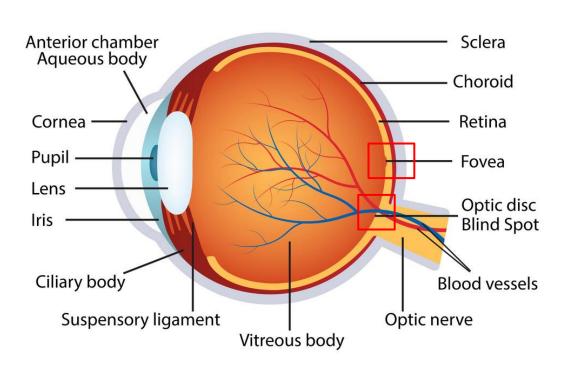
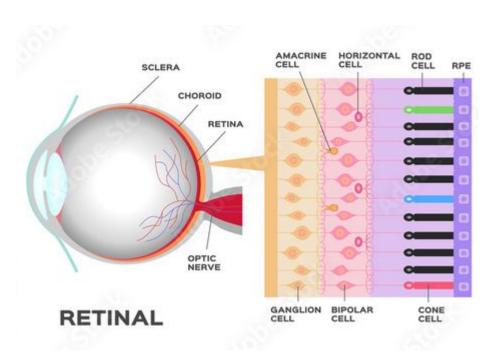
# COGS 17 WEEK 4 WINTER 2024, A04

#### THE EYE



- Fovea -- Small central area of high concentration of Cones only, for HIGH DETAILS
- Retina -- Senses light, send information to the brain through >>
- Optic nerve
- Blind Spot -- No Receptors here

#### THE RETINA



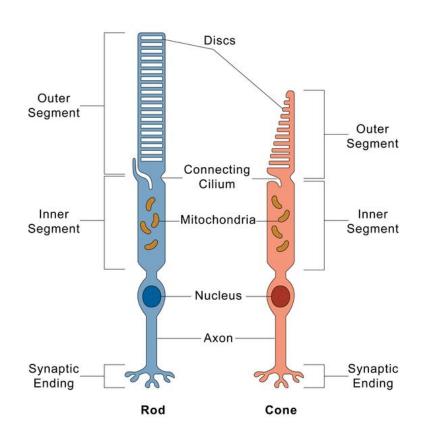
- •Receptors -- Rods & Cones
- Bipolar cells -- Postsynaptic to Receptors
- •Ganglions -- Axons of the Ganglion Cells form the Optic Nerve
- •Interneurons -- perpendicular to pathway, influence above neurons
- Pigment Epithelium -- Non-neural cells, feed & recycle nutritions from receptors;

helps reflect/maximize light

#### ISOMERIZATION

- •Converting light into a neural signal
- Photopigment -- Made of Opsin & Retinal
- •11-Cis Retinal absorbs photon of light, changes shape >> detaches from Opsin >>
- Activates second messengers in receptor >> ion gates closing >> modifying GRADED release of NT
- Photopigment regeneration -- using Enzymes from Pigment Epithelium, requires time

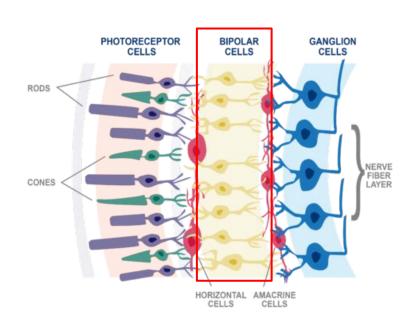
#### VISUAL RECEPTORS



- •Rods -- being larger, have MORE
  photopigment but only 1 kind >>
  DO NOT code color; high
  sensitivity; poor acuity;
  excellent for motion detection
- •Cones -- smaller, have 3 kinds of
  photopigments >> DO code color;
  low sensitivity; excellent
  acuity; poor for motion detection
- Receptors show Spontaneous firing,
   Graded notentials release

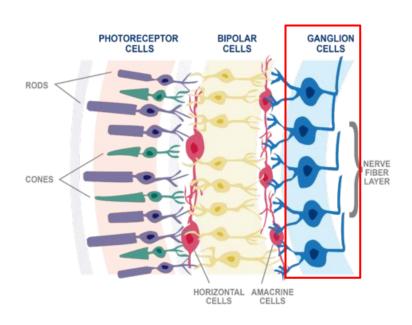
Graded potentials, release
Inhibitory NT

## BIPOLAR CELLS



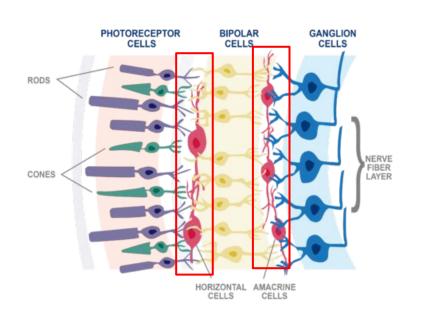
Postsynaptic to Receptors, showSpontaneous firing, GradedPotentials, release Excitatory NT

# GANGLION CELLS



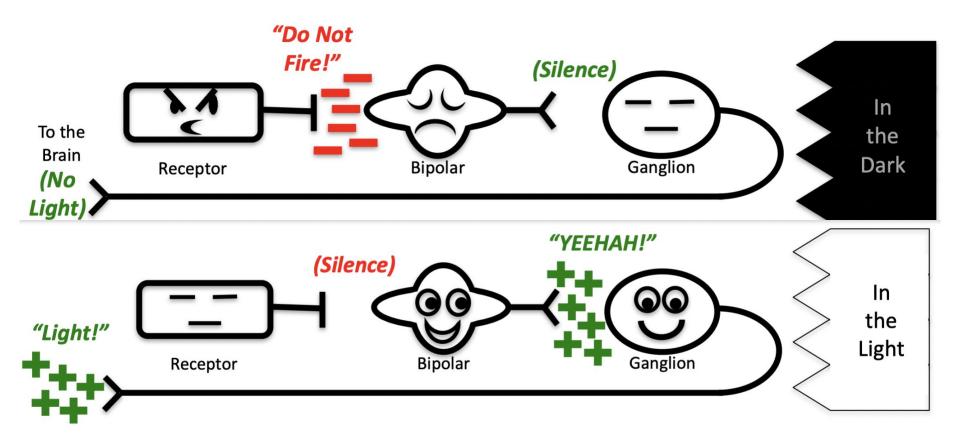
Postsynaptic to Bipolars, showAction Potentials, releaseExcitatory NT

#### INTERNEURONS



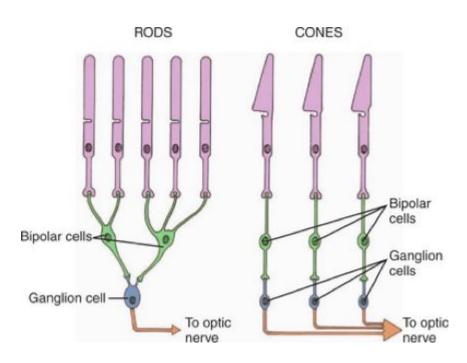
- Horizontal Cells -- Graded
   Potentials, mostly Inhibitory
   NT, modify interface of
   Receptors and Bipolars
- Amacrine Cells -- Graded
   Potentials, mostly Inhibitory
   NT, modify interface of Bipolars and Ganglions

# RECEPTORS ARE TURNED OFF BY LIGHT



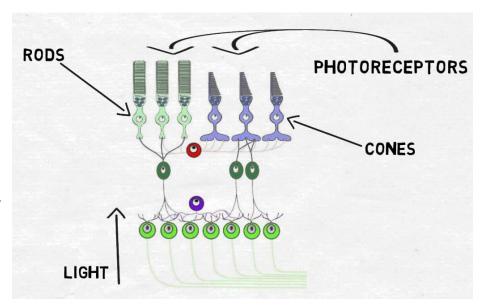
# CONNECTIVITY PATTERNS

 Play a critical role in information transmission functions



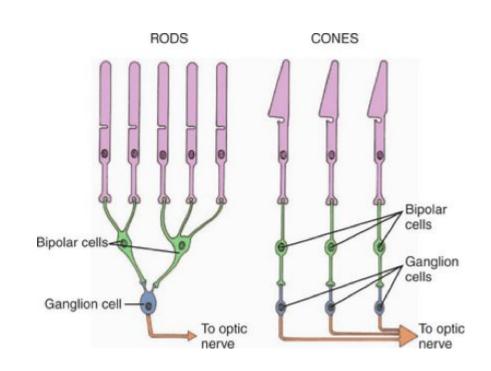
#### CONVERGENCE

- •Receptors converge (via their Bipolars) onto Ganglion cells
- Rods -- High Convergence, avg.120:1 Ganglion
- ●Cones -- **Low** Convergence, avg. 6:1 Ganglion
- ●In Fovea: **Very Low**, often only 1:1 Ganglion

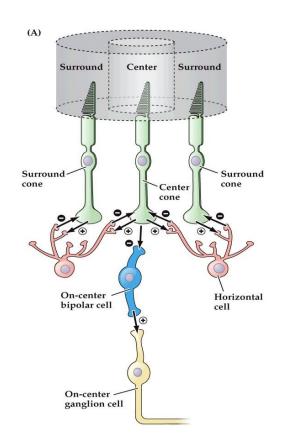


#### CONVERGENCE

- Helps to account for acuity and sensitivity differences between rods & cones
- Also, Rods are LARGER and have more Photopigment than Cones do, and this also contributes to sensitivity

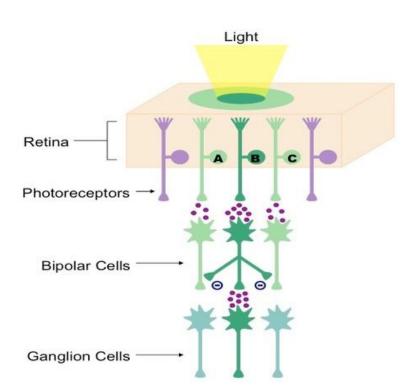


#### RECEPTIVE FIELDS



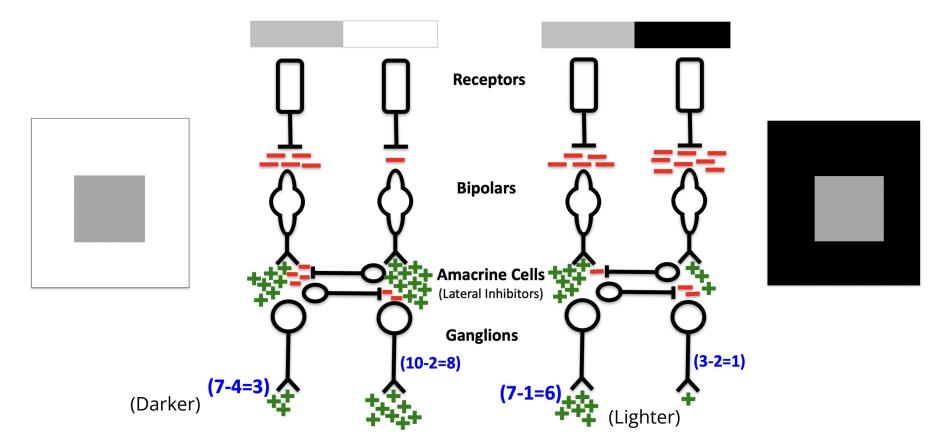
- •Set of Receptors whose activity influences the activity of a "Target" cell
- •Size and type of a Target's RF is determined
  by patterns of Convergence and Lateral
  influences
- •Example 1: Ganglion along path from converging Rods has large RF, while Ganglion along path from Cones has small RF
- •Example 2: Some Ganglions, LGN, and V1 cells have Center-Surround RFs

#### LATERAL INHIBITION

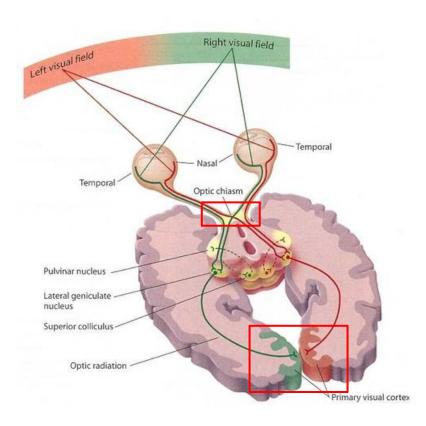


- •A neuron's response to a stimulus is **inhibited** by the excitation of a **neighboring** neuron
- •Mainly to exaggerate differences
- •Example: simultaneous contrast

# SIMULTANEOUS CONTRAST

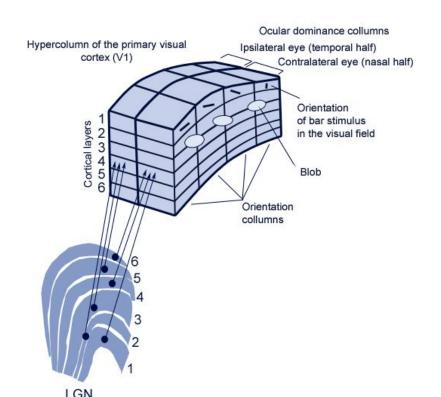


## VISUAL CROSSOVER



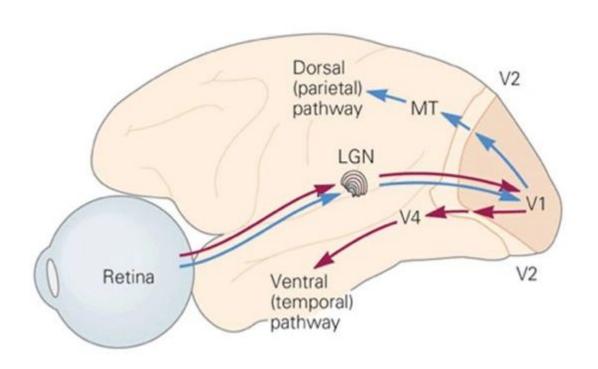
- Each Optic Nerve, from each eye, divides and goes to both sides of the brain
- •Info from Left Visual Field =>
  Retina on right side of RIGHT
  eye => right LGN => right Visual
  Cortex
- •Info from Right Visual Field =>
  Retina on left side of RIGHT eye
  => crossover at the Optic Chiasm
  => right LGN => right Visual
  Cortex
- Info from cortex exchange across corpus callosum

#### VISUAL CORTEX



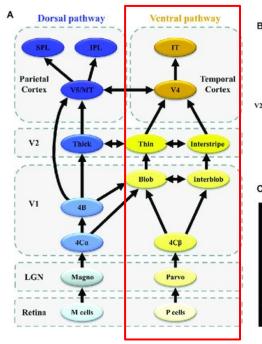
- •Cells in all 6 layers that respond to same "preferred" stimulus
- •E.g. lines of a particular Orientation
- ●Hypercolumn -- One set of orientation column w/same Receptive Field
- •All cells within a given Hypercolumn have same Receptive Field
- •One hypercolumn includes columns set of full orientations, plus Blobs for color processing

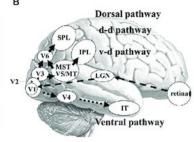
#### VISUAL PATHWAYS

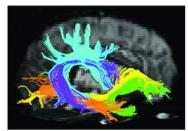


- ParvocellularPathway (Who/WhatPathway, VentralPathway) -- Foridentifying stimuli
- Magnocellular
   Pathway
   (Where/How Pathway,
   Dorsal Pathway) For visual-spatial
   mapping

#### PARVOCELLULAR PATHWAY

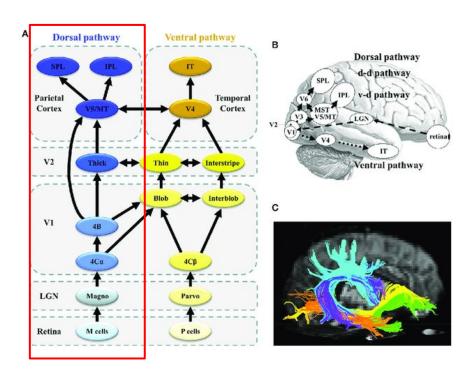






- Specialized for color & detail
- Begins at Cones in and nearFovea
- Mostly Parvocellular (small)Ganglions, with small RFs
- •Basic pathway: Retina >> Top 4
  Layers of LGN in Thalamus >>
  V1 >> V2 >> V3 >> V4 (all in
  occipital
  - lobe) >>Inferior Temporal Cortex

#### MAGNOCELLULAR PATHWAY



- Specialized for detecting motion, locating objects, navigating & manipulating environment including gross outline
- Begins at Rods & Cones in periphery of Retina
- Basic pathway: Some of info to Superior Colliculus of Midbrain (e.g. for "Blindsight"), then to LGN;

Most go directly to LGN >> All info >> V1 >> V2 >> Medial Temporal Cortex >> Medial Superior Temporal Cortex >> Posterior Parietal Cortex

# QUESTIONS?

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Office Hours: Mon 5-6 pm
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To get the section slides:
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https://github.com/JasonC1217/COGS17\_A04\_Wi24

OR:

