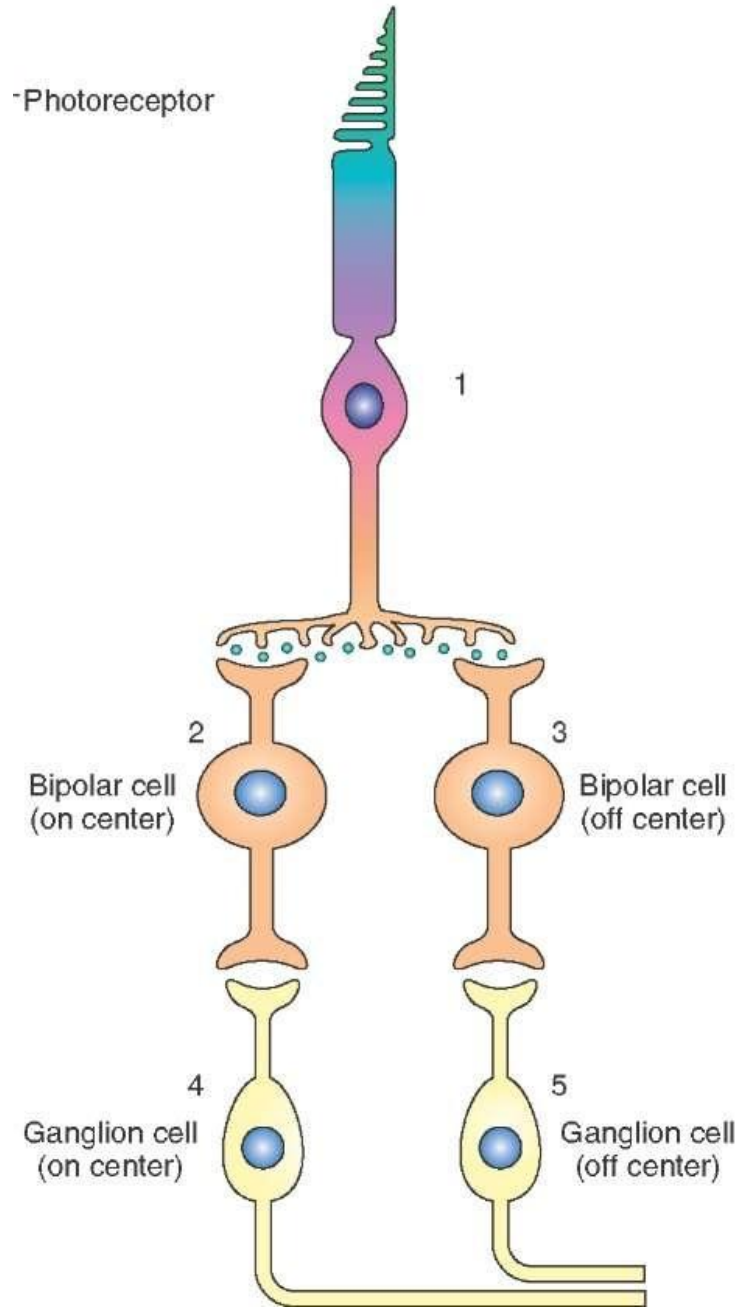
ON → hyperpolarizes



OFF → depolarizes



why?



ON = metabotropic receptor (mGluR6, g-protein coupled!)

OFF = ionotropic receptor (AMPA)

glutamate has opposite effects on these receptors

In the light:

ON → less glutamate around, so voltage-gated Na⁺ channels are open

OFF → less glutamate around, so less Na⁺ enters via channels

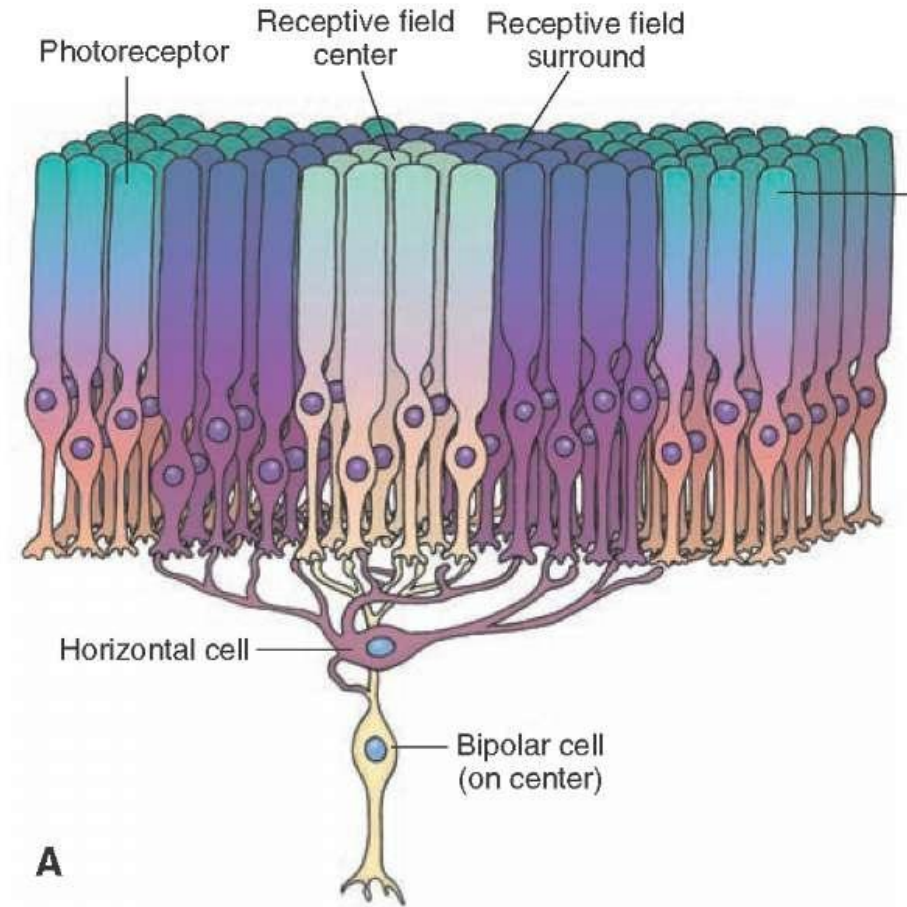
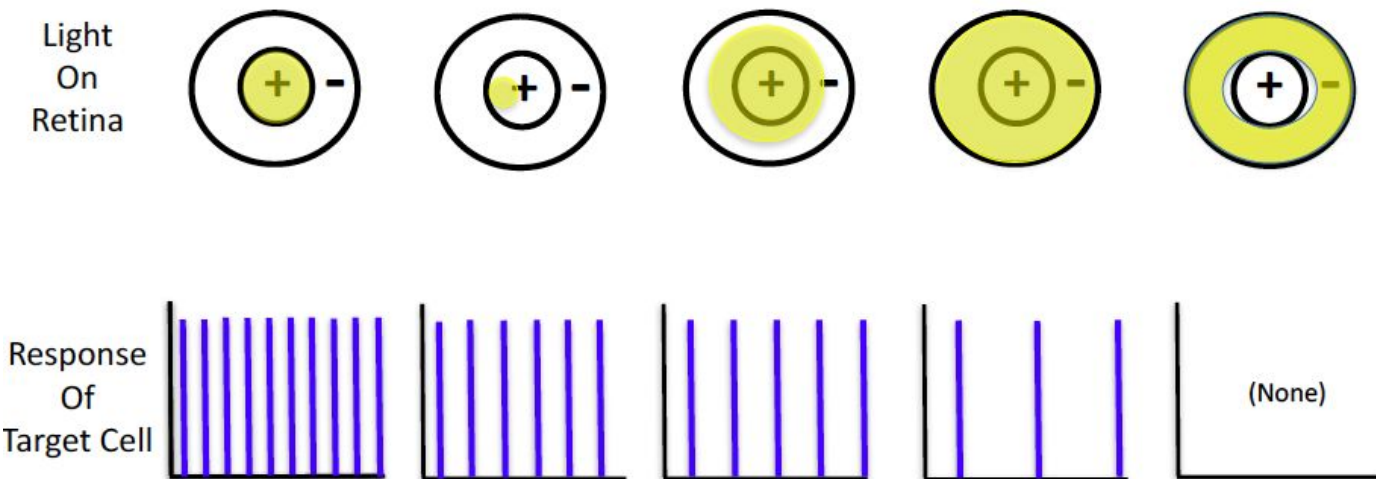
In the dark:

ON → more glutamate, so voltage-gated Na⁺ channels are closed

OFF → more glutamate, so more Na⁺ enters via channels

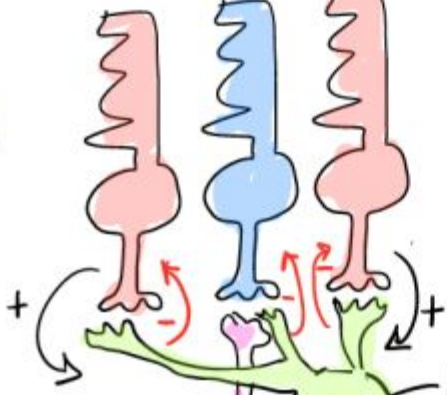
Center-Surround

- Excitatory center and inhibitory surround receptive fields
- Refers to receptive fields for cells in a place on the retina
- Plus and minus refer to what the target cell is doing
- Receptive fields are overlapping thus many receptor cells contribute to many receptive fields



Mach

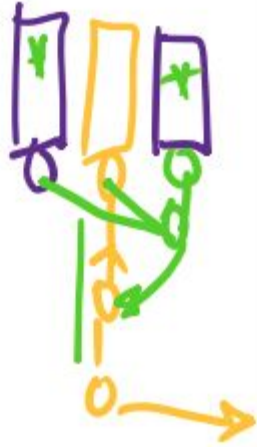
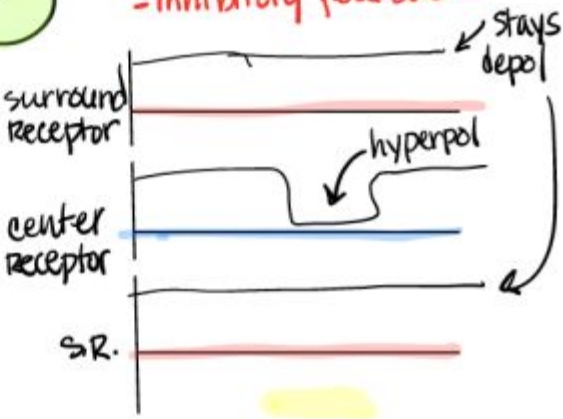
surround center surround



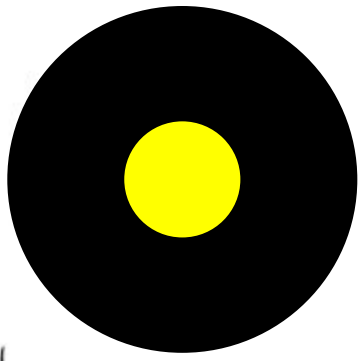
ON BIPOLAR

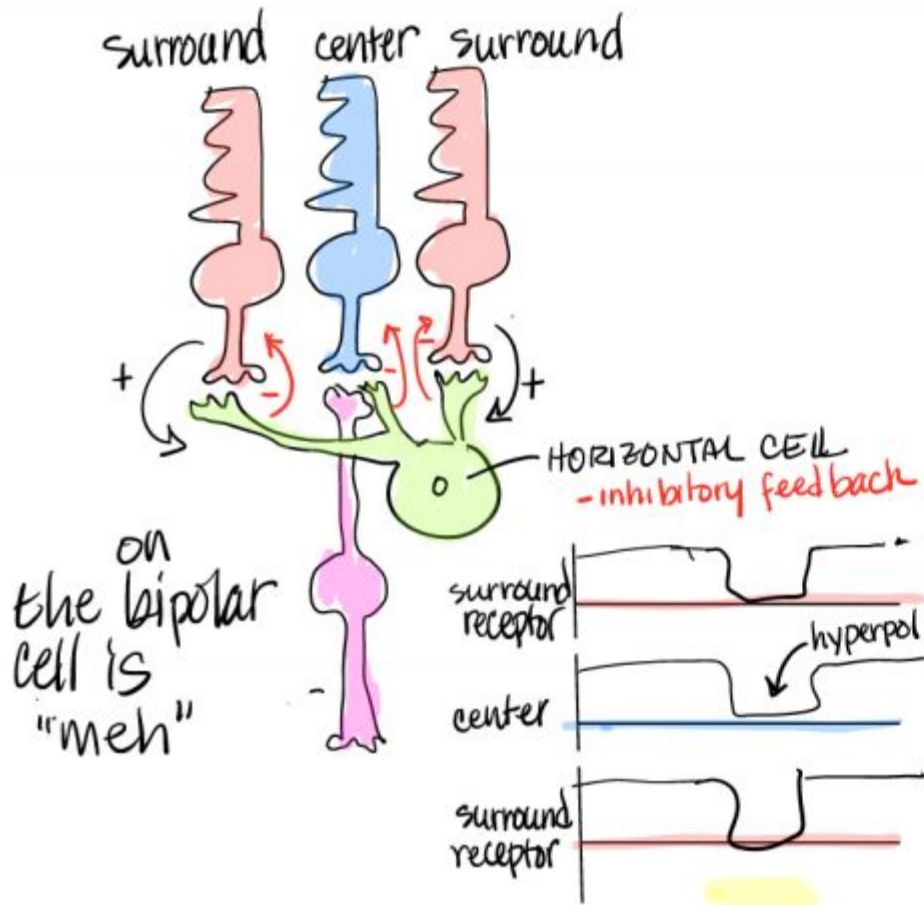
HORIZONTAL CELL -inhibitory feedback

maximally excited



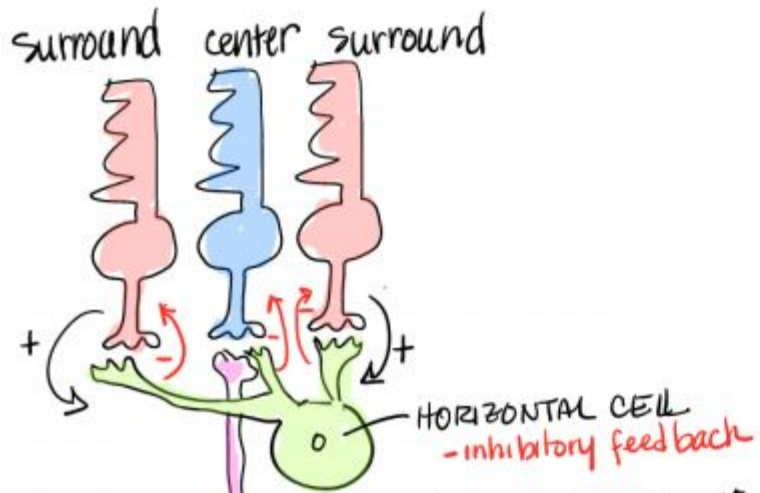
* center cone releases less xmttr due to horizontal inhibition



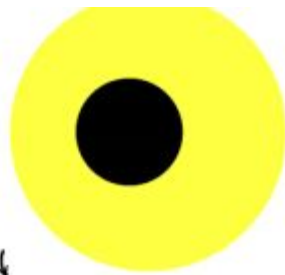
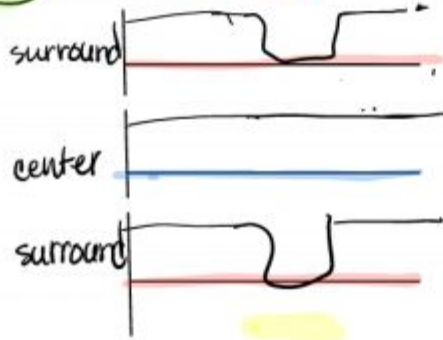


*horizontal cell is less excited
 -inhibition to center receptor is reduced

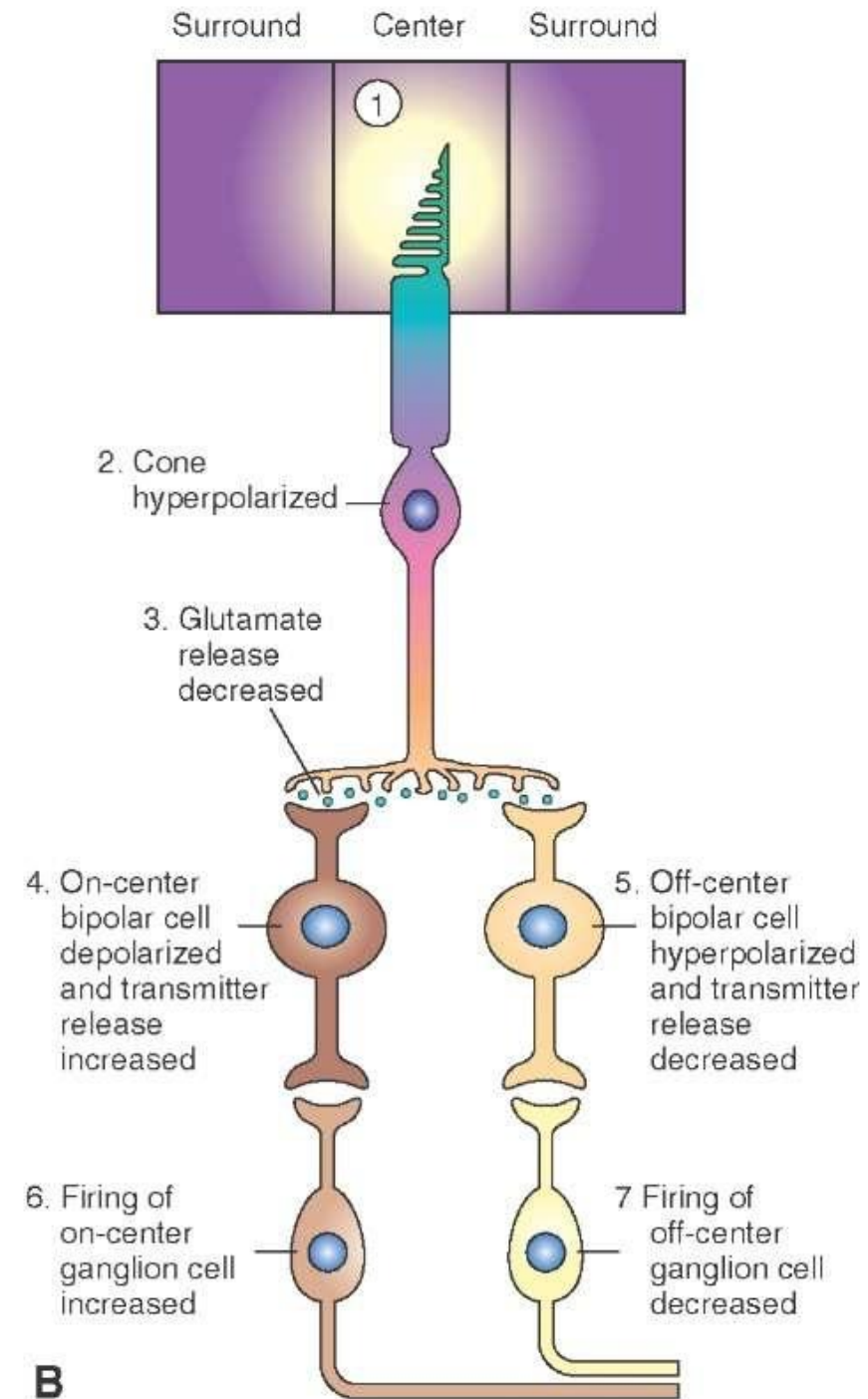
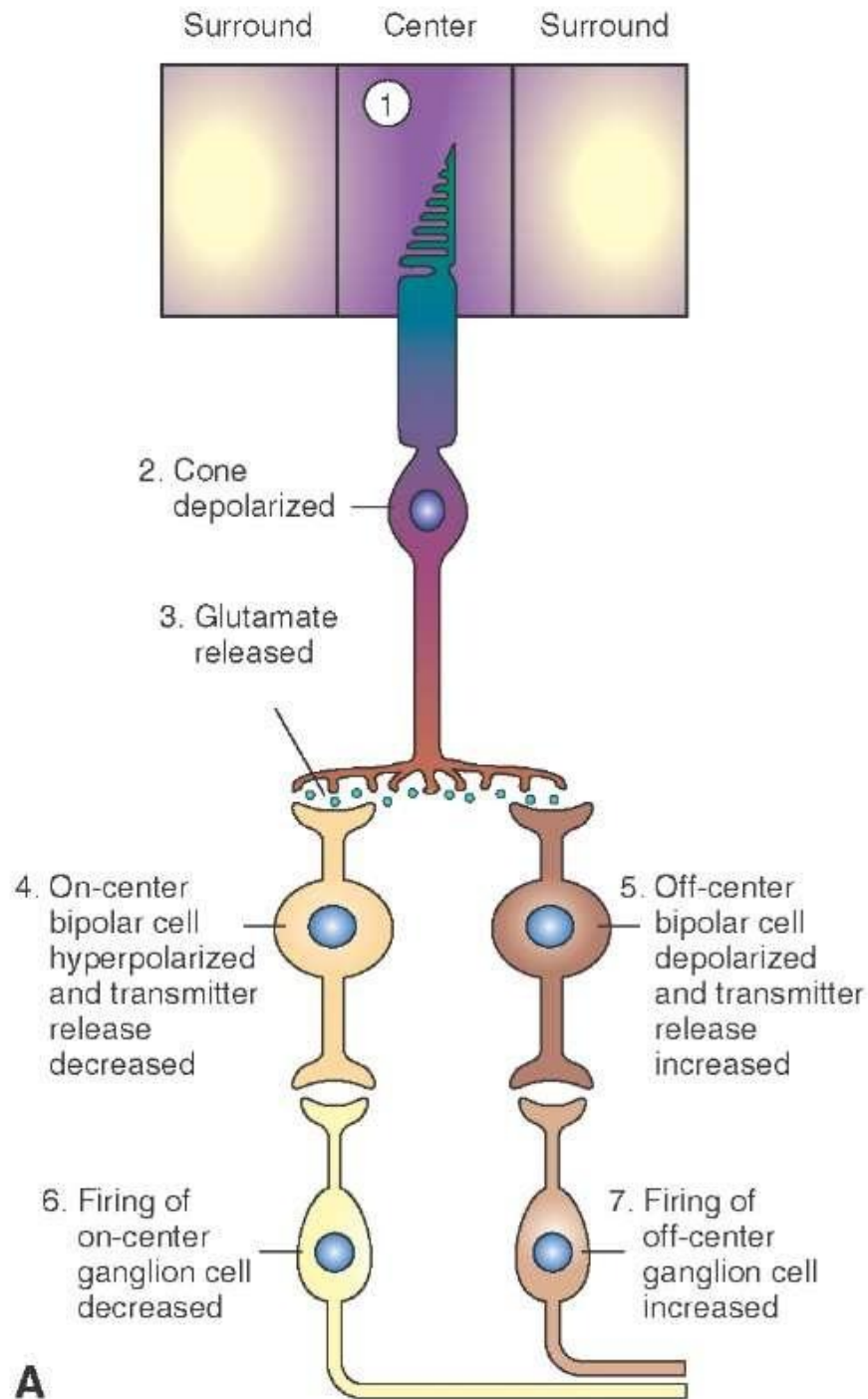


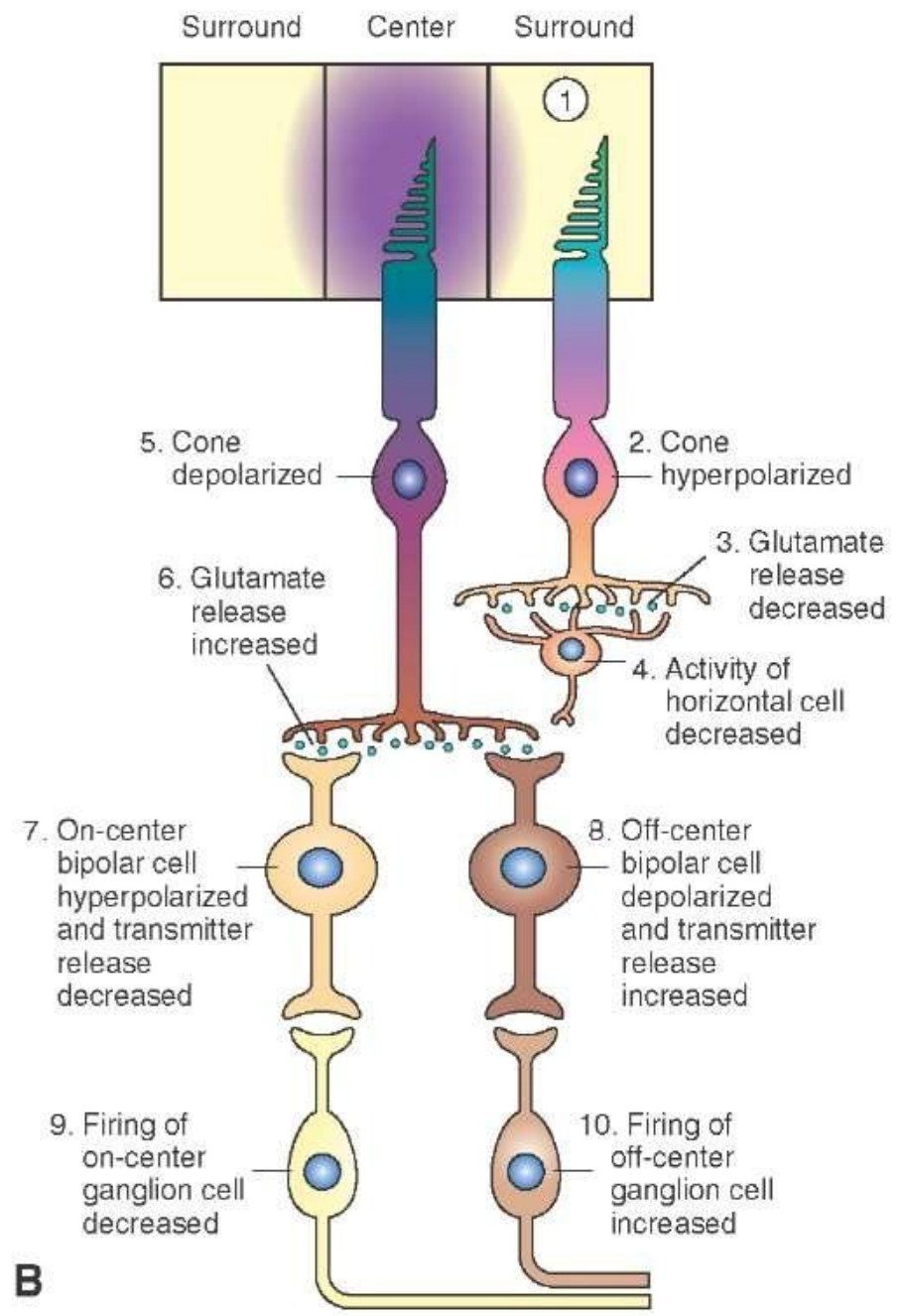
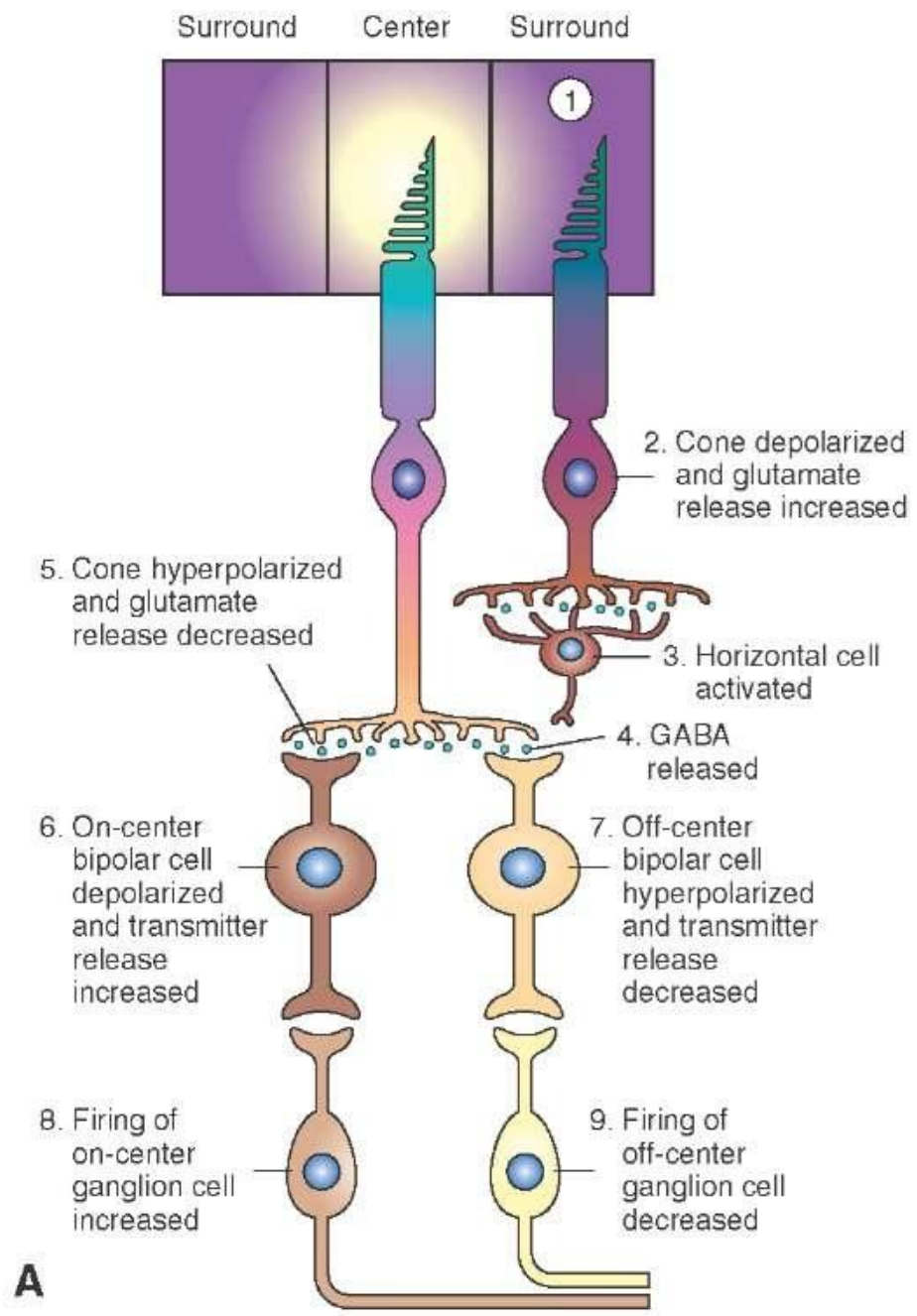


the bipolar cell is the most inhibited because receptor is depol & he is not inhibiting

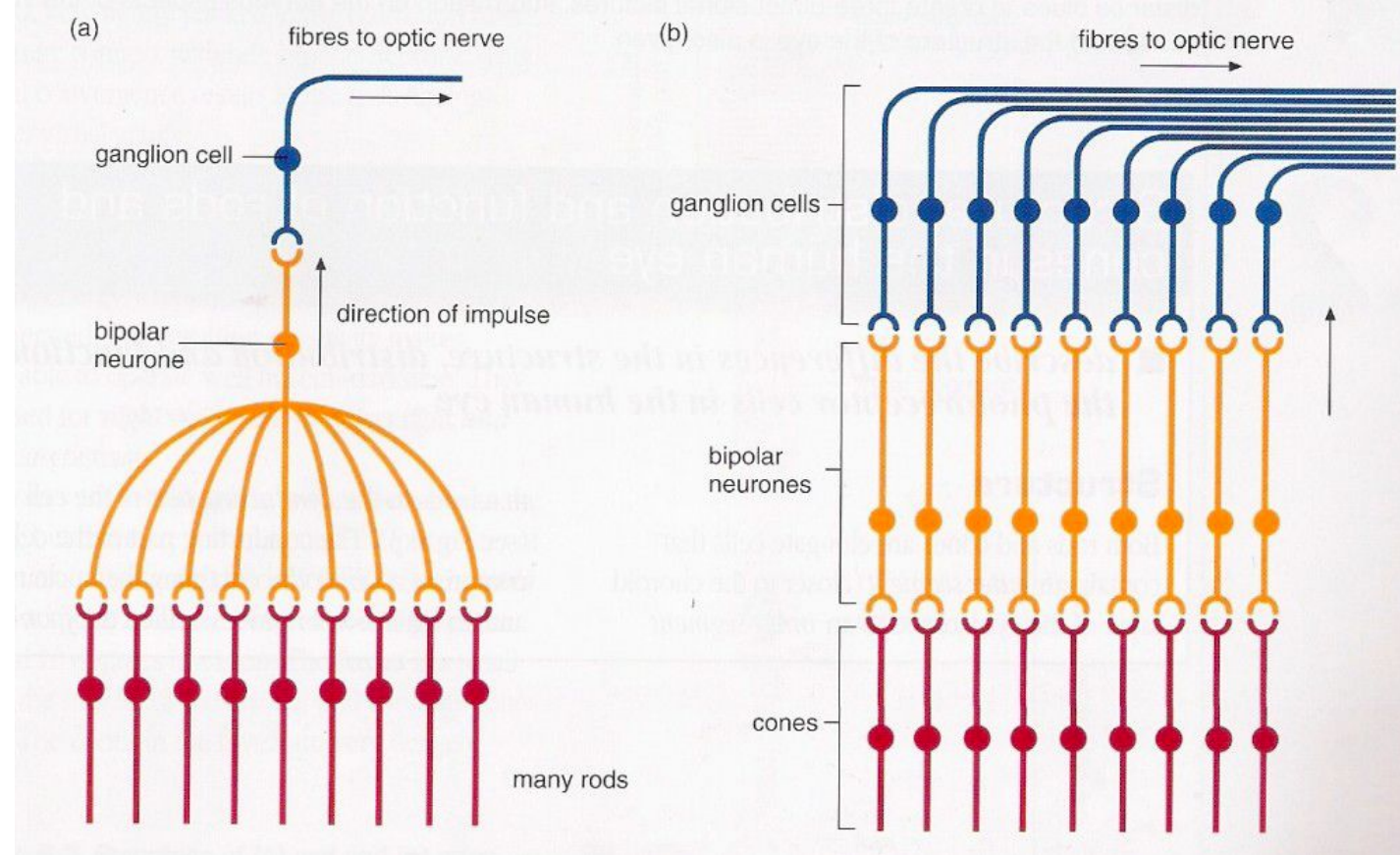


* surround receptors \rightarrow hyperpol \rightarrow horizontal cell minimally excited
 \therefore inhibitory signals are low

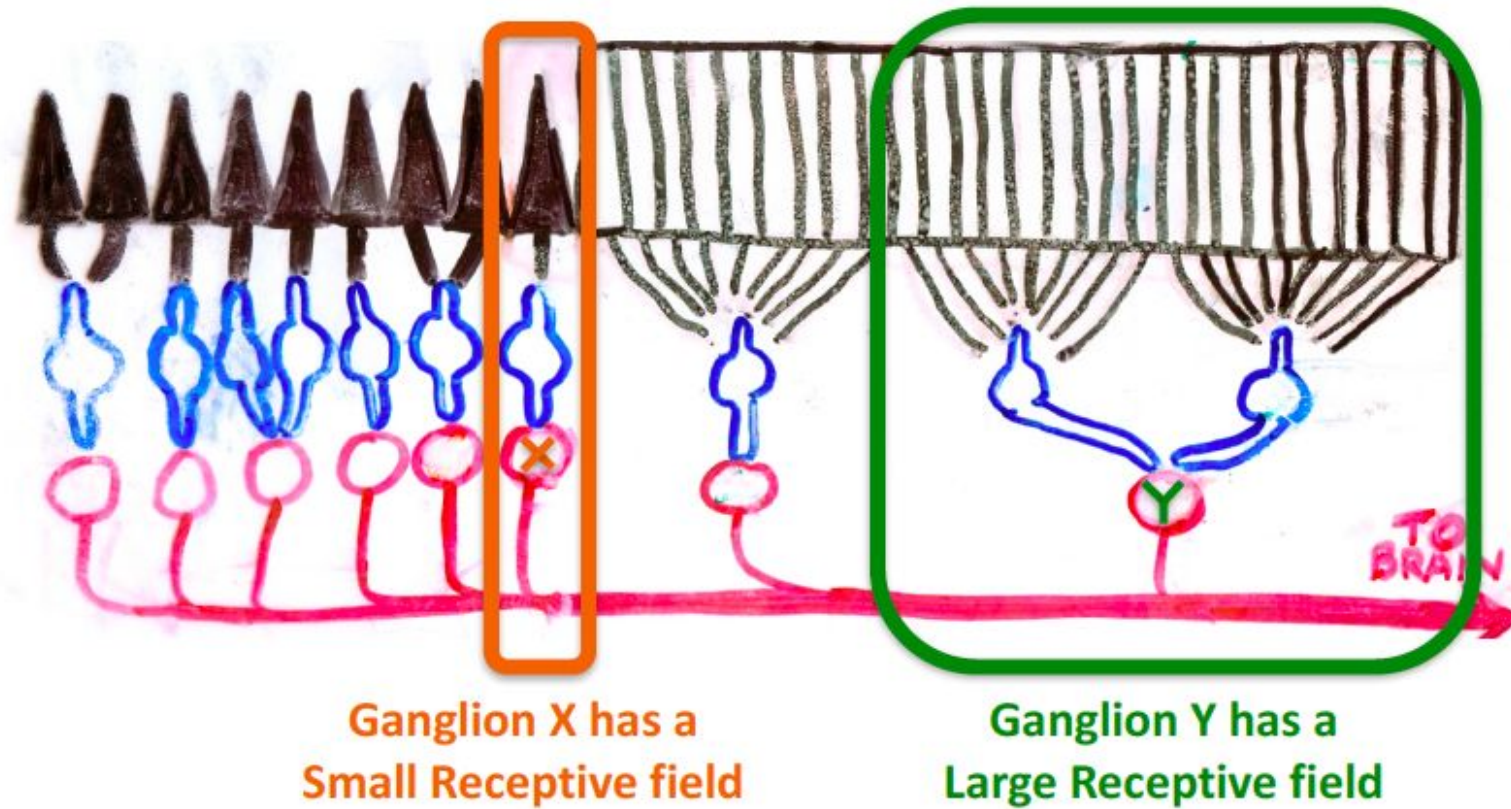




- Cones
 - *Low convergence*
 - *1:1 or few : 1*
 - *High acuity*
 - *Information about details gets preserved*
- Rods
 - *High convergence*
 - *Many : 1*
 - *Poor acuity*
 - *Details can be lost*
- In dim light (cones)
 - *Each cone reacts slightly*
 - *Not enough activity for each bipolar cell to cross the threshold for ganglion cell to fire*
 - *Still too much inhibition coming from the rods*
- In dim light (rods)
 - *Little bit of excitation from each bipolar cell*
 - *Ganglion cell will have enough input to cross threshold and start to fire*



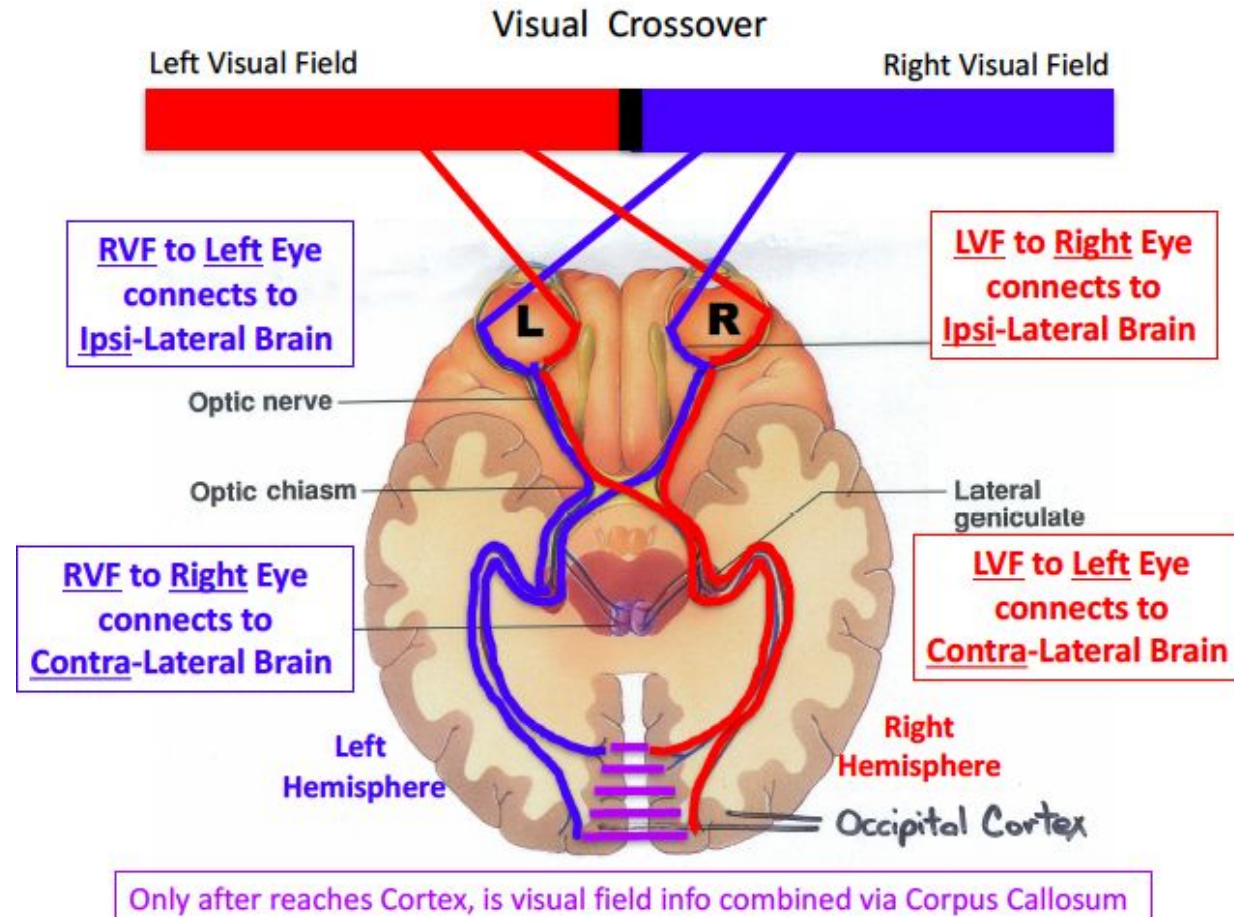
convergence



- Receptive field: set of receptors whose activity influences the activity of a “target” cell
- Better acuity when a cell has a smaller receptive field

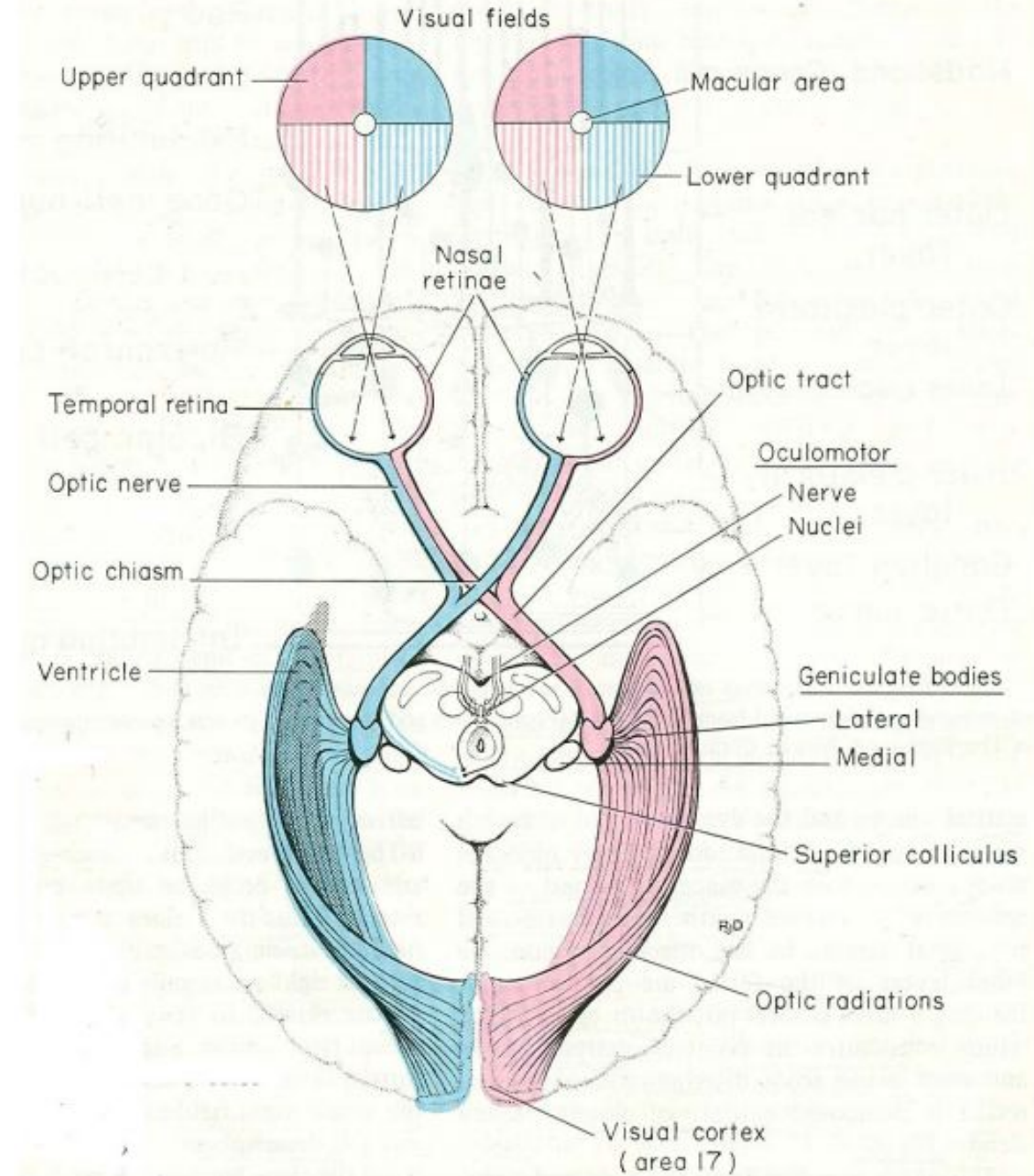
visual crossover

- Information from the right visual field goes to the left side of the eye
- Information from the left visual field goes to the right side of the eye
- Right visual field information goes to left side of the brain and vice versa
- Each optic nerve divides and goes to both sides of the brain

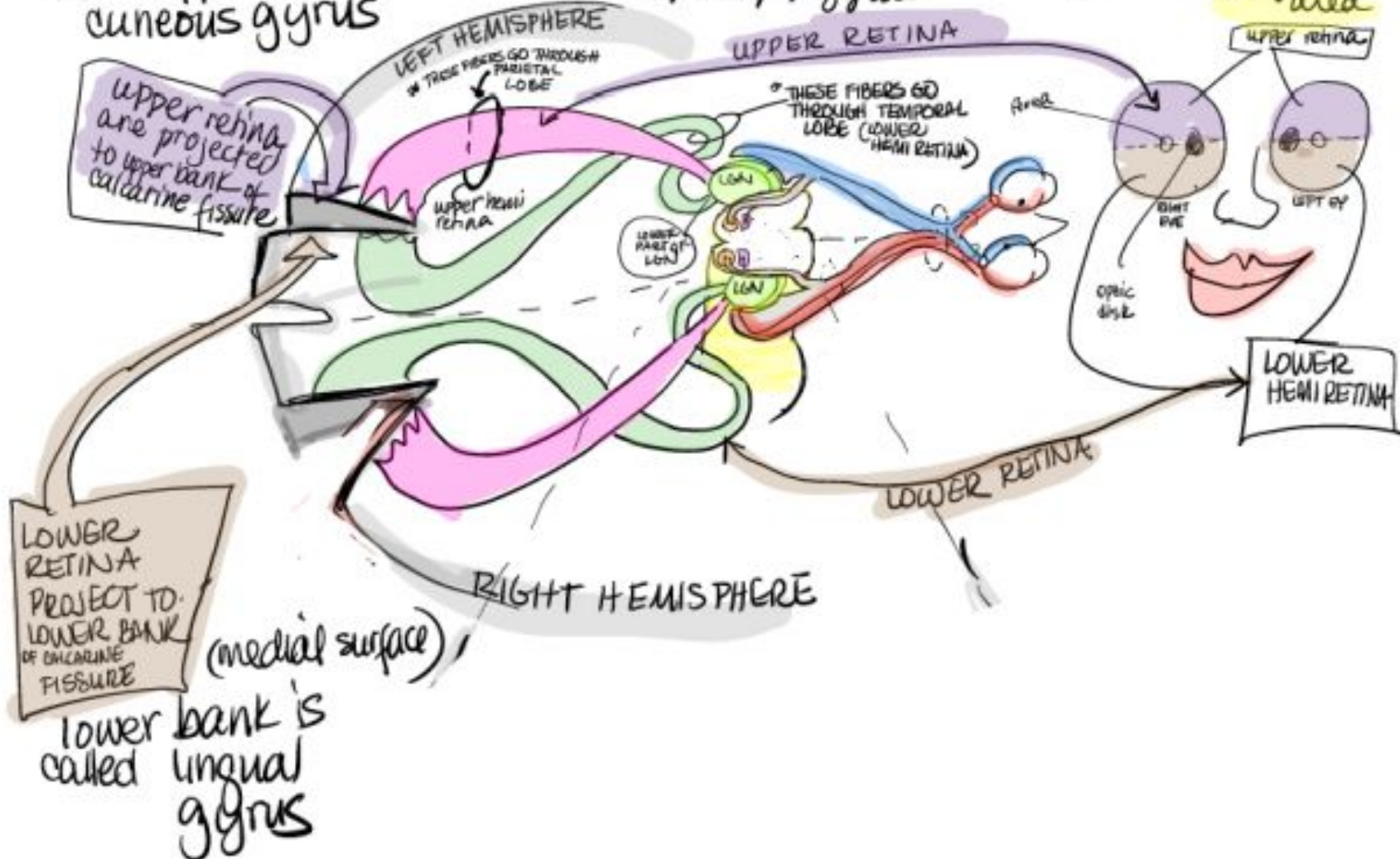
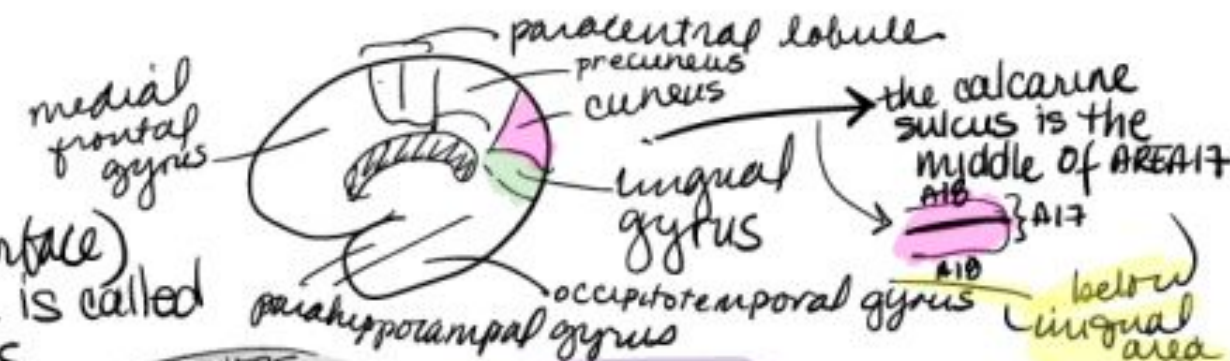


visual crossover

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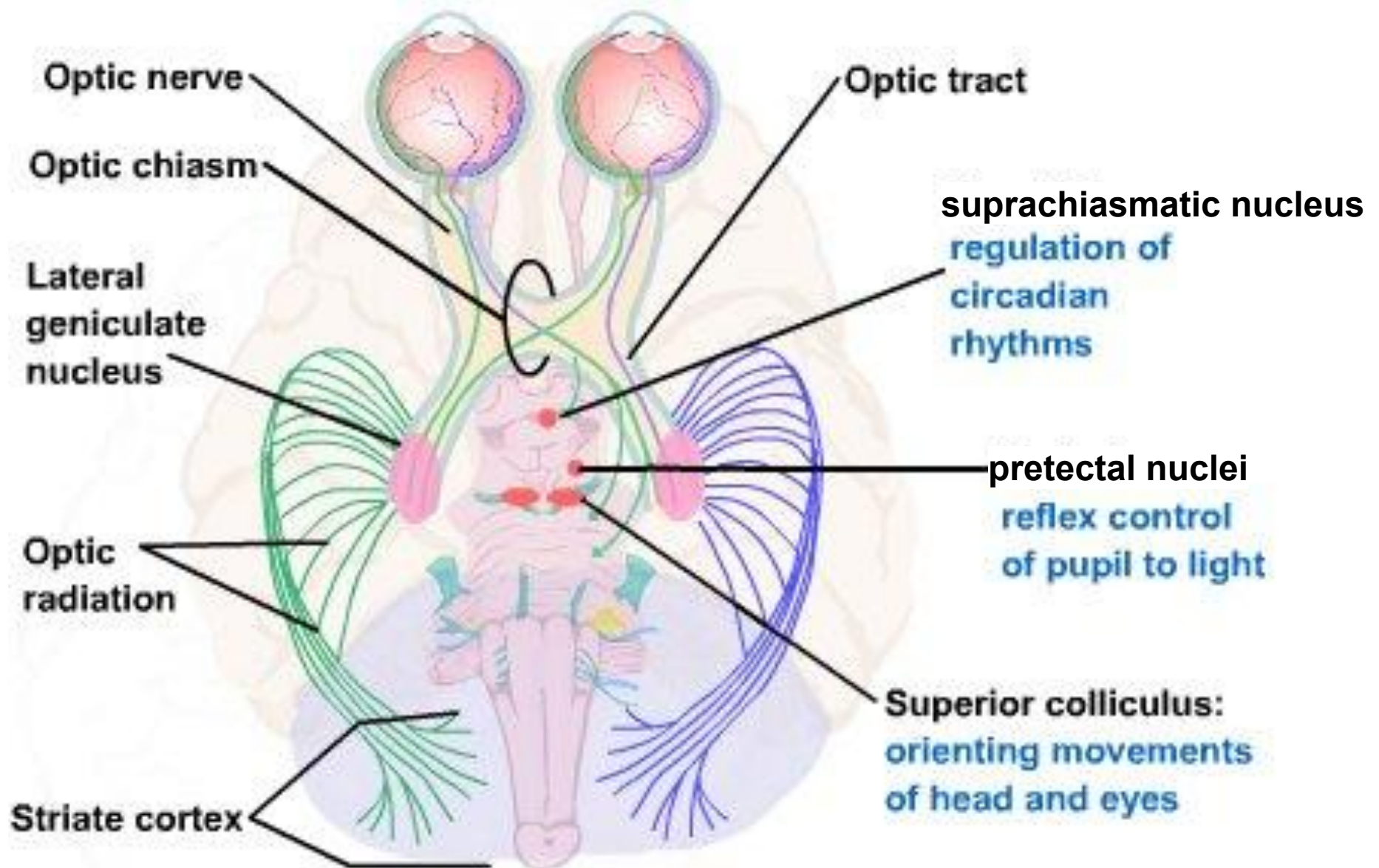
(medial surface)
note: upper bank is called
cuneus gyrus

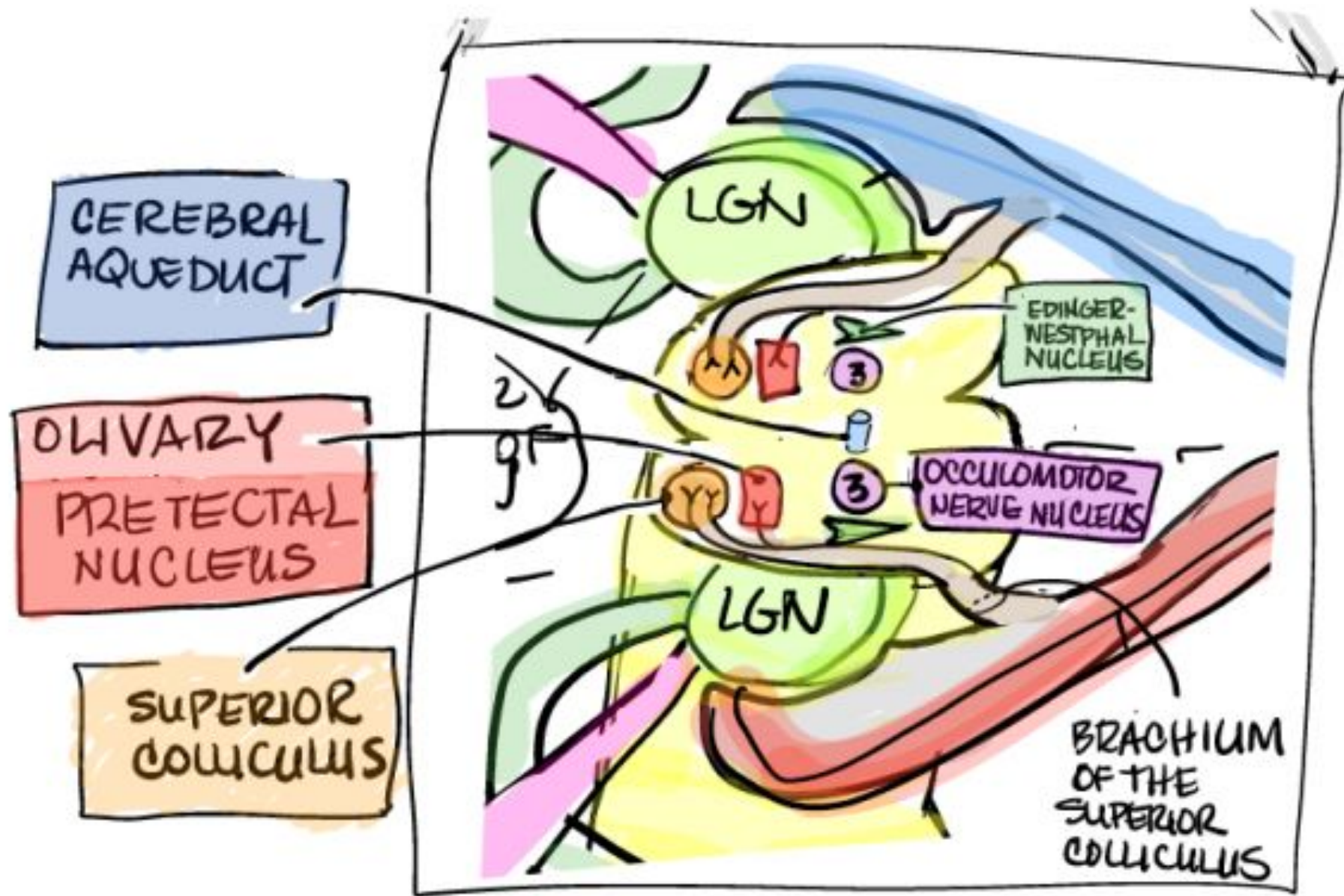


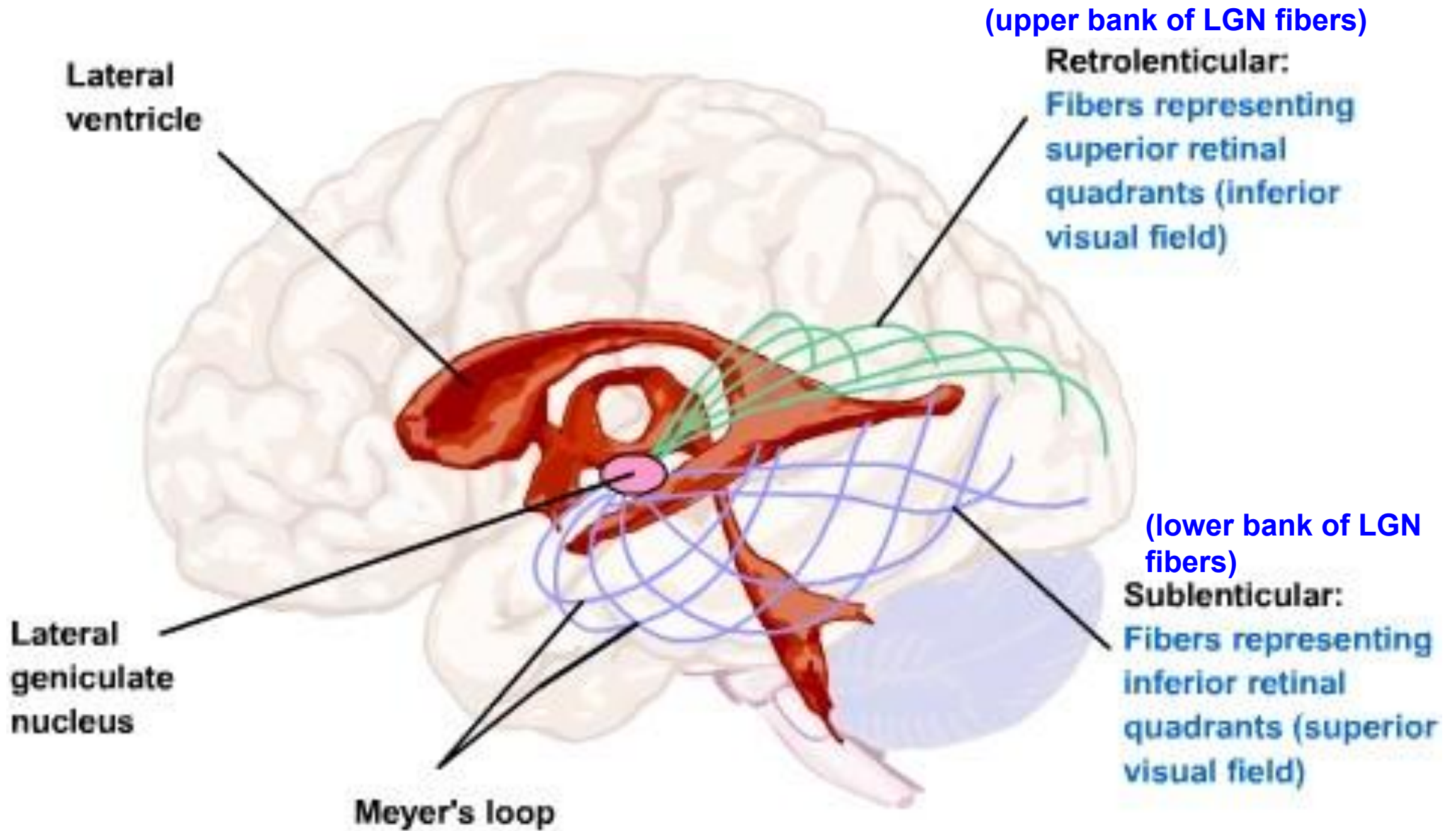
upper retina
are projected
to upper bank of
calcarine fissure

LOWER
RETINA
PROJECT TO
LOWER BANK
OF CALCARINE
FISSURE

(medial surface)
lower bank is
called lingual
gyrus

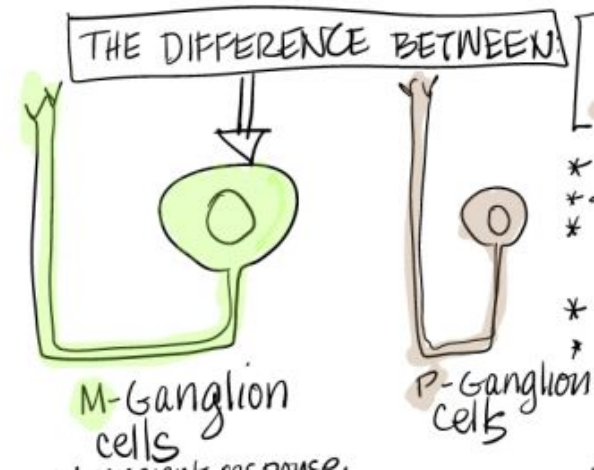
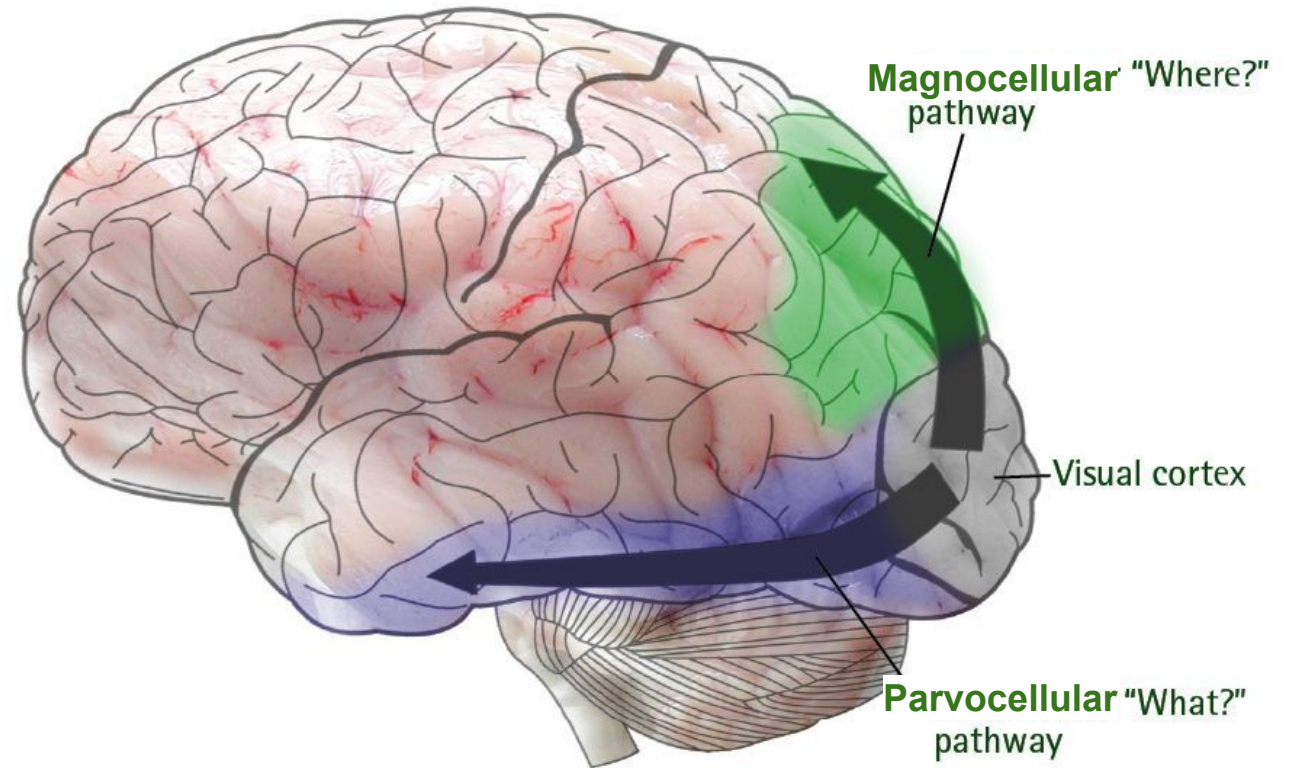






visual pathways

- Dorsal Pathway
 - “Where/how”
 - Motion and depth
 - Magnocellular pathway
 - Primarily rods
 - ‘Large’ ganglion cells
- Ventral Pathway
 - “Who/what”
 - Color and detail
 - Parvocellular pathway
 - Primarily cones
 - ‘Small’ ganglion cells



visual pathways

- Optic nerve goes to lateral geniculate nucleus (LGN) of the thalamus
- Optic chiasm: location where the axons cross (don't synapse)
- LGN
 - 6 layers
 - *Magnocellular pathways project to layers 1 and 2*
 - *Parvocellular pathways projects to layers 3-6*
- Some axons from magnocellular pathway project to the super colliculus

