

# Color Vision

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**Nicole Weisschuh**  
**Molecular Genetics Laboratory**



**Lecture "Sensory Systems – Basics and Principles II**  
**05.07.2010**

# Lecture aims: To understand the following

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## SHOCKING CONFESSION

Daniel B. (24): „I am colorblind!“

**Young & Helmholtz state: „Color vision is trichromatic!“**

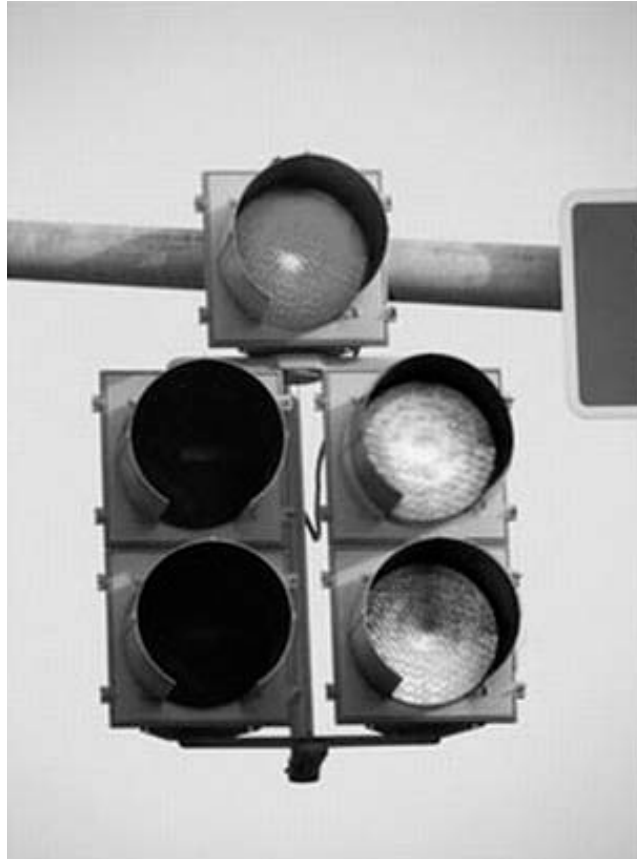
The principle of univariance

Cone mystery resolved

**Color opponency: Hering declares his  
4 „Urfarben“ to be responsible**

**How dependent are we on color vision?**

Imagine you are in a hurry...

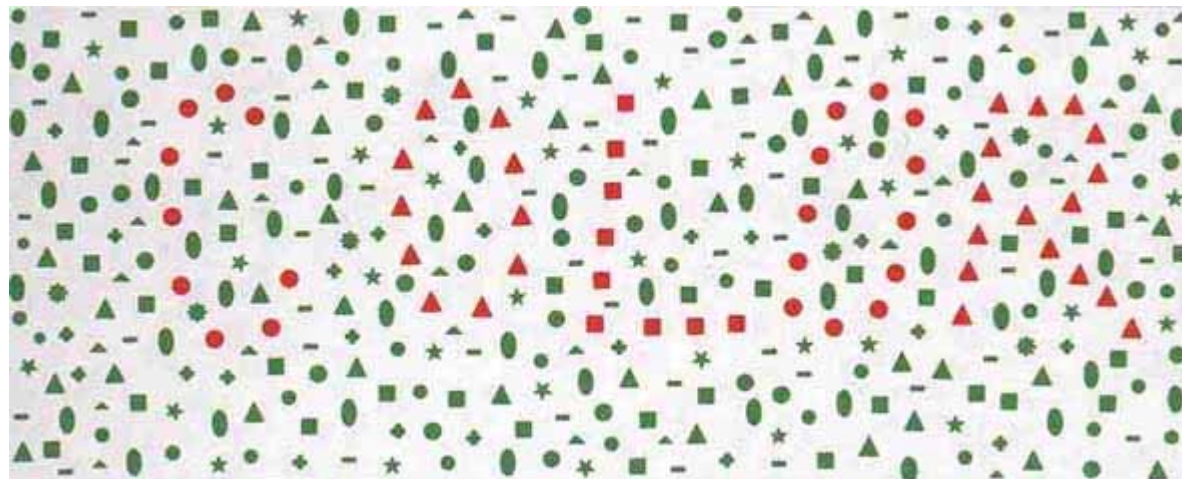
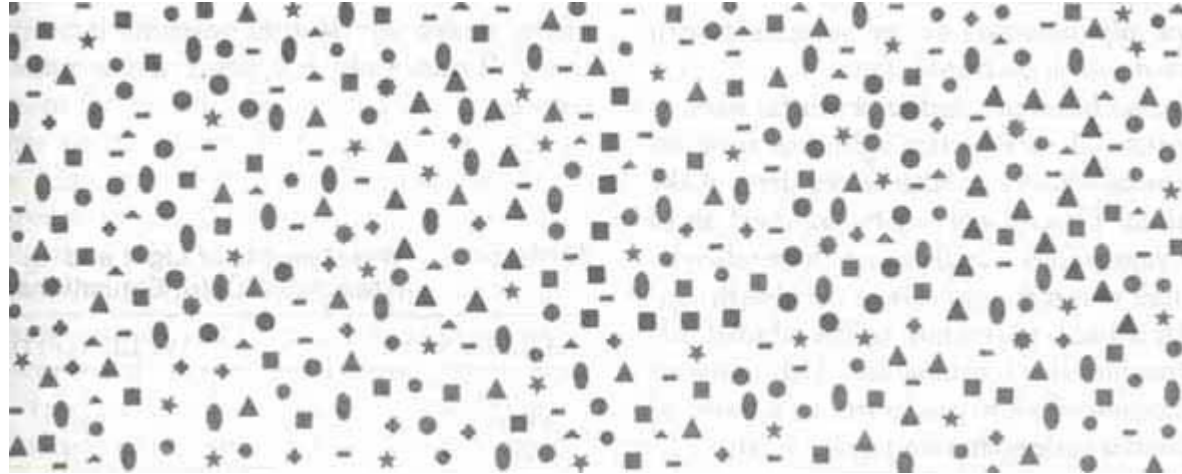


**... or you are hungry**

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# Color helps!



**Split the image into...**

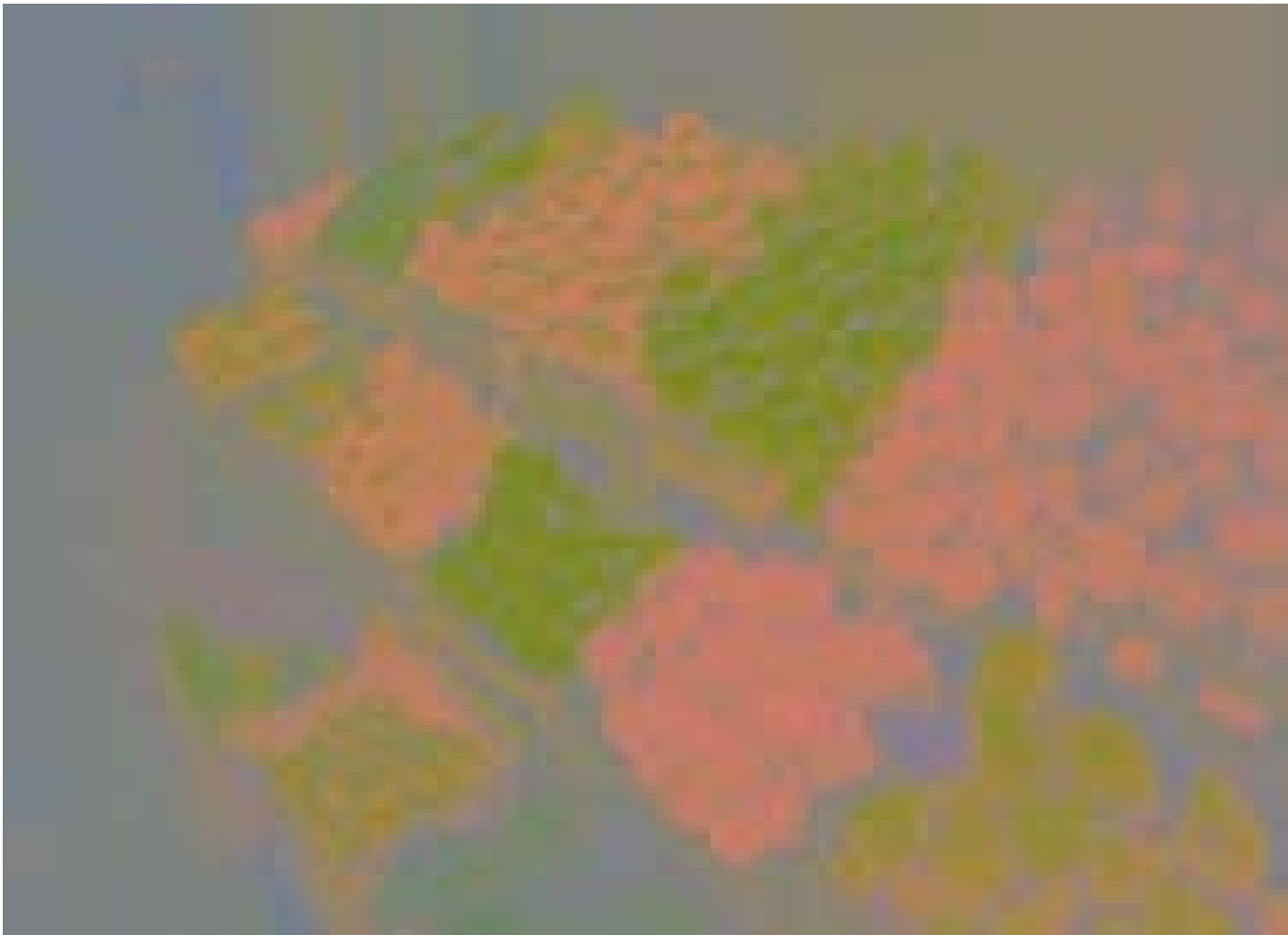


**Chromatic components**



**Achromatic components**





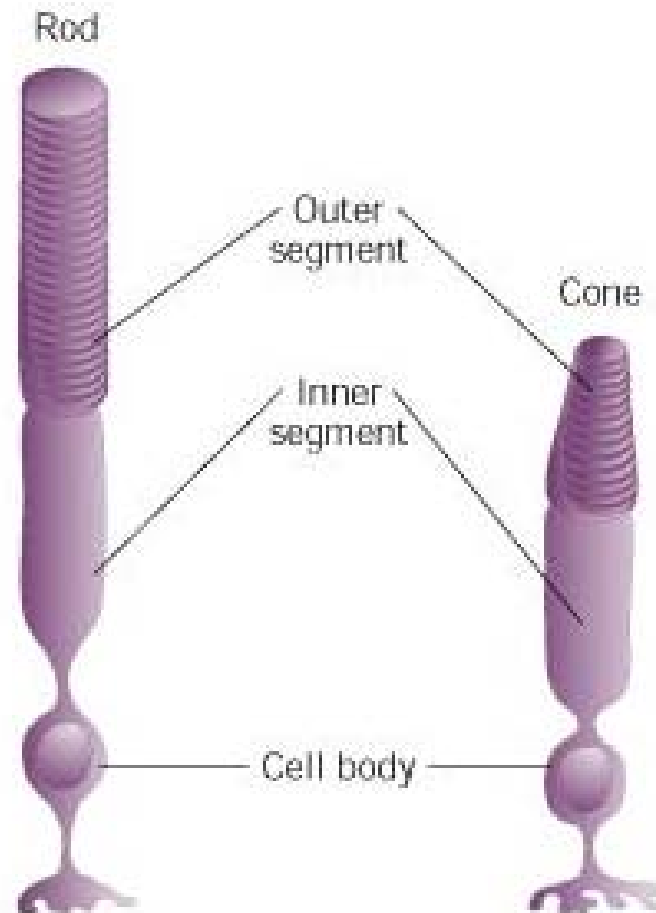




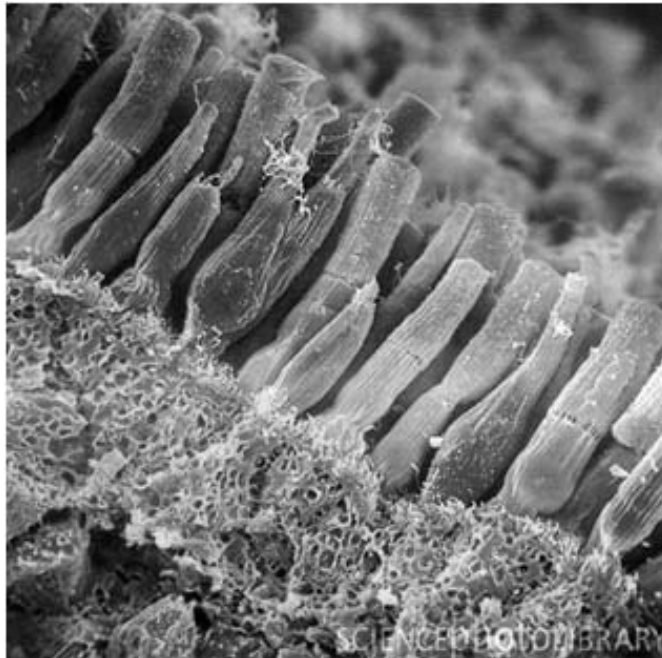
**How do we see colors?**

# Photoreceptors

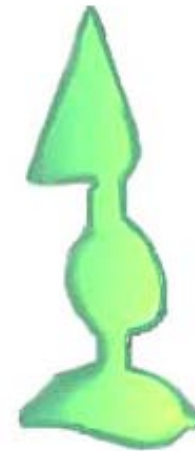
There are two types of photoreceptor cells in the human retina, **rods** and **cones**.



# Color vision is mediated by cones



**Blue cone**  
„S-cone“

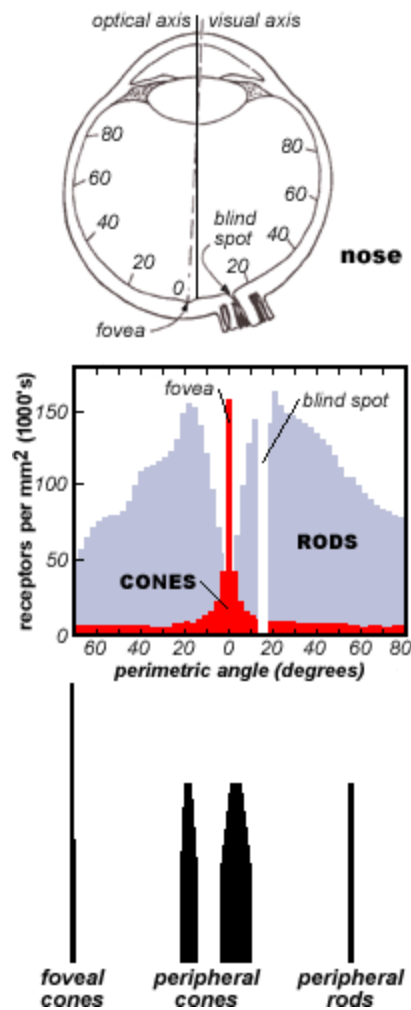


**Green cone**  
„M-cone“

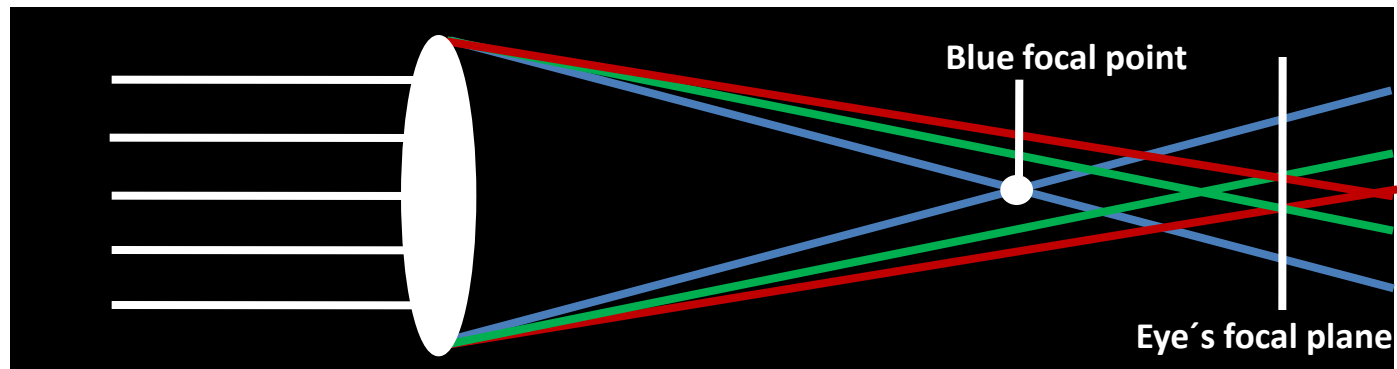
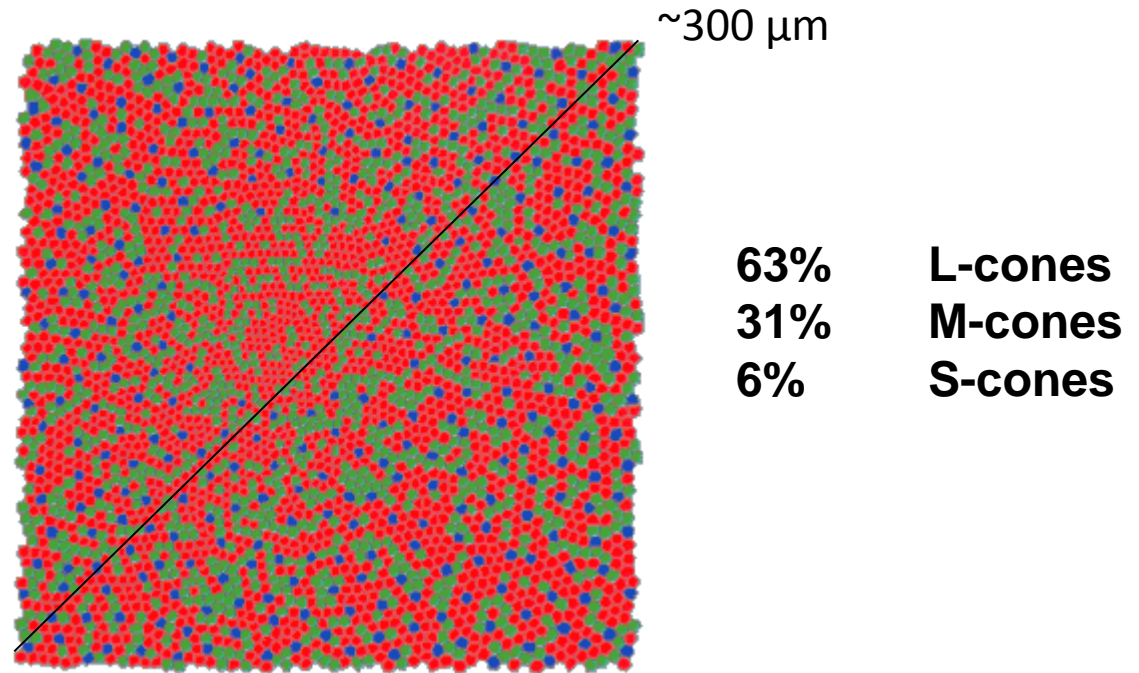


**Red cone**  
„L-cone“

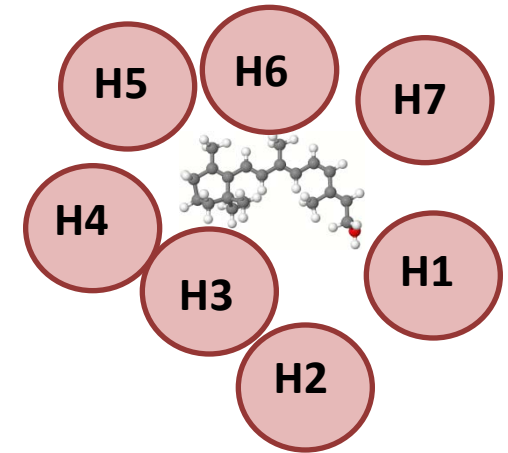
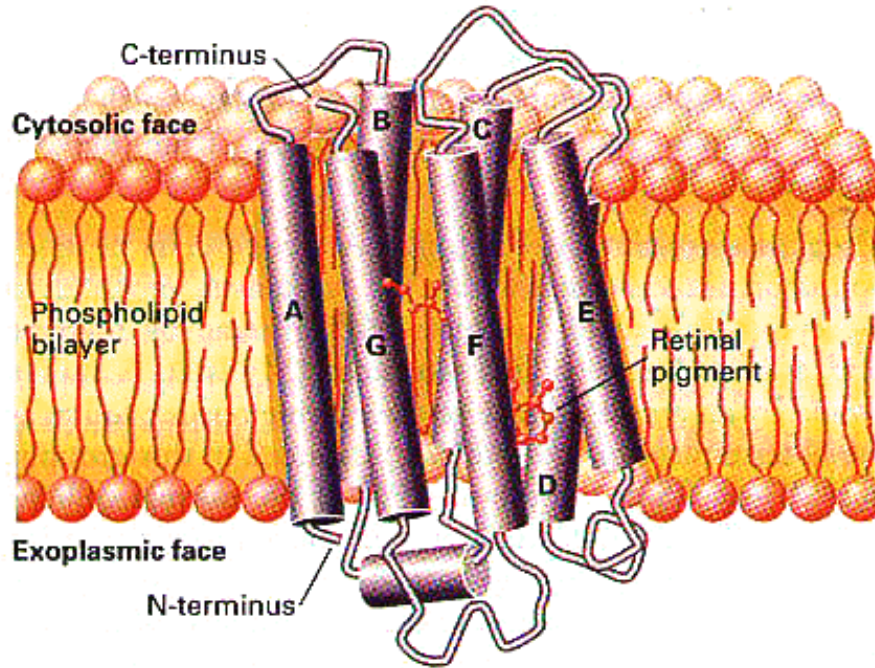
# Distribution and size of photoreceptors in the retina



# The cone mosaic of the rod-free inner fovea



# Opsin structure

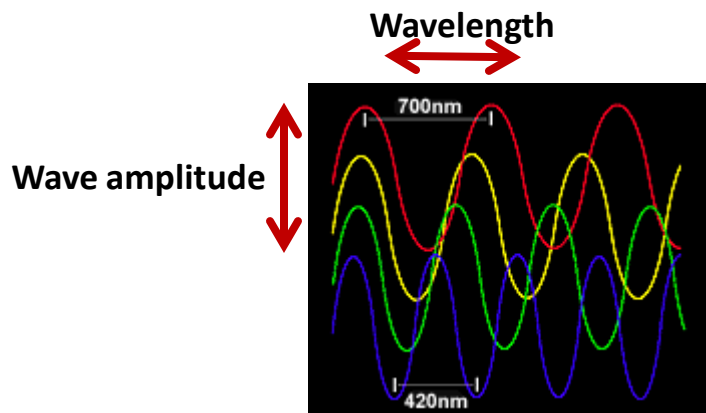


**Note:** Opsins have a  $\lambda_{\max}$  below 300 nm. Retinal has a  $\lambda_{\max}$  of  $\sim 380$  nm. The broad absorbance spectrum of 400-700 nm is created by the binding of both components. The  $\lambda_{\max}$  of the absorbance band depends on the genetically determined aa sequence of the respective opsin and the relationship of the opsin with the chromophore.

**Is colour, as we perceive it,  
mainly a property of physics or  
biology?**

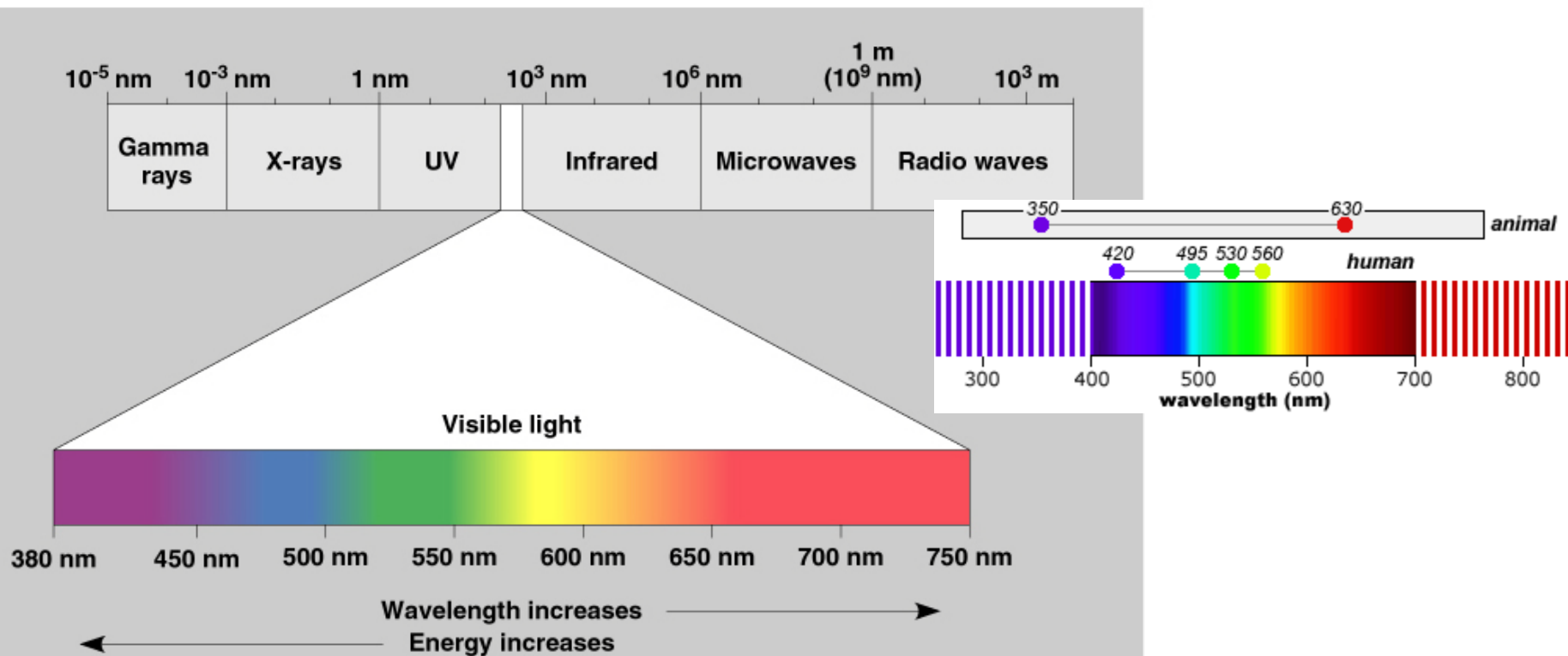


# Visible spectrum



Physical properties of light	Related perception
Wavelength	Color
Amplitude	Brightness

Visible light is a small part of the electromagnetic spectrum



# Who knows these gentlemen?



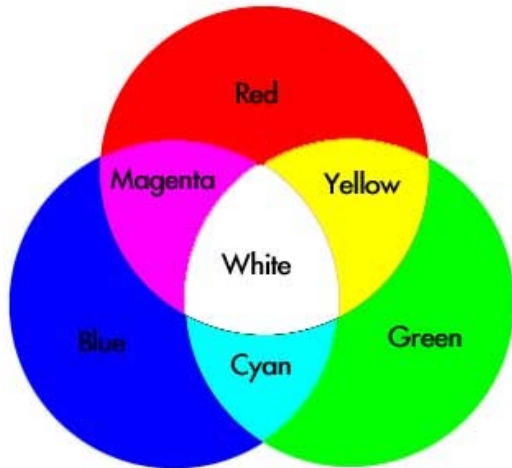
**Thomas Young 1773–1829**



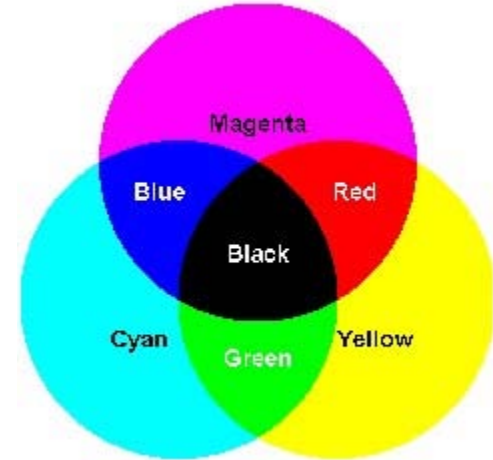
**Hermann Ludwig Ferdinand  
von Helmholtz 1821-1894**

# Human vision is trichromatic

Additive color mixing (RGB)



Subtractive color mixing (CMYK)



**Why is normal human vision trichromatic?**

# 1. Three types of cones

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**Blue cone**  
„S-cone“



**Green cone**  
„M-cone“



**Red cone**  
„L-cone“

## 2. Univariance

„I just absorbed 1 photon of 450 nm  
and 1 photon of 600 nm“

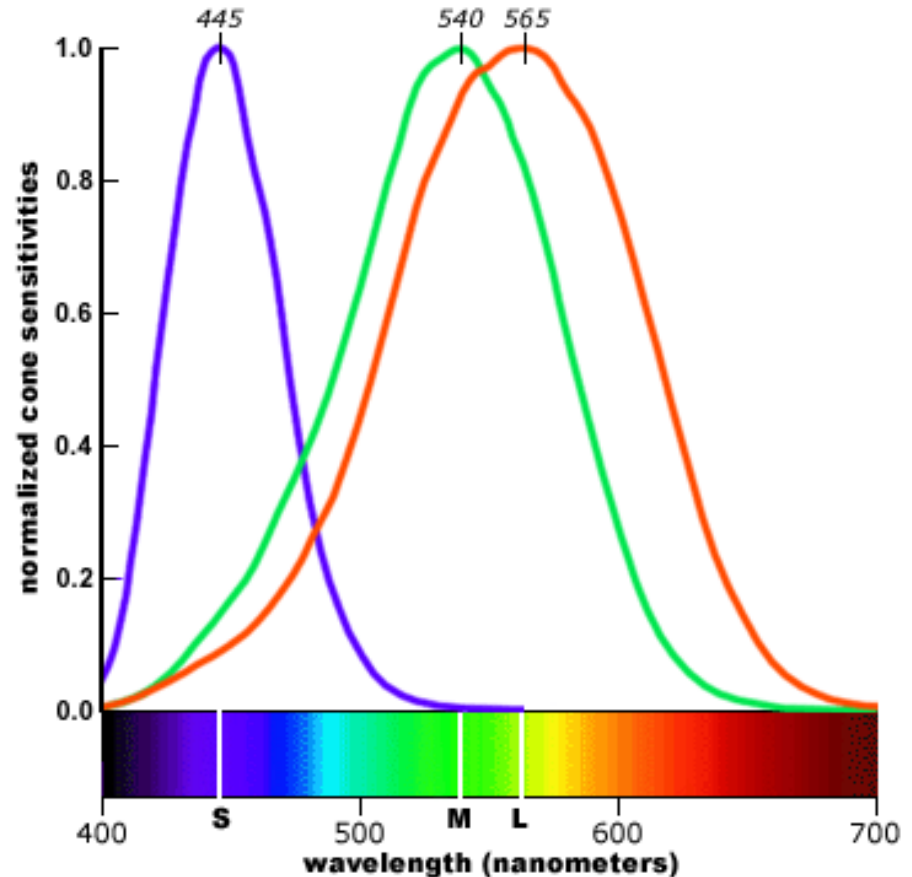
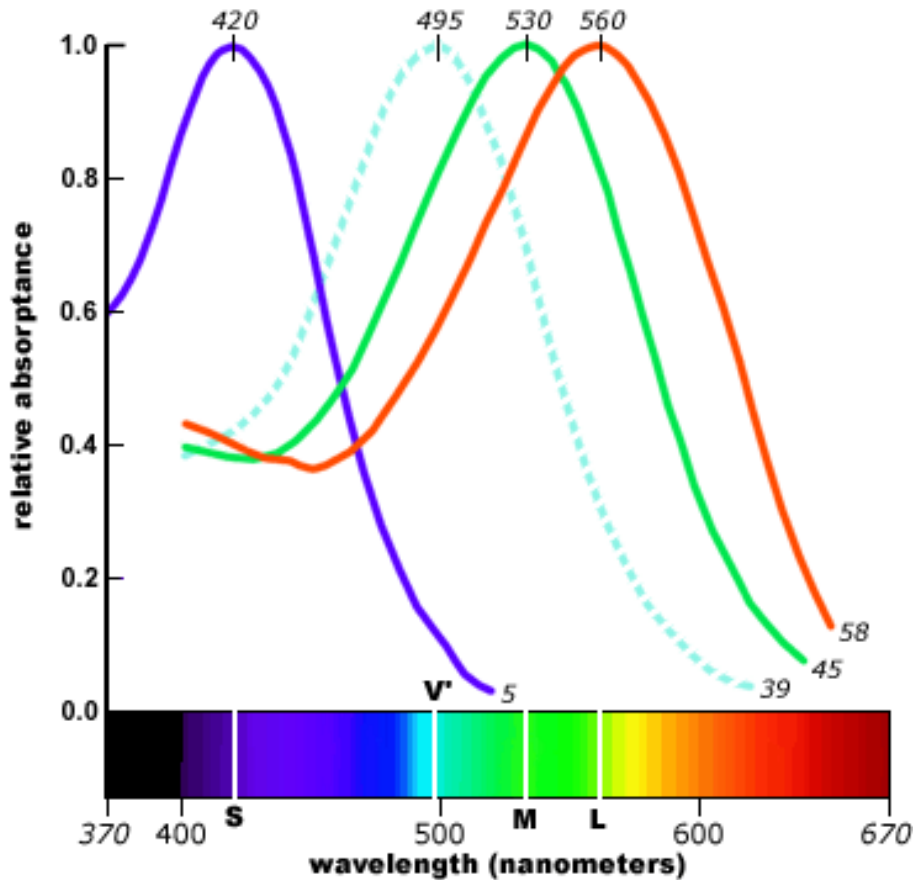


„I just absorbed 2 photons  
and I have no idea what  
their wavelengths are“



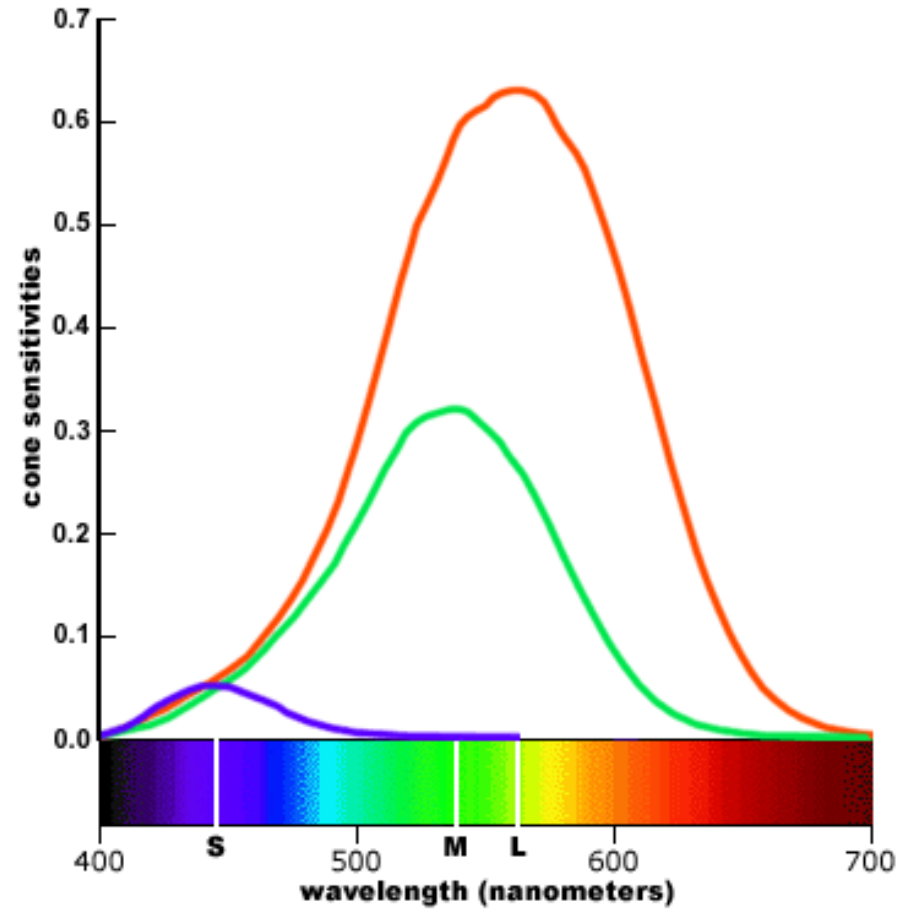
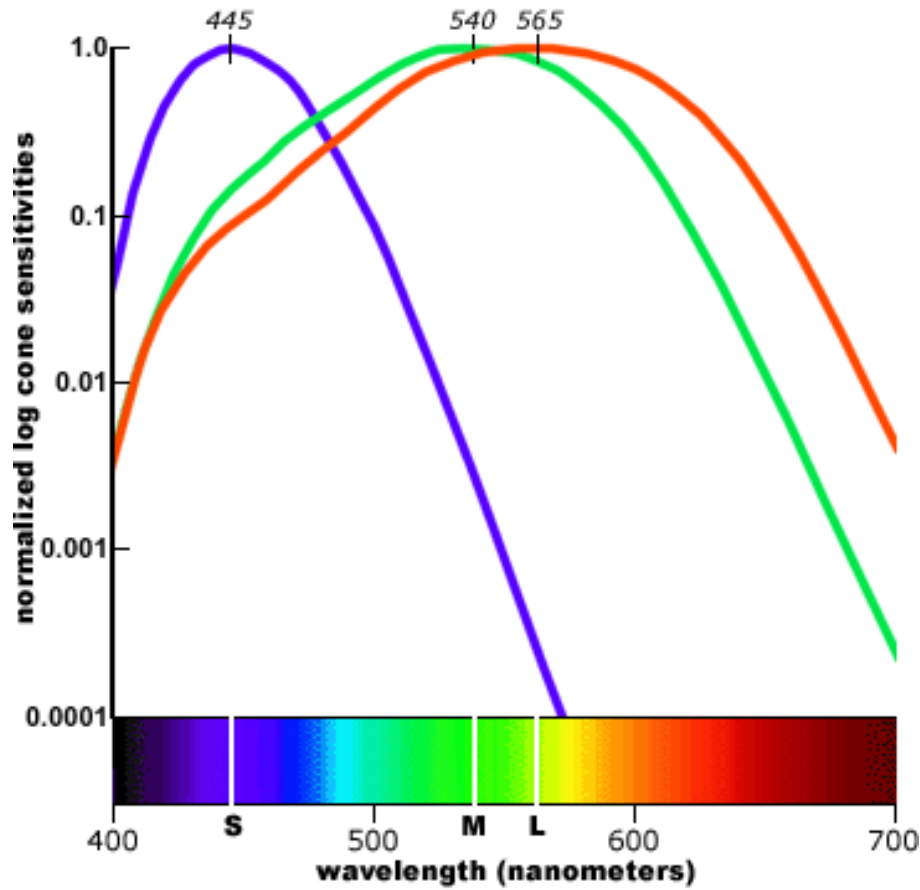
What do you think is true?

# Absorption spectra



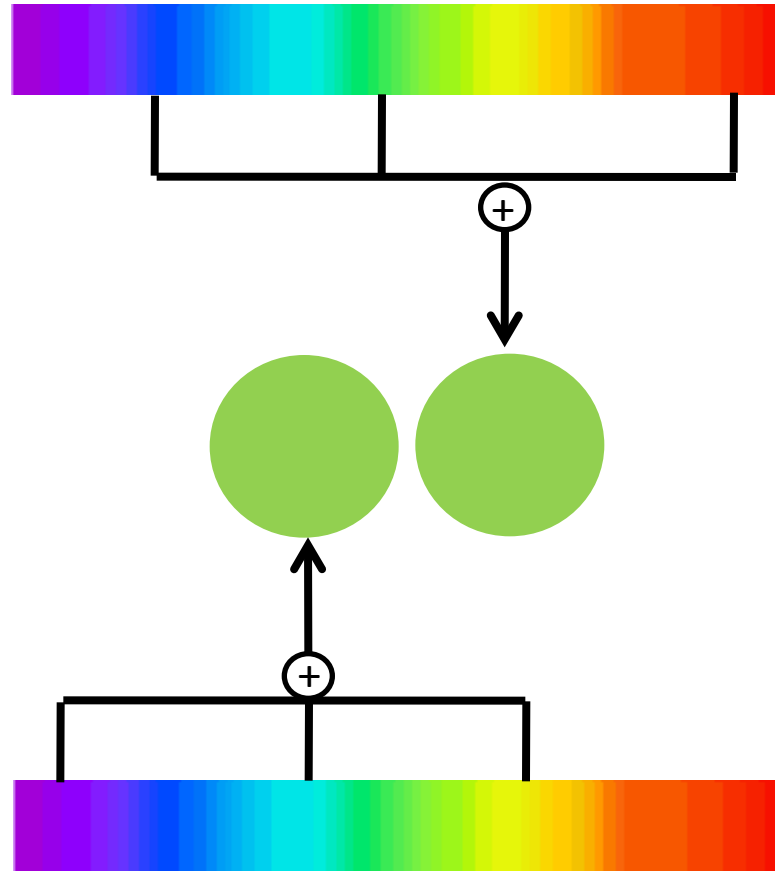
Note: The  $\lambda_{\max}$ 's are shifted *in vivo* to 445, 540 and 565 nm. This is due to the transmission properties of the intervening ocular media (lens, macular pigment).

# Absorption spectra





# Trichromacy means our color vision is limited

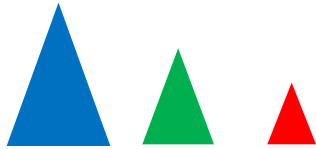


**So, if each photoreceptor is color-blind, how do we see color?**

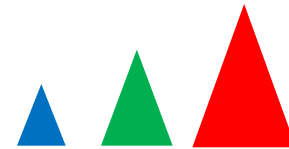
**The perception of color is created by postreceptoral pathways,  
but we will come to that later...**

# Colors as relative responses (ratios)

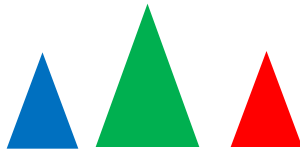
Blue light



Red light



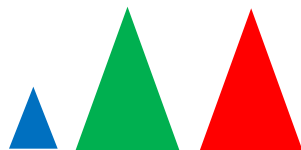
Green light



Purple light



Yellow light



White light

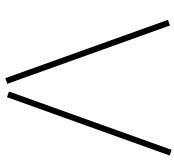


# Color vision deficiencies

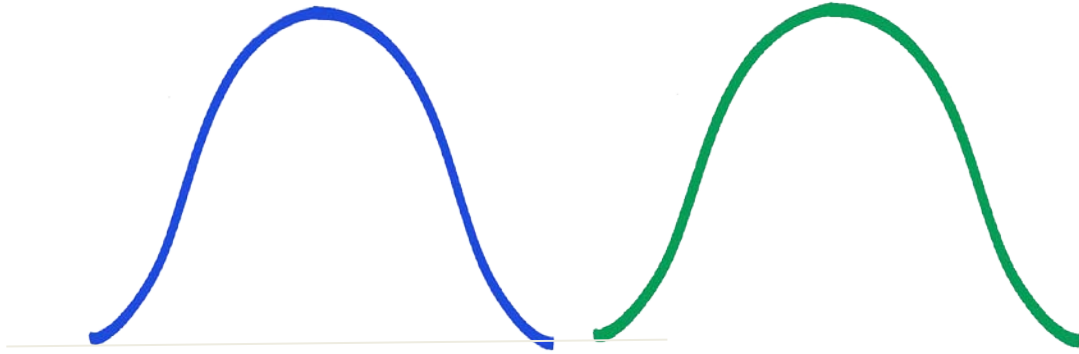
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**Protan**  Protanope (missing red cone pigment)  
Protanomalous trichromat (shifted red cone pigment absorption)

**Deutan**  Deuteranope (missing green cone pigment)  
Deuteranomalous trichromat (shifted green cone pigment absorption)

**Tritan**  Tritanope (missing blue cone pigment)  
Tritanomalous trichromat (shifted blue cone pigment absorption)

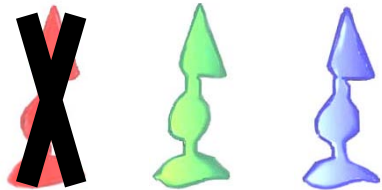
# Visualizing Protanopia



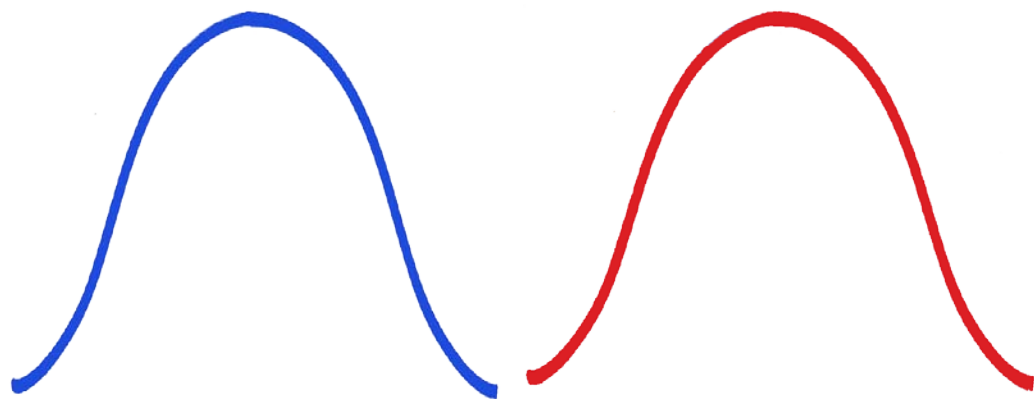
Google™



Google™



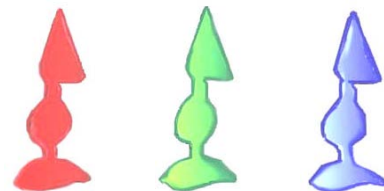
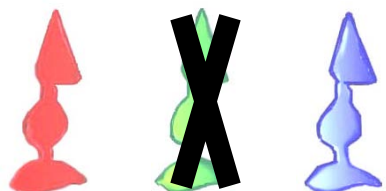
# Visualizing Deuteranopia



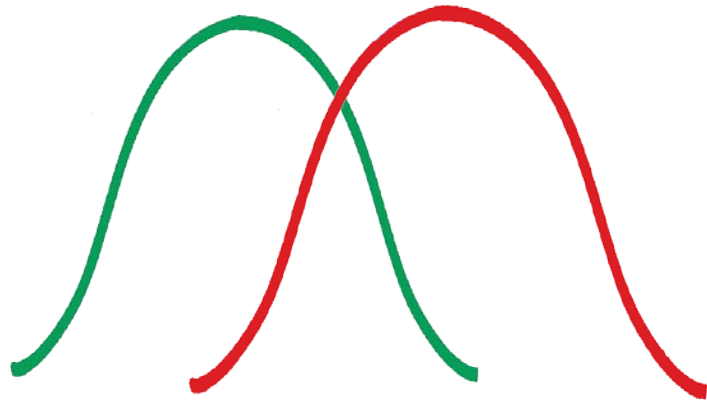
Google™



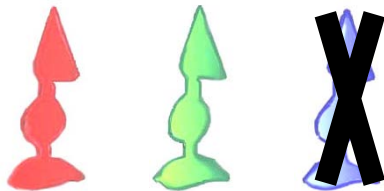
Google™



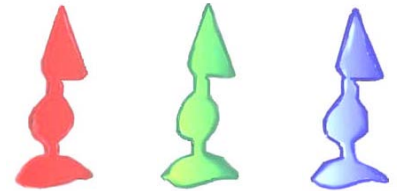
# Visualizing Tritanopia



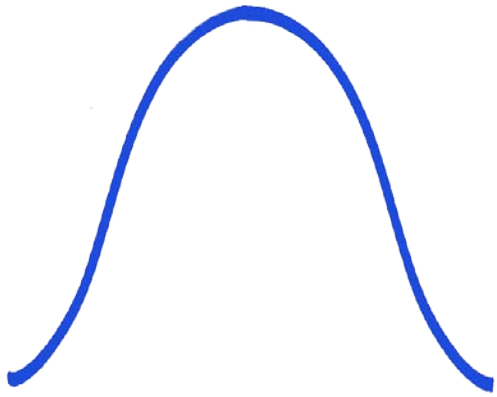
Google™



Google™



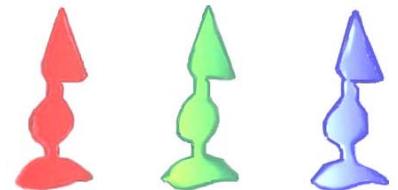
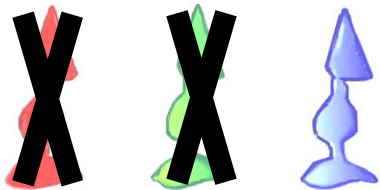
# More color vision deficiencies...



Google™



Google™







# Consequences of color vision deficiencies

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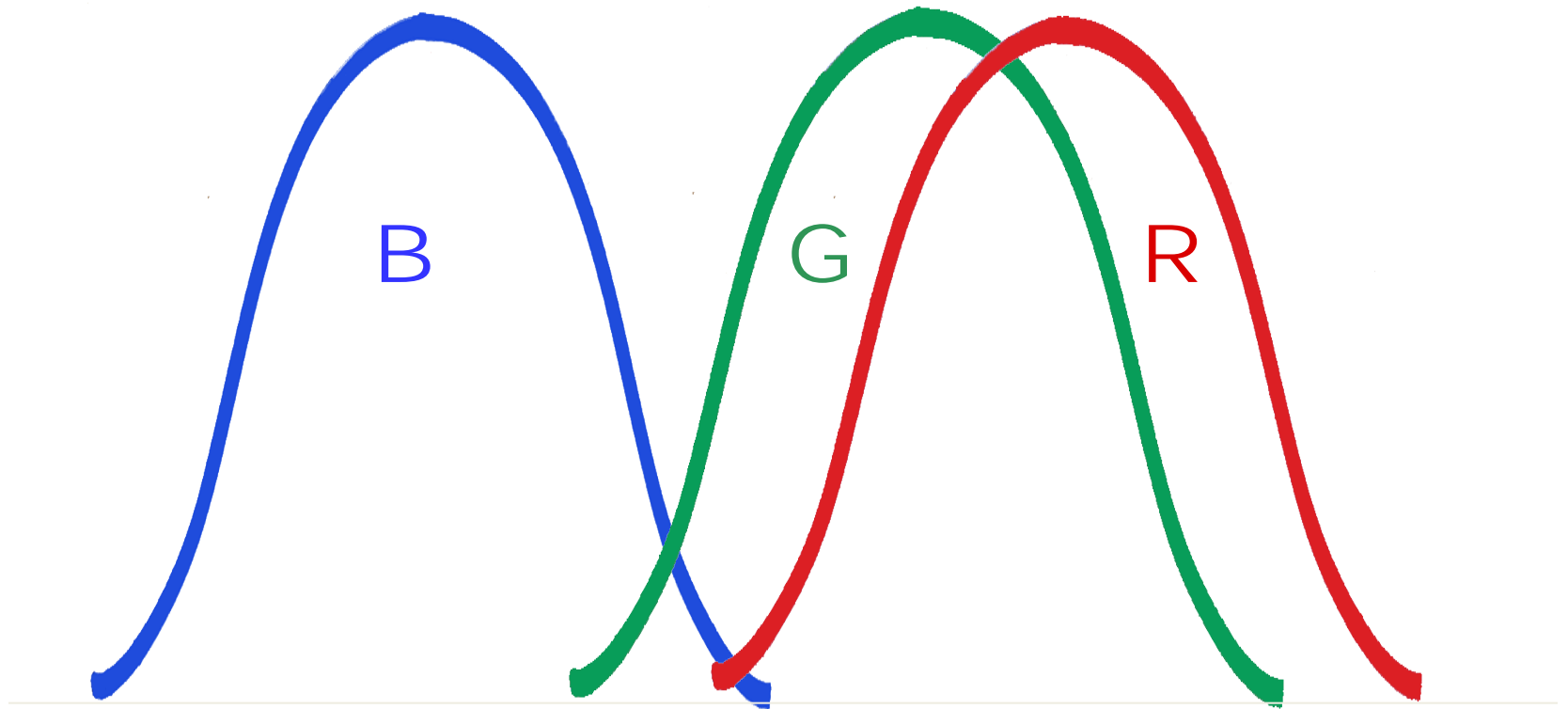
**Normal trichromats can distinguish between 150 distinct wavelengths**

**Protanopes can only distinguish between 21 distinct wavelengths**

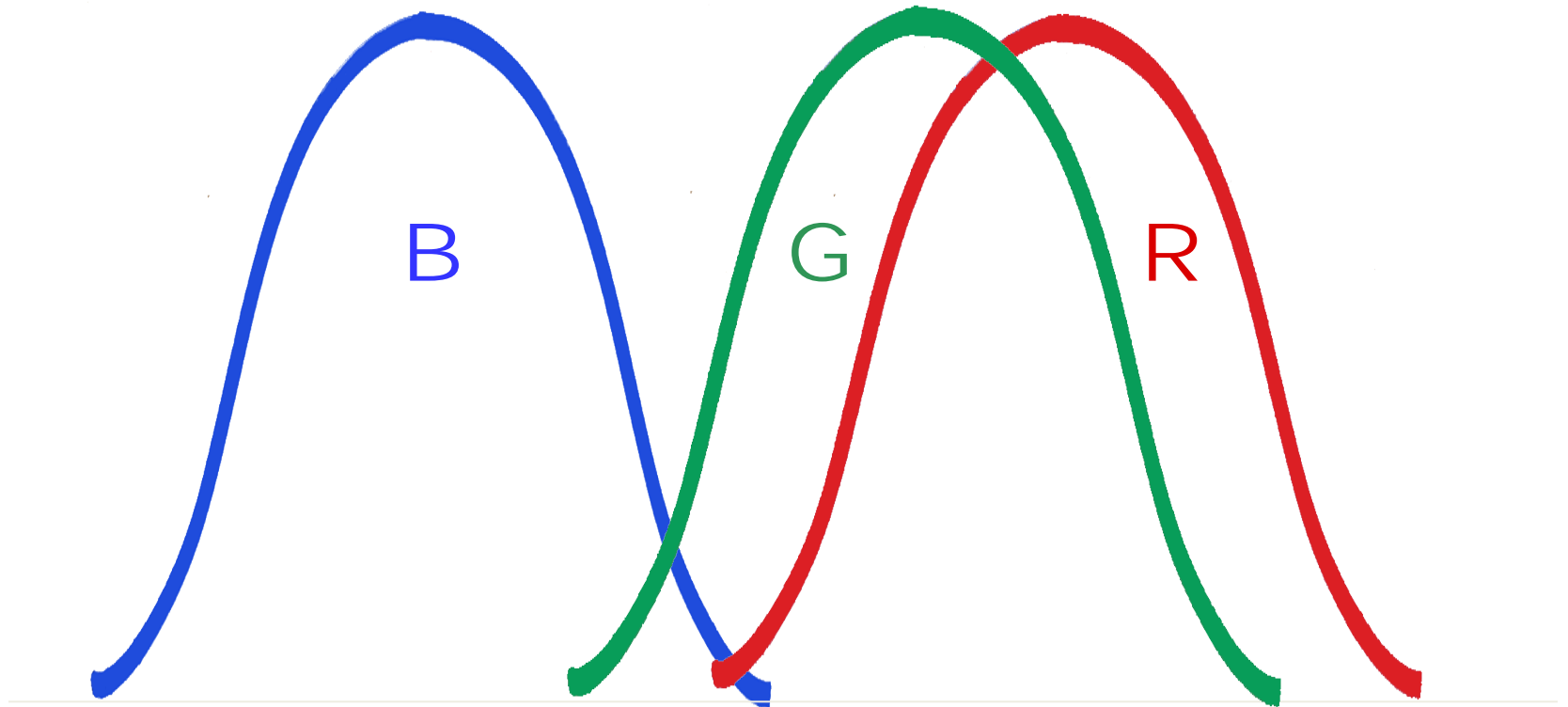
**Deuteranopes can only distinguish between 31 distinct wavelengths**

**Tritanopes can only distinguish between 44 distinct wavelengths**

# Protanomaly (red shifted towards green)

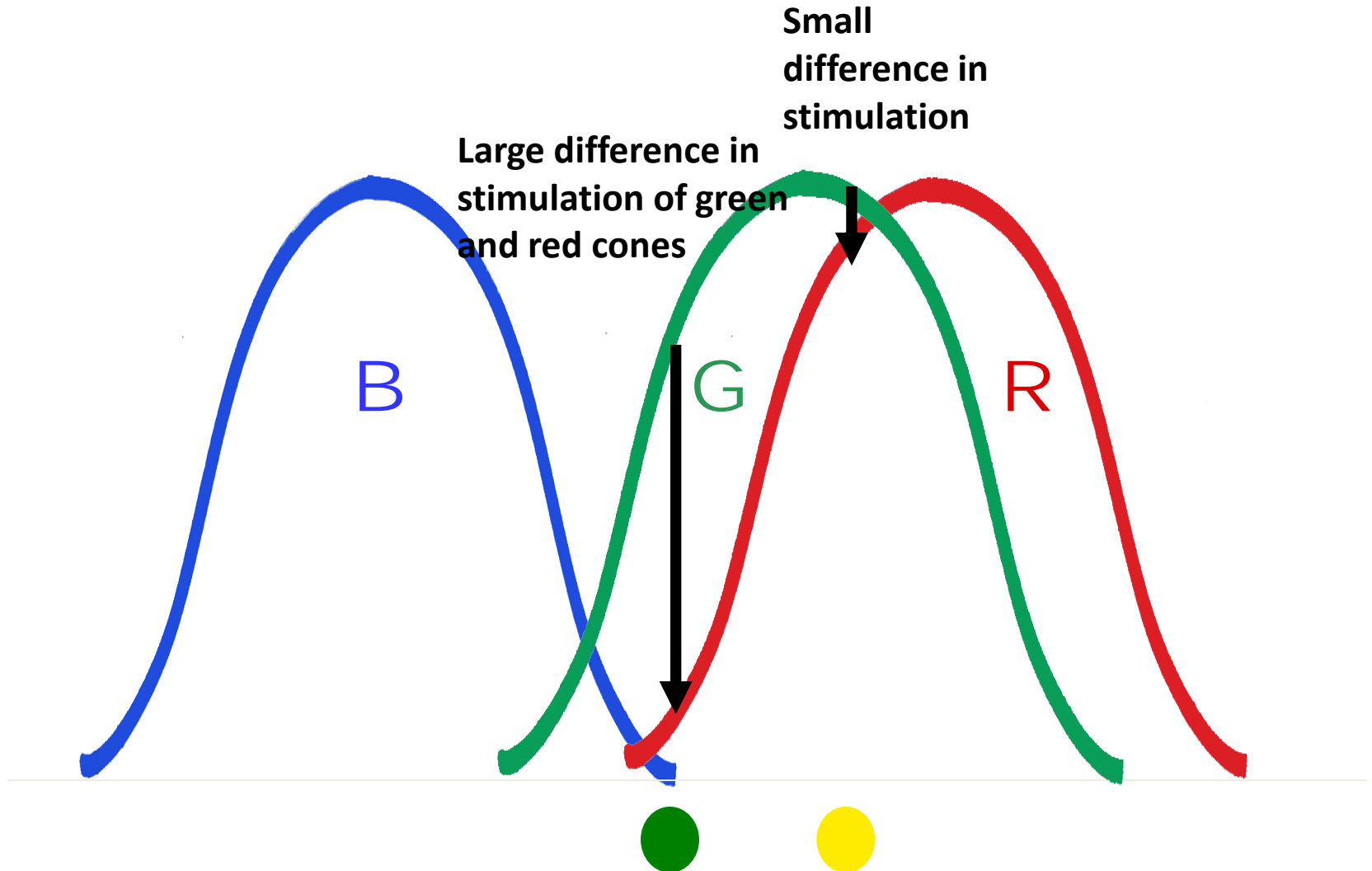


# Deuteranomaly (green shifted towards red)

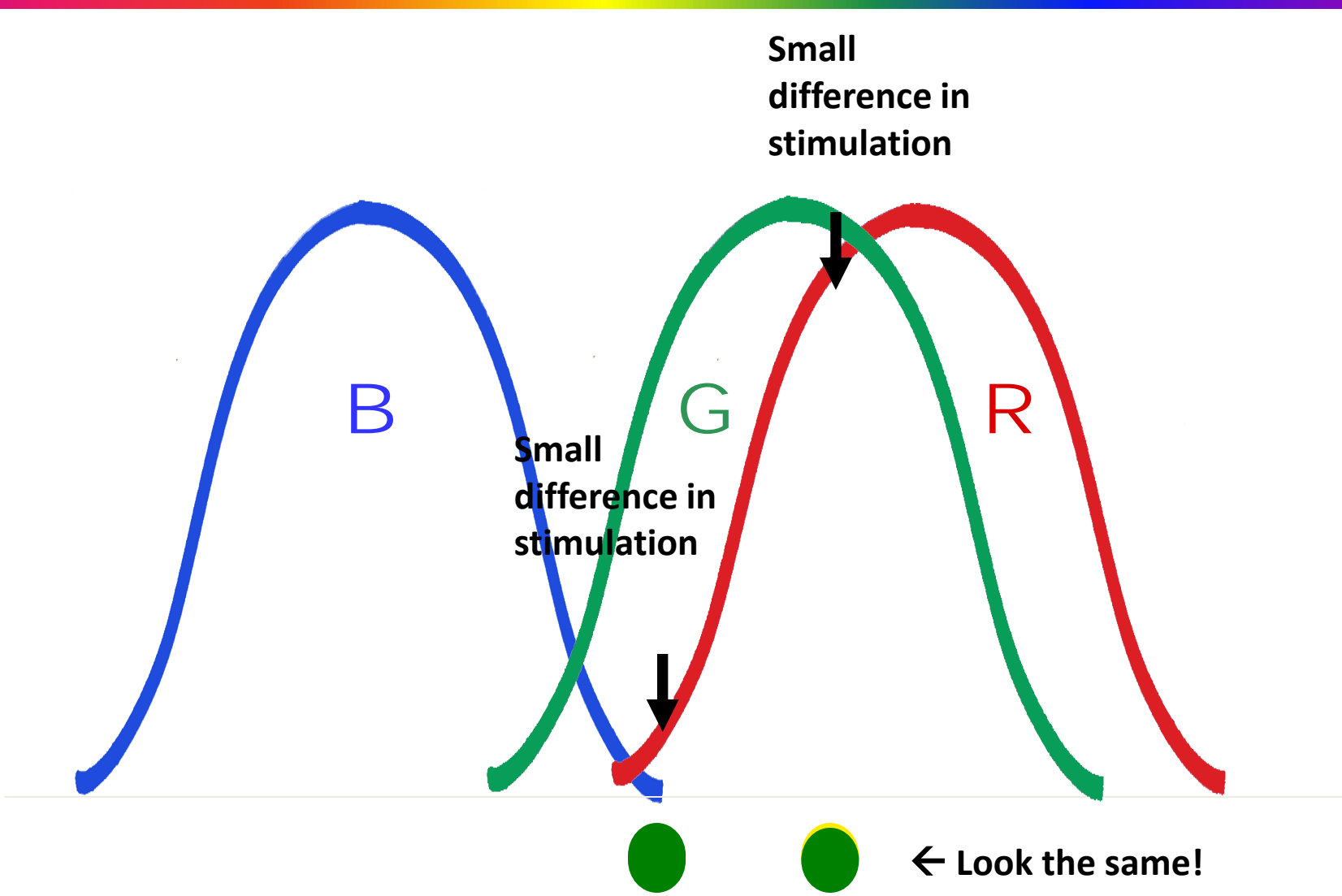


**Why do colors that look different to us appear the same to color deficient individuals?**

# Consider a green versus a yellow light...



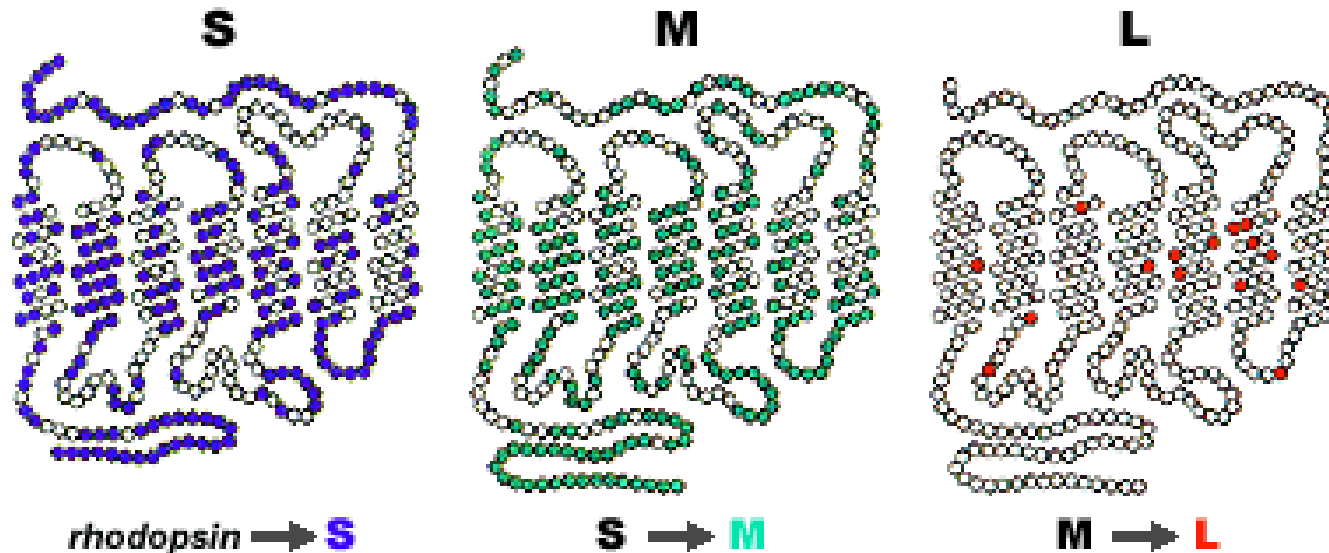
# ...this is the perception of a deuteranomalous trichromat



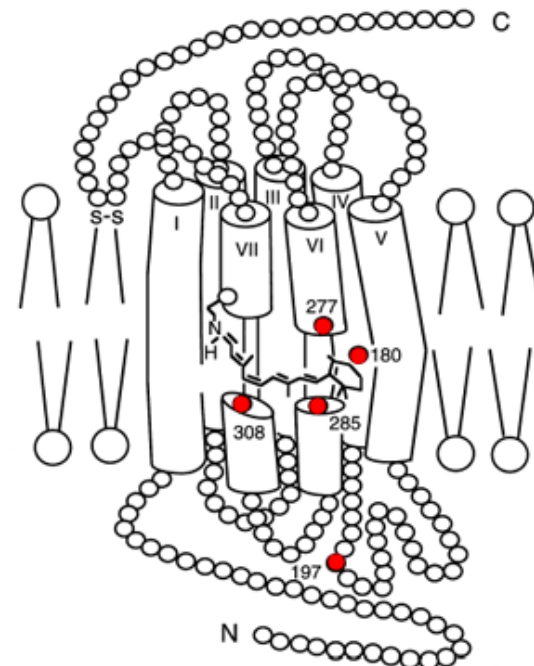
# Prevalences of color deficiencies

Color vision deficiency	Males	Females
Protanomaly	1%	0.03%
Protanopia	1%	0.02%
Deuteranomaly	5%	0.4%
Deuteranopia	1%	0.01%
Tritanomaly	Rare (if at all)	Rare (if at all)
Tritanopia	0.008%	0.008%

# Genetic background of color vision



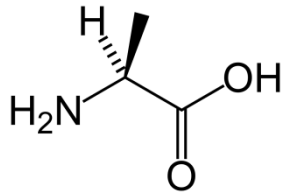
● spectral tuning sites: 180, 277, 285





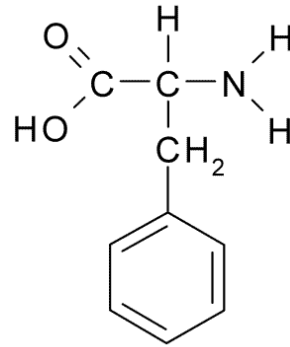
# Spectral tuning sites shift the $\lambda_{max}$ of the respective opsin

N  
O  
N-  
P  
O  
L  
A  
R



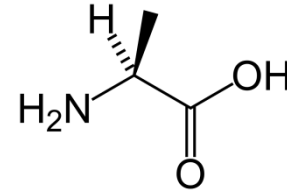
Alanine

**A180S**



Phenylalanine

**F277Y**

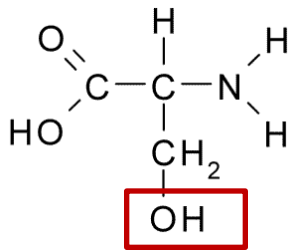


Alanine

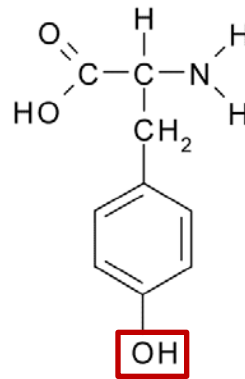
**A285T**

green cone opsin

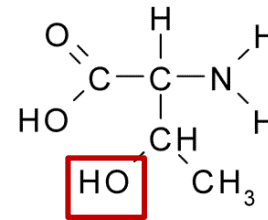
P  
O  
L  
A  
R



Serine



Tyrosine

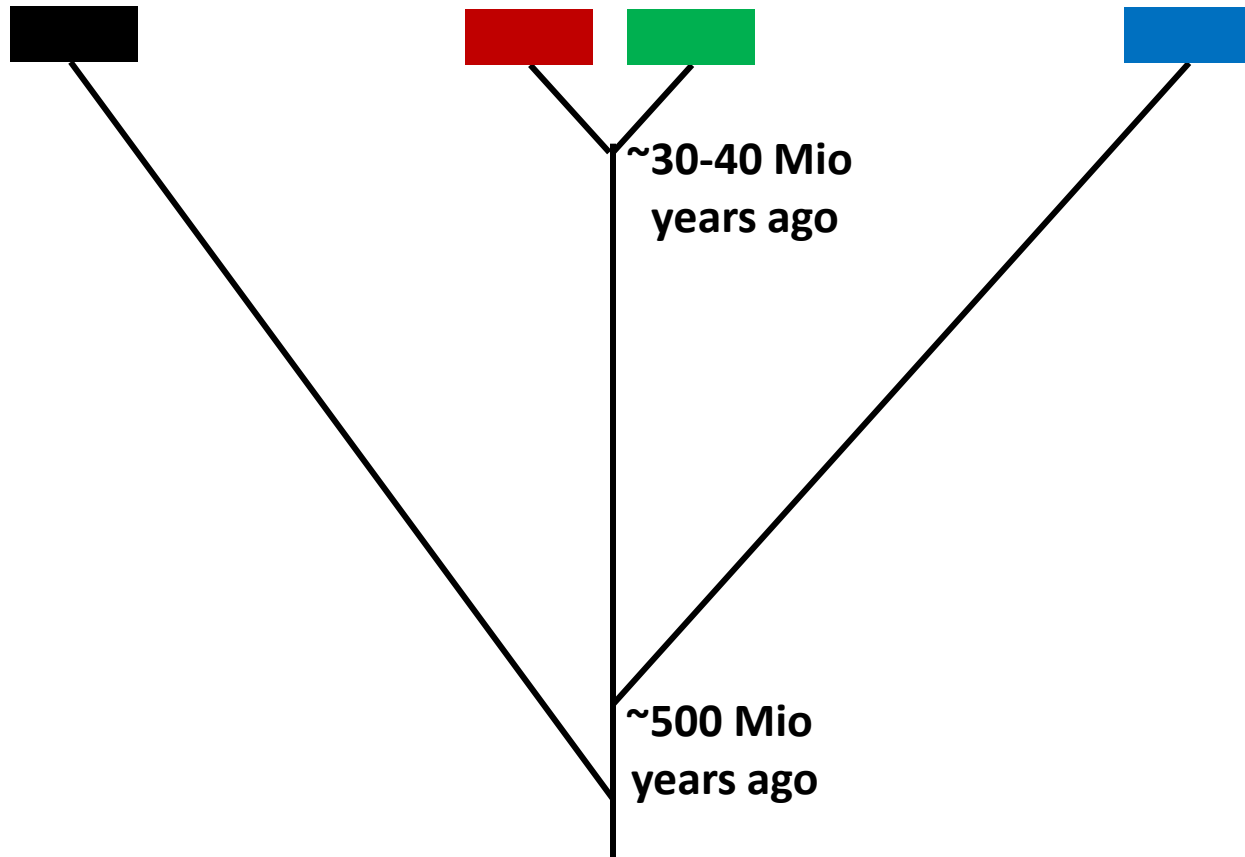


Threonine

red cone opsin  
all with OH group

**Why are the M- and L-cone opsins so similar?**

# Evolution of trichromacy



The selective advantage of trichromatic vision is thought to be the ability to detect ripe fruits against a background of dense green foliage.

## No red-green discrimination



## Red-green discrimination



# Red and green cone opsin genes



X

Crossing over



Hybrid gene

OR



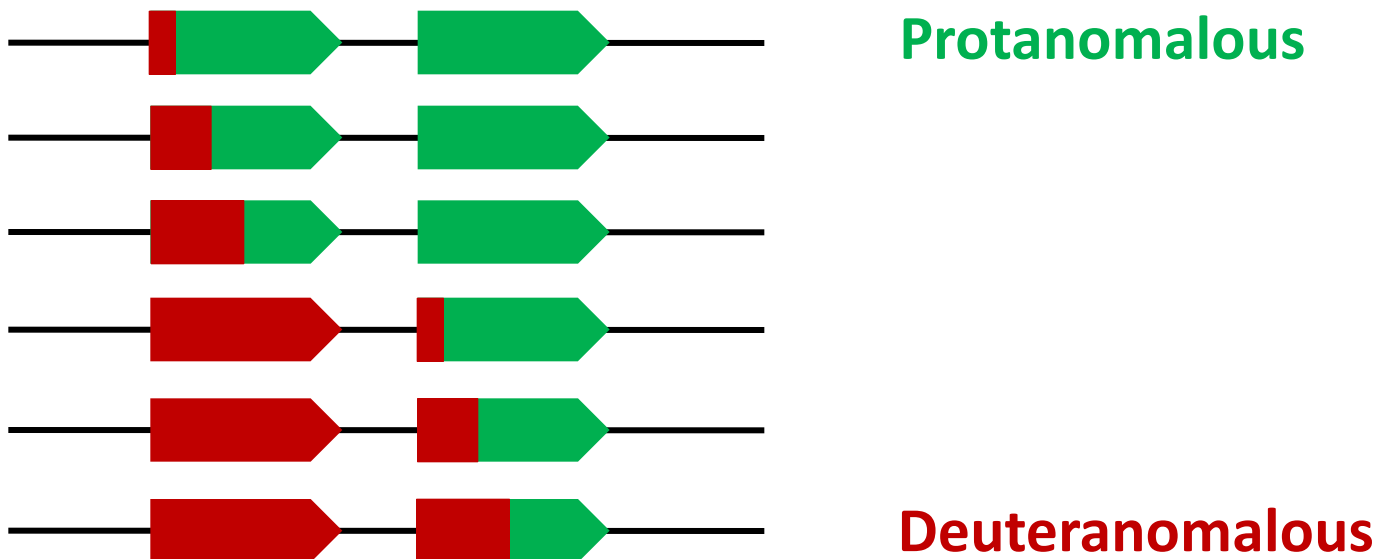
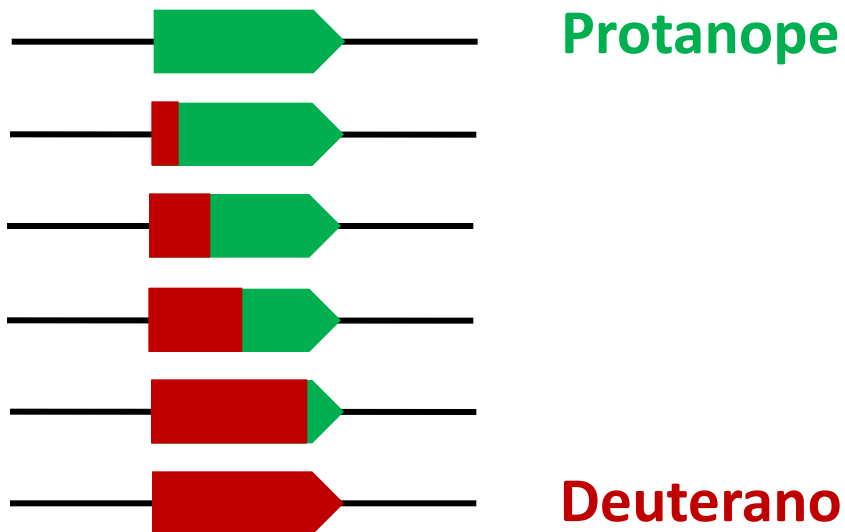
Loss of gene

OR



Gene duplication

# Red and green color deficiencies

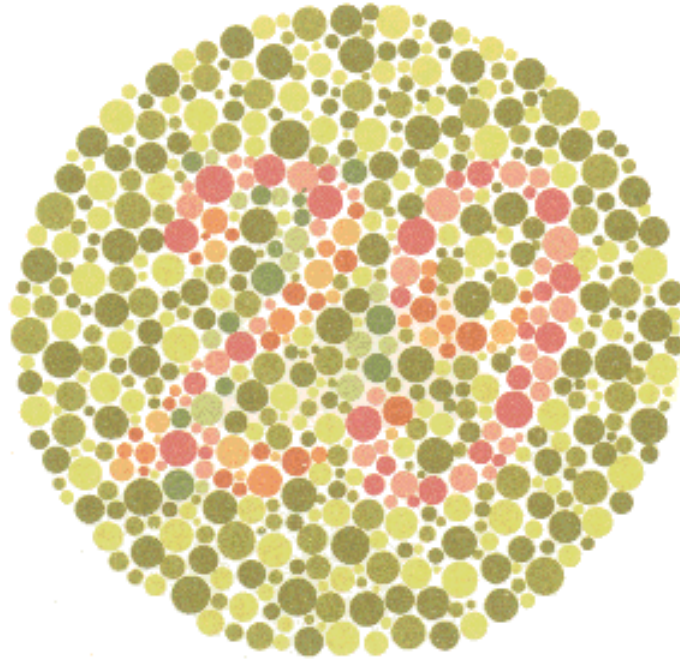


## **Diagnosing color vision deficiencies**



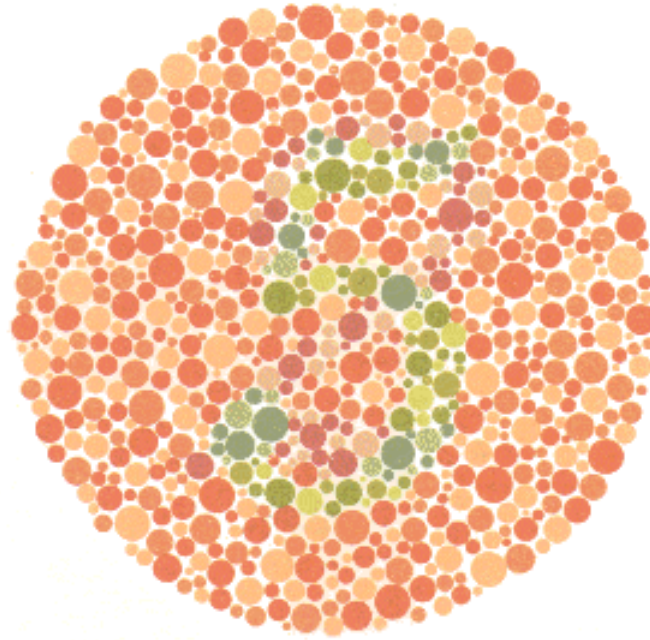
# A quick color vision test...

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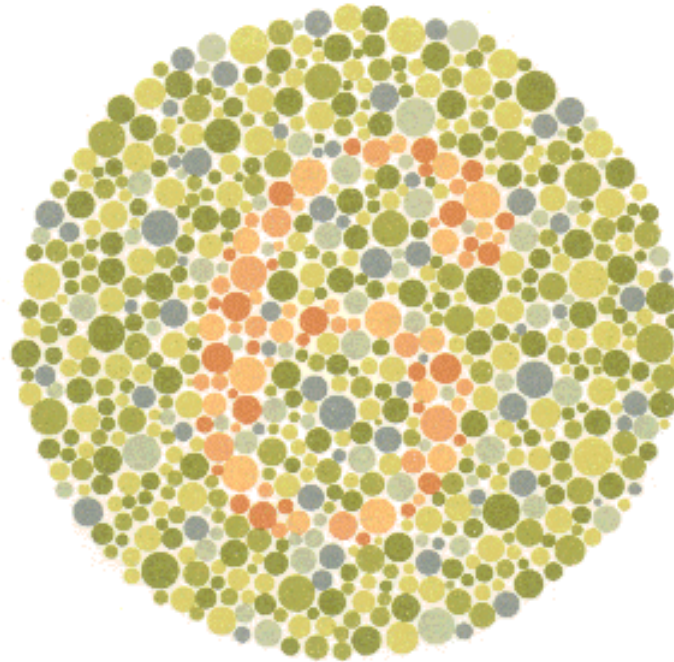


# A quick color vision test...



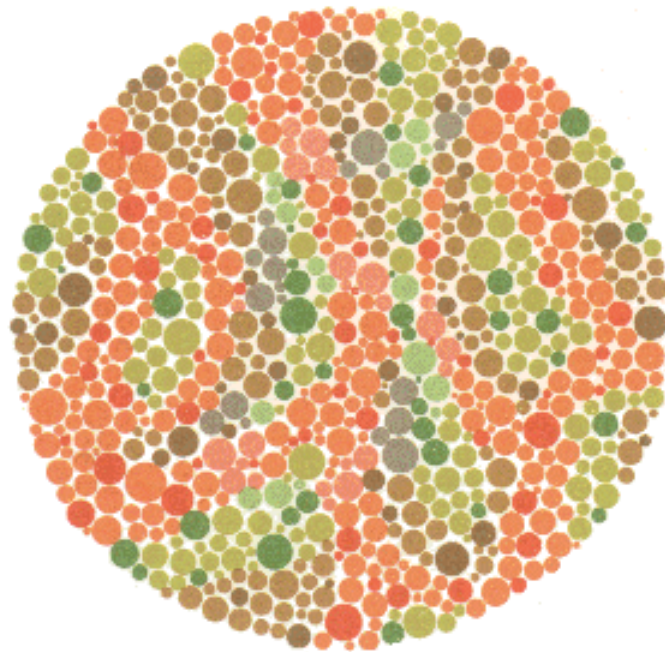
# A quick color vision test...

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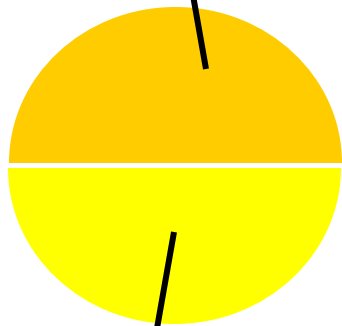
# A quick color vision test...

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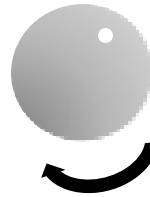
# Diagnosis of red-green color deficiencies: Anomaloscope

red/green mix (=yellow)



pure yellow

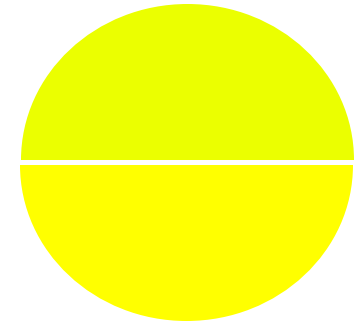
Green



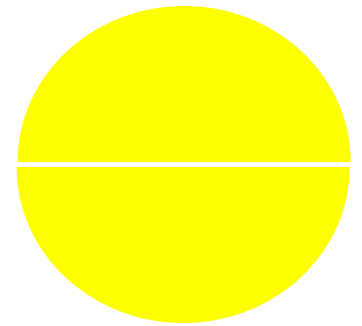
Red



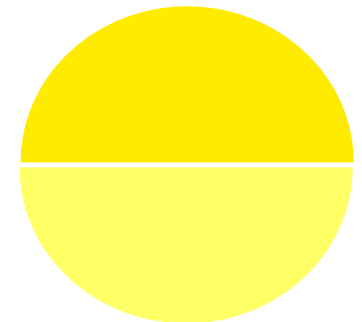
Yellow intensity



Deuteranope match



Normal trichromate match



Protanope match

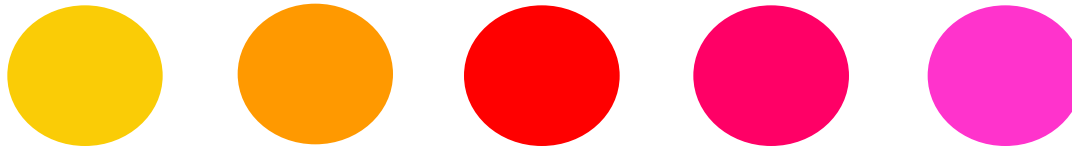
## **Postreceptoral color vision**

# Who knows this gentleman?

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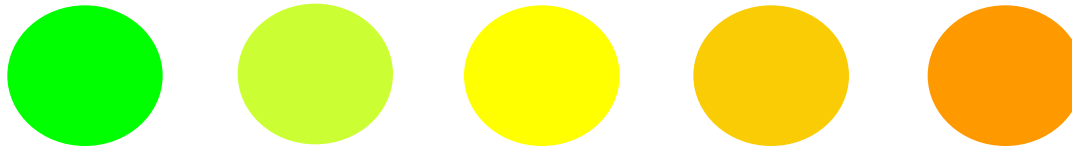


**Ewald Hering 1834–1918**

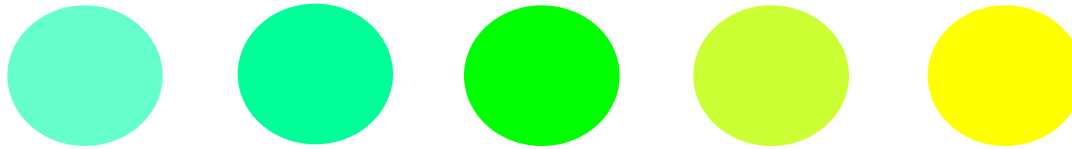


**Reds can get bluer or yellower but not greener**

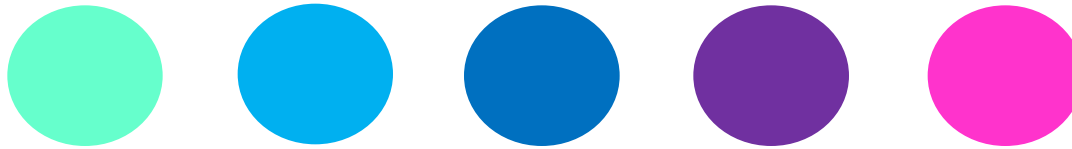




**Yellows can get greener or redder but not bluer**



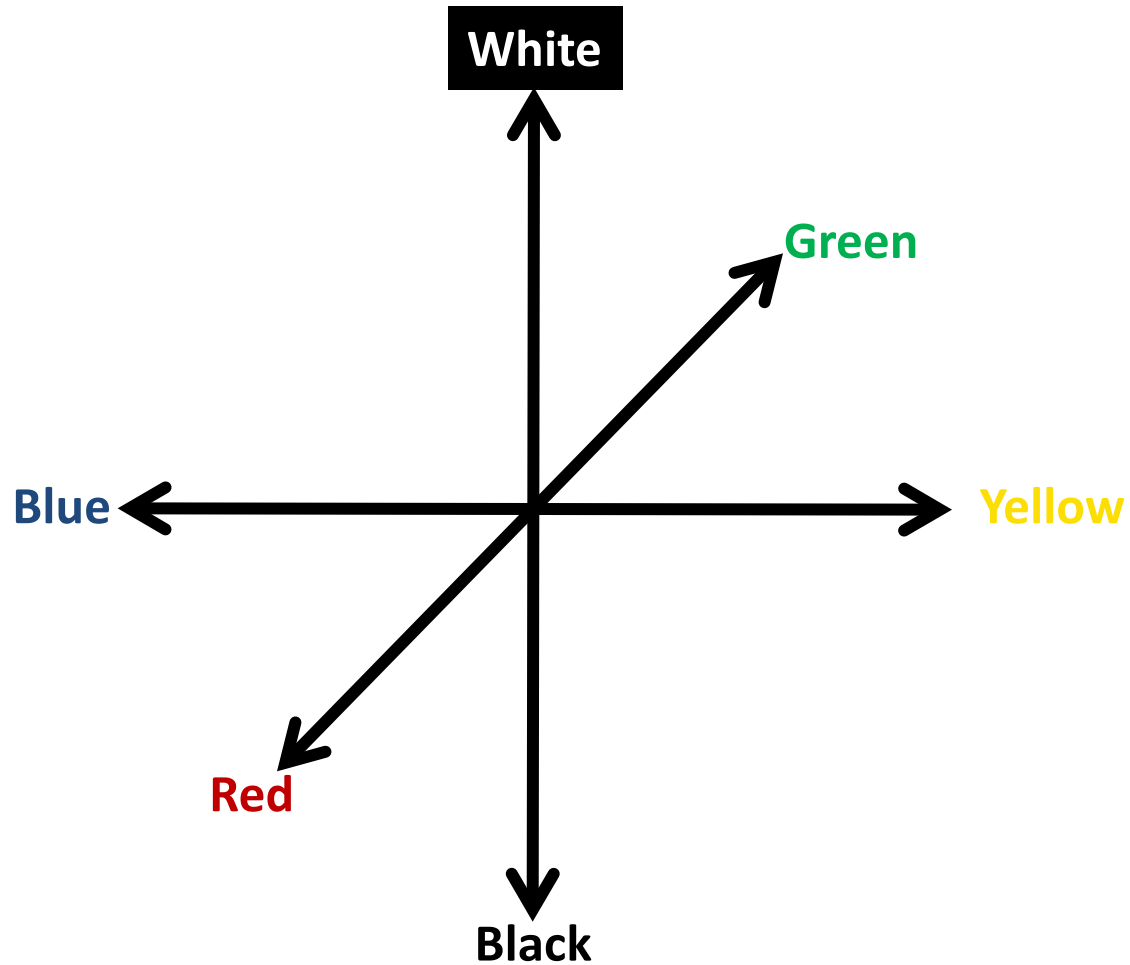
**Greens can get bluer or yellower but not redder**



**Blues can get greener or redder but not yellower**

# The color opponent theory of Hering

Four „Urfarben“ are arranged in two opponent processes



# Opponent channels



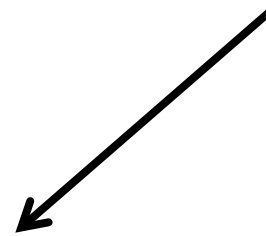
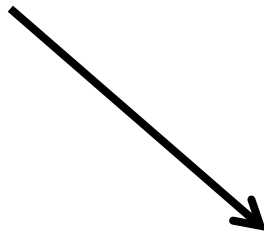
B-W channel



R-G channel



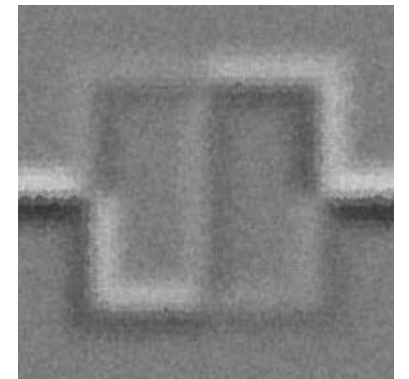
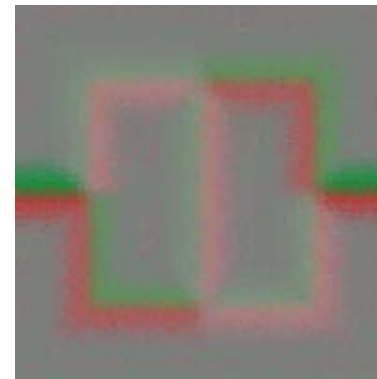
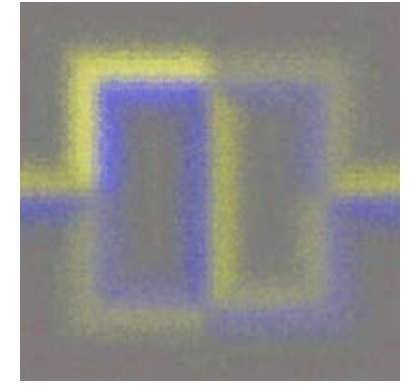
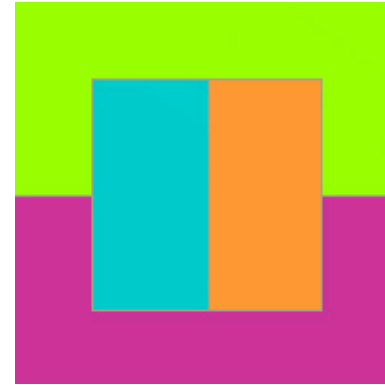
B-Y channel





# Illustration of how the opponency channels work in your perception

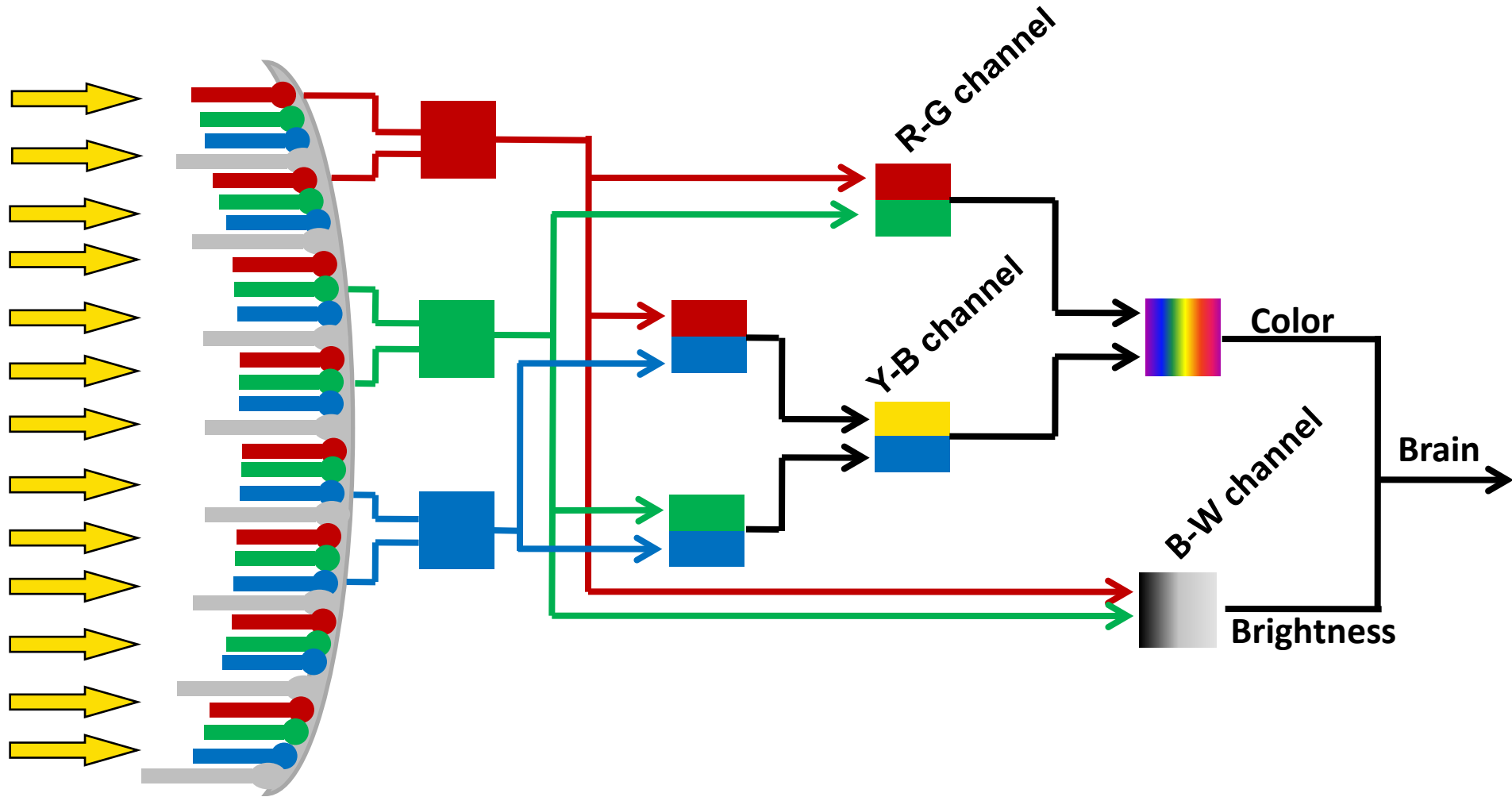
- Here are the *enhanced edges* resulting from your *y-b chromatic channel*
- Here are the *enhanced edges* resulting from your *r-g chromatic channel*
- Here are the *enhanced edges* resulting from your *r-g chromatic channel*



The artist Van Gogh often used opponent colors to enhance them

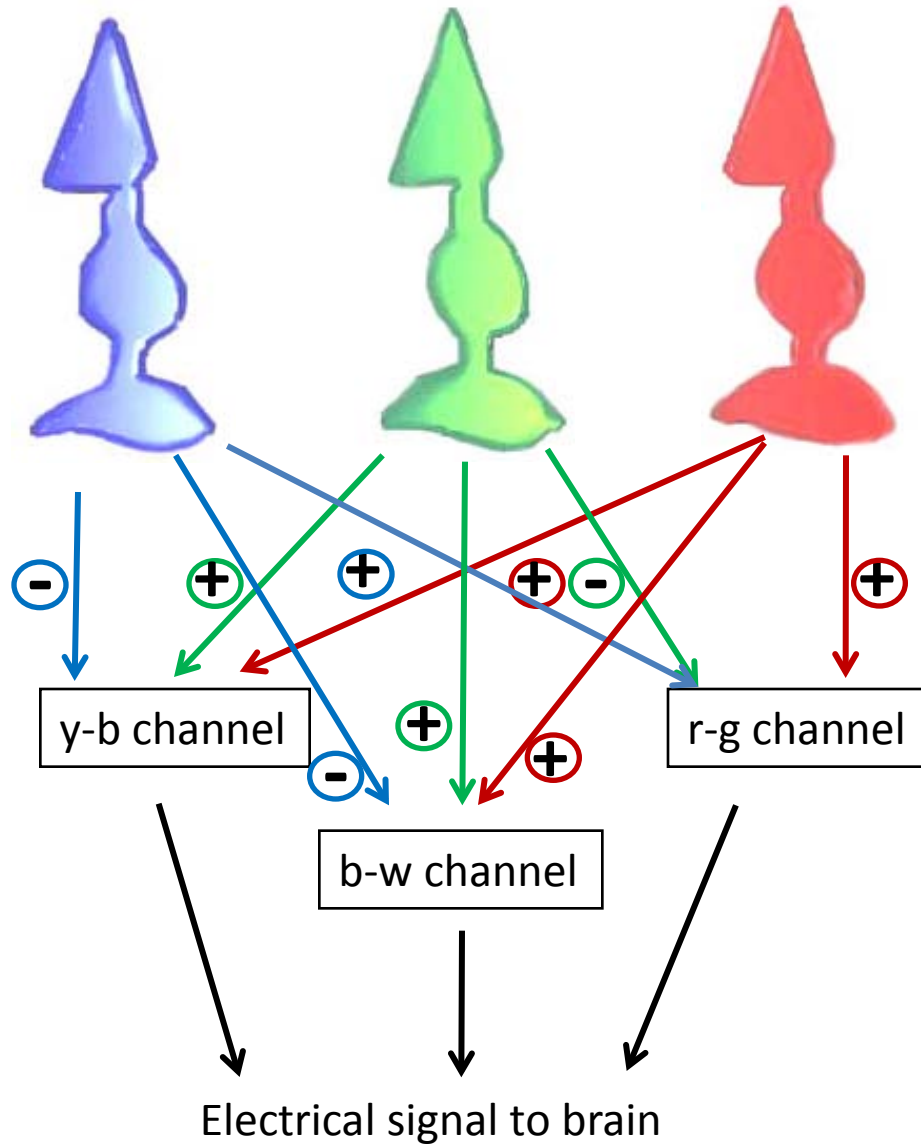


# How are cone outputs organised at subsequent stages of visual processing?





# Opponent channels



**So far, we've mainly been talking about the colours of isolated patches of light. But the colour of a patch depends also upon:**

- What precedes it (in time)**

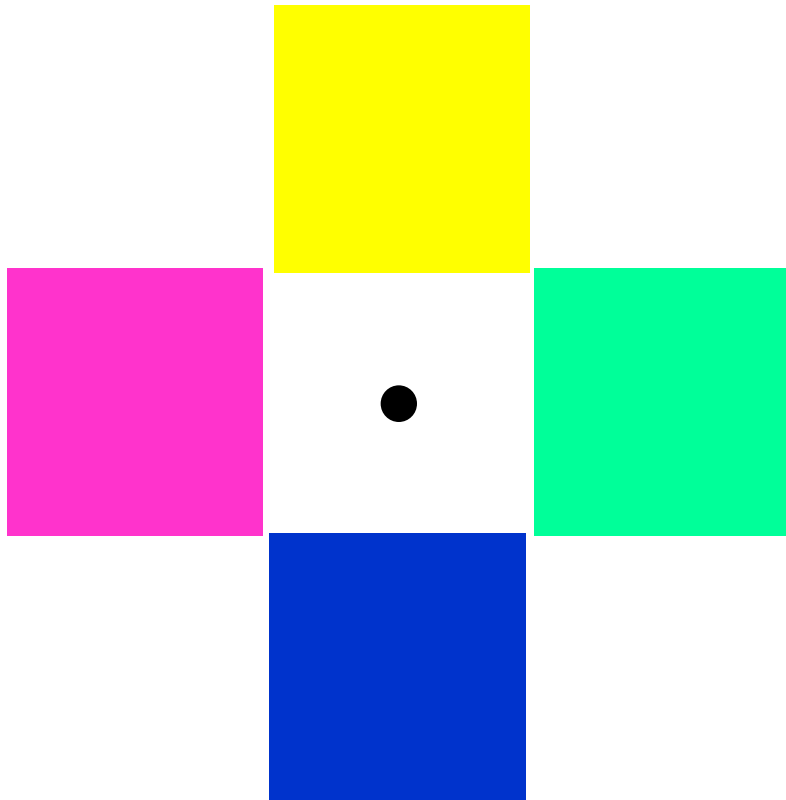
**COLOR AFTER-EFFECTS**

- What surrounds it (in space)**

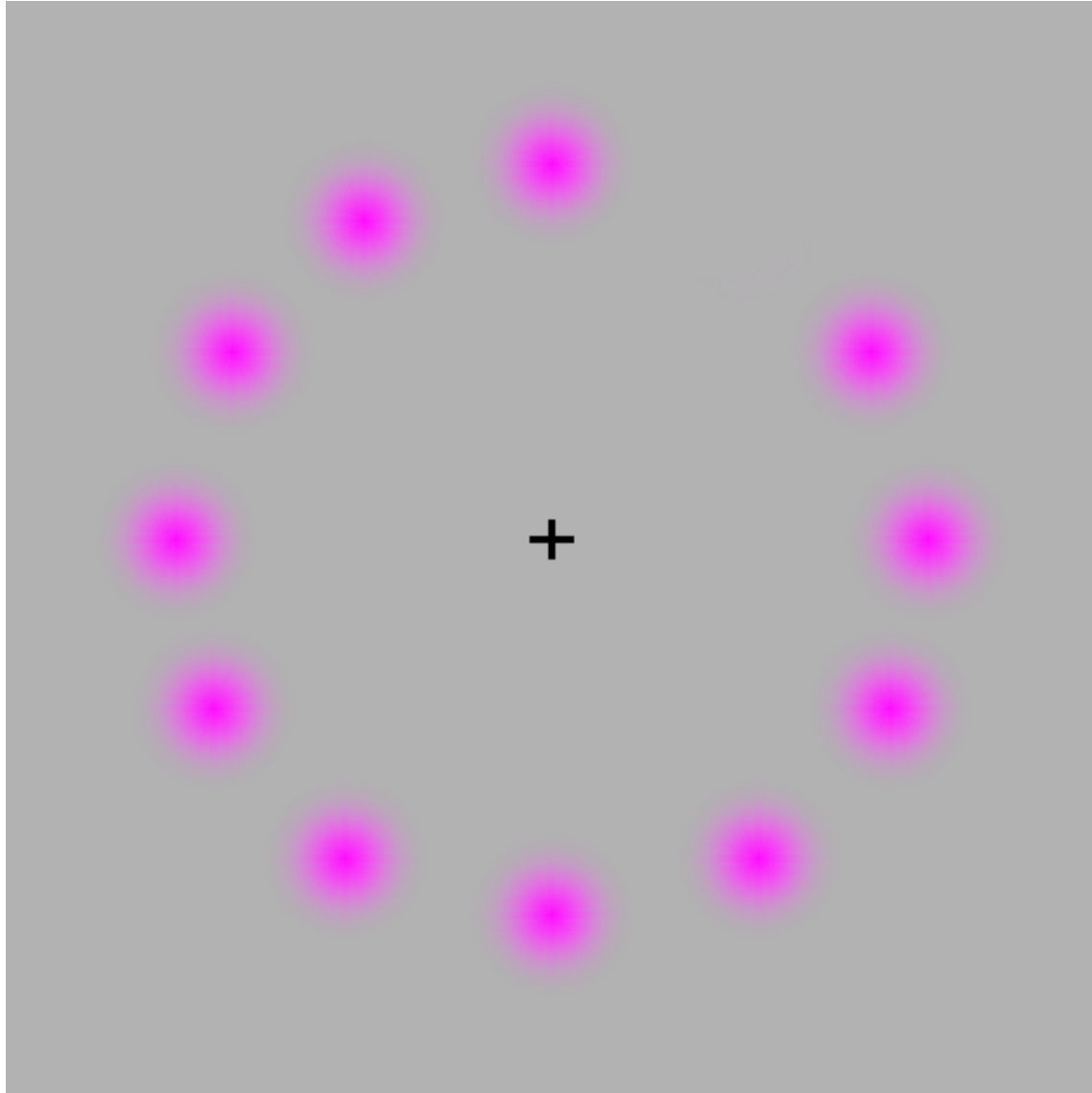
**COLOR CONSTANCY**

# Color after-effect: Successive contrast

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# Color after effect: The lilac chaser

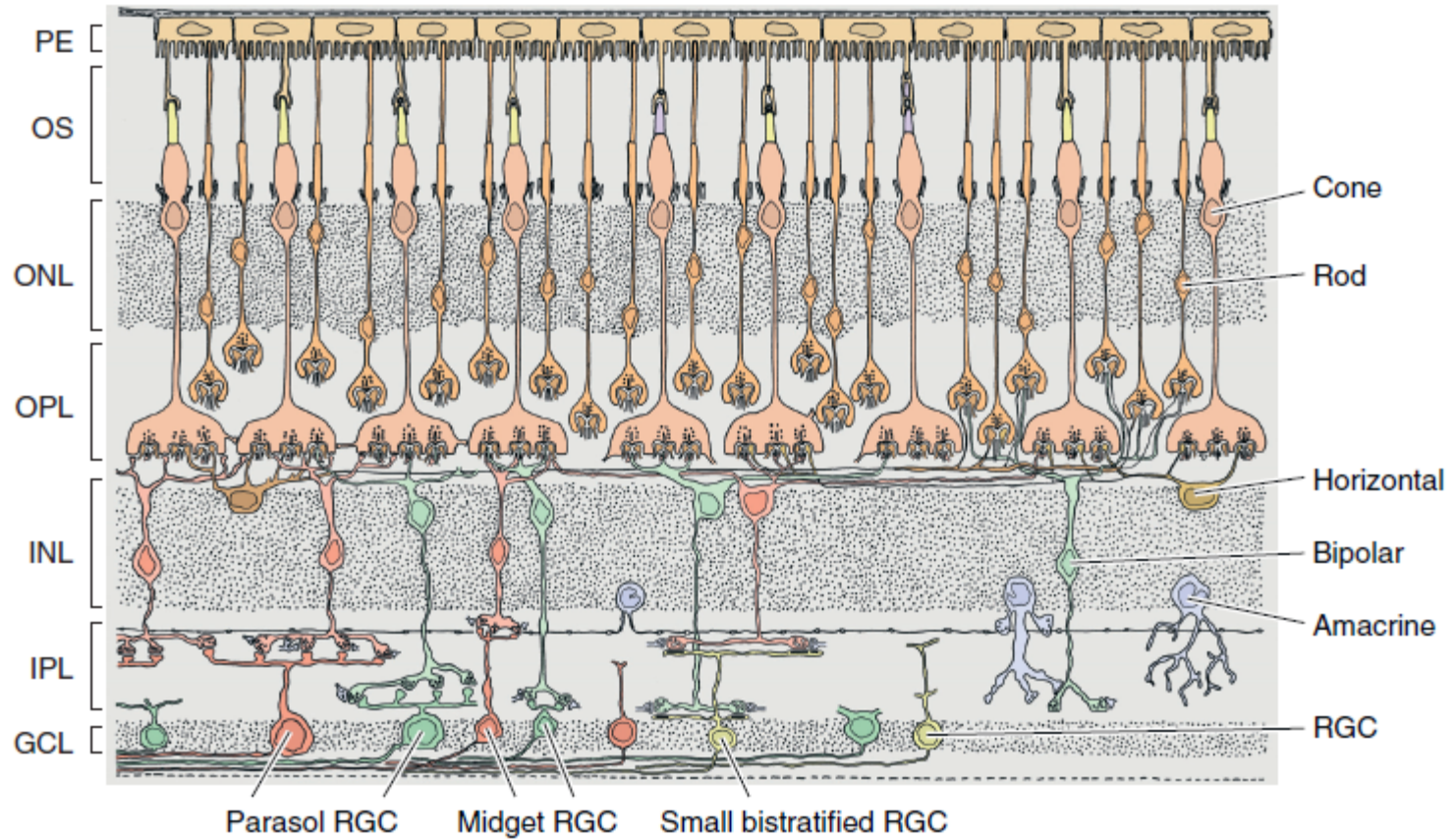


# Color vision is a two stage model

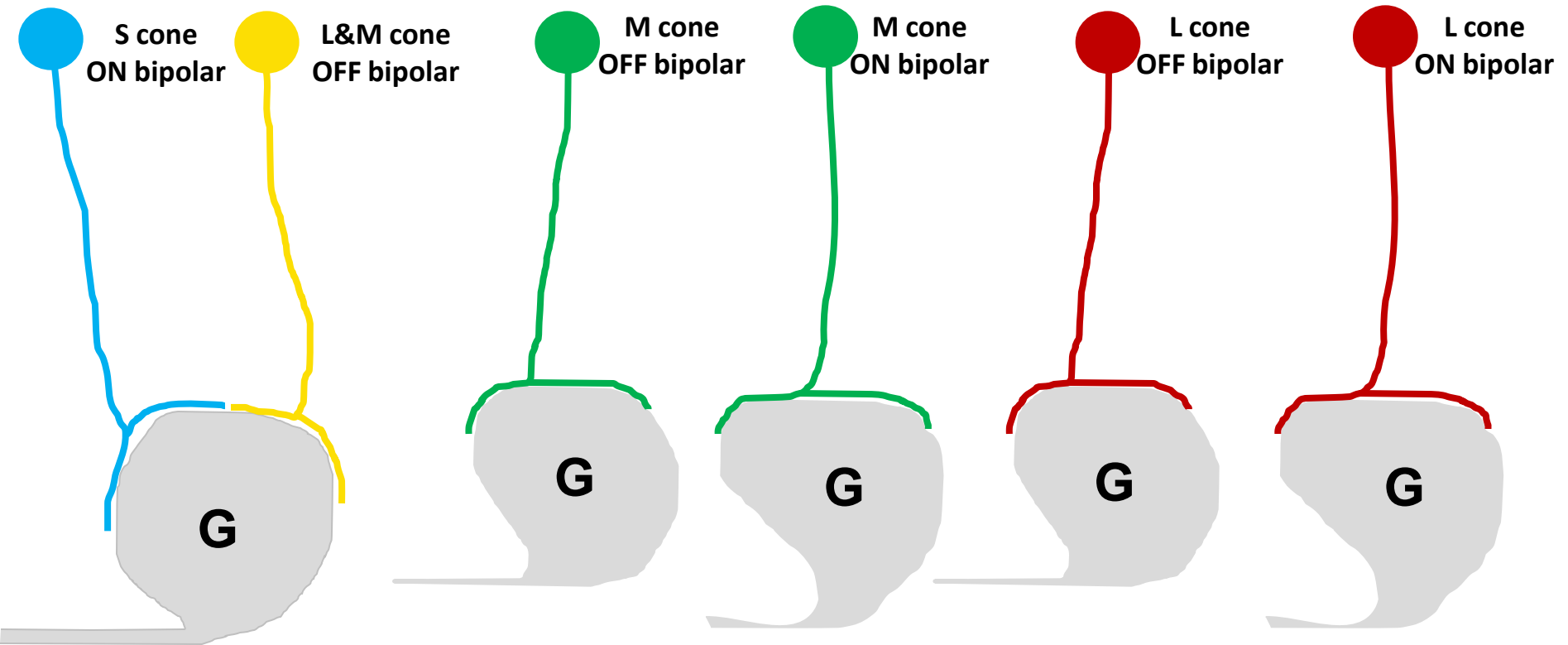
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- **Early processing is trichromatic**
- **Later on it is opponent processing**

# The physiological basis of opponency



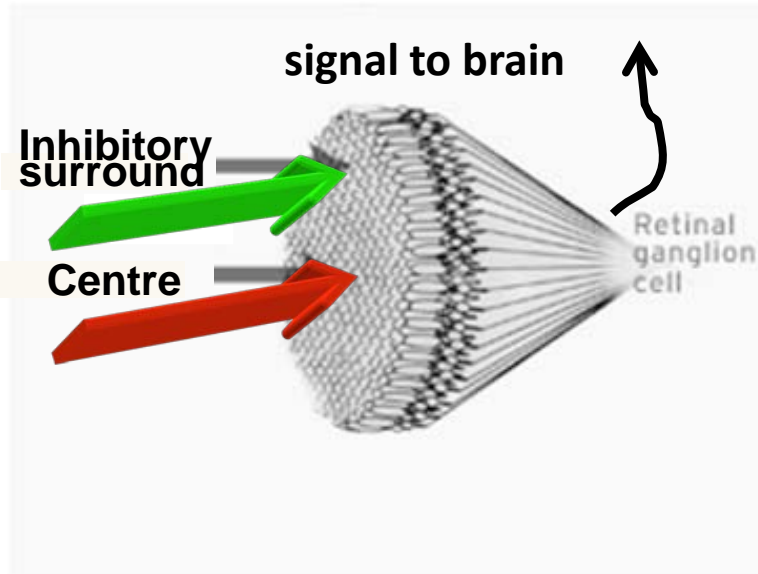
# Ganglion and bipolar cells of trivariant color vision



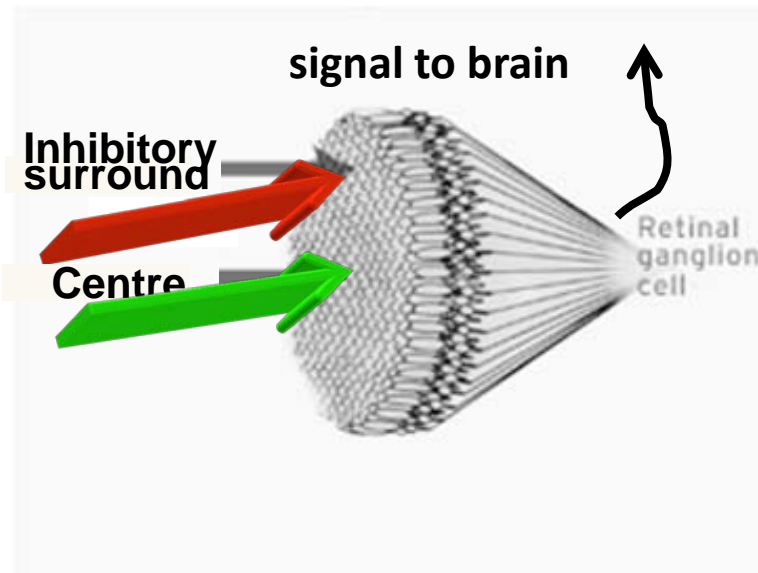
**Bistratified S cone  
ON cell**

**Midget cell system in the fovea**

# Opponent receptive fields in our retina



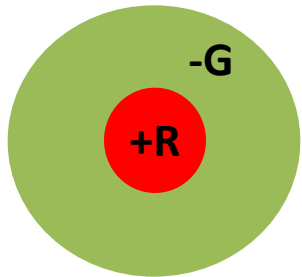
RED



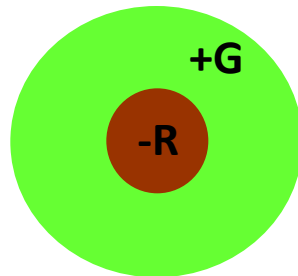
GREEN



# The physiological basis of opponency: opponent retinal ganglion cells



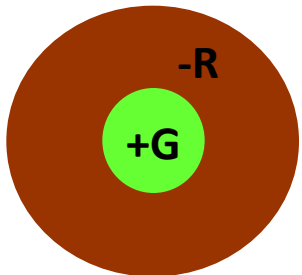
OR



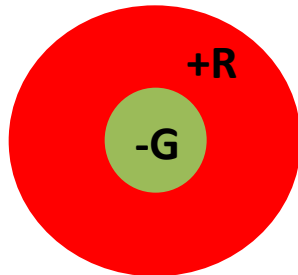
→ signal increase → „red“

red ON/green OFF

red OFF/green ON



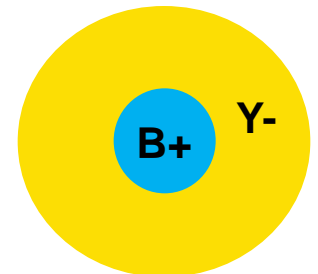
OR



→ signal decrease → „green“

green ON/red OFF

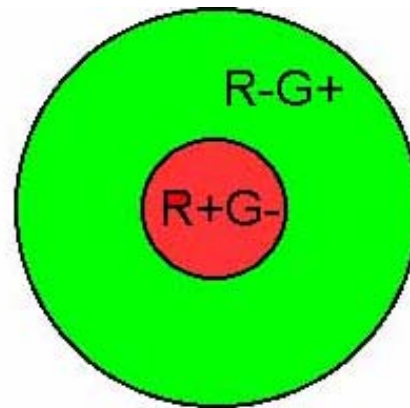
green OFF/red ON



blue ON/yellow OFF

# Double opponent cells

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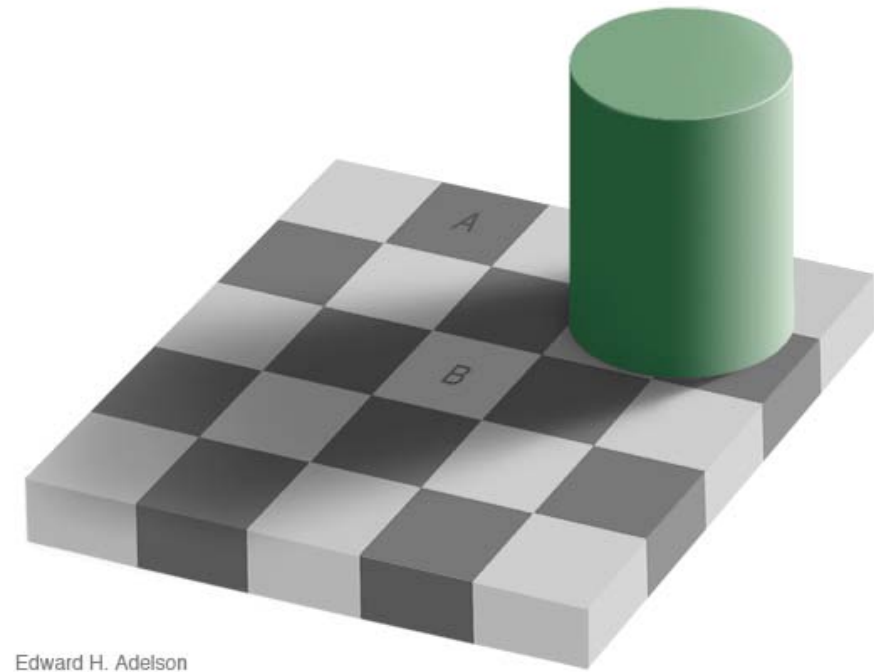


# Color constancy

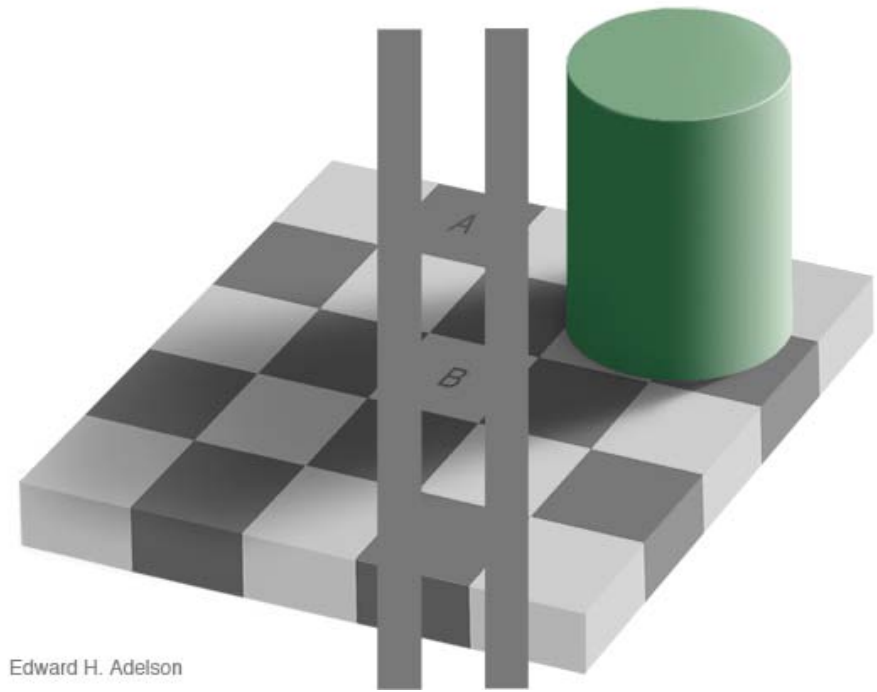




# Color constancy



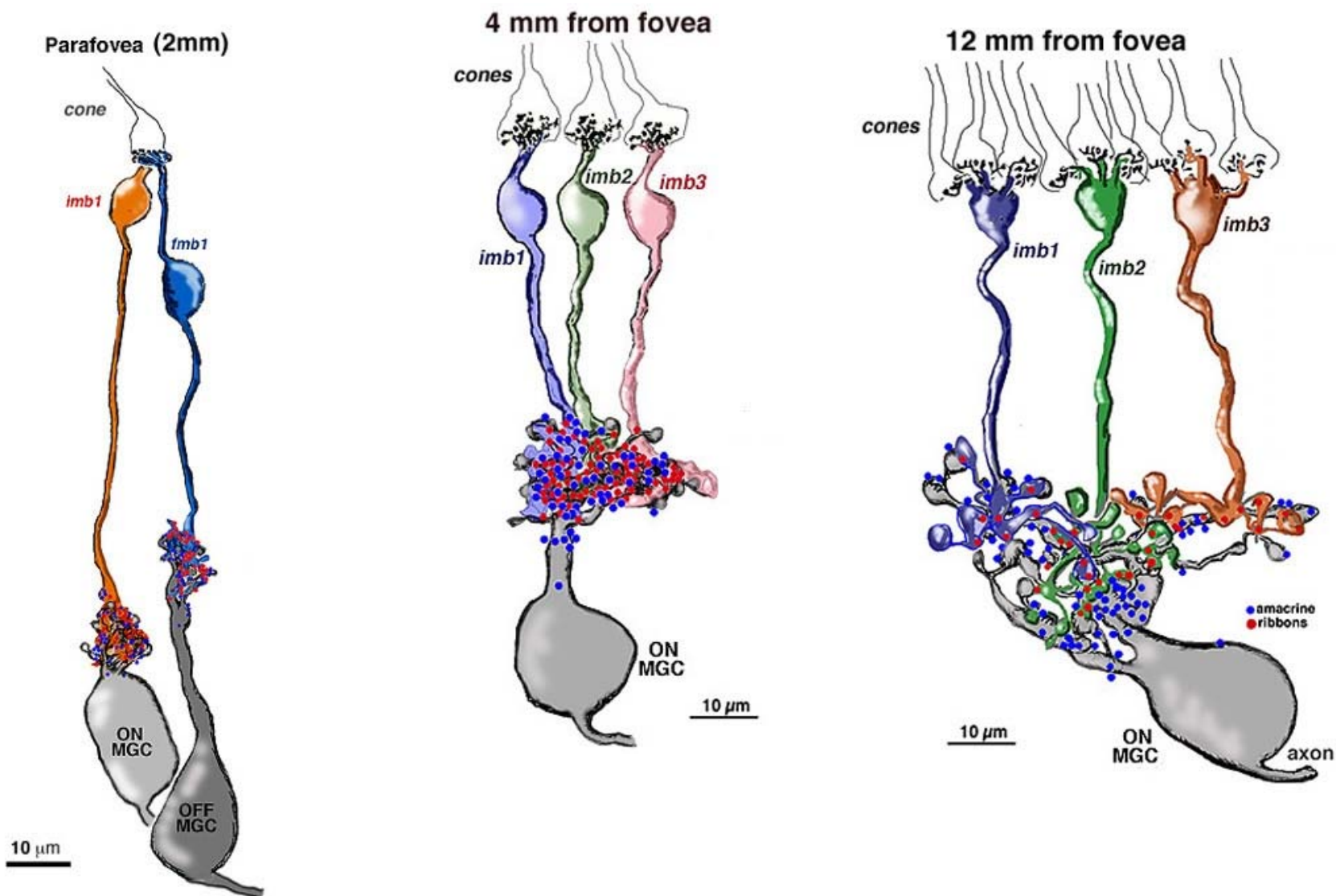
Edward H. Adelson



Edward H. Adelson

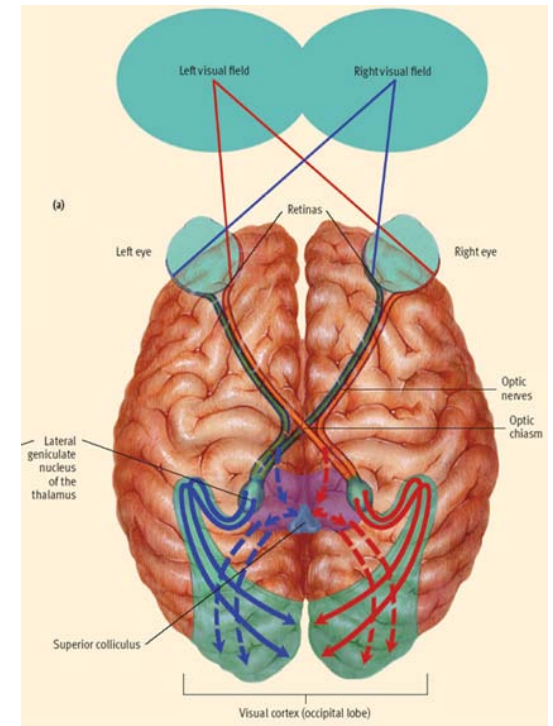
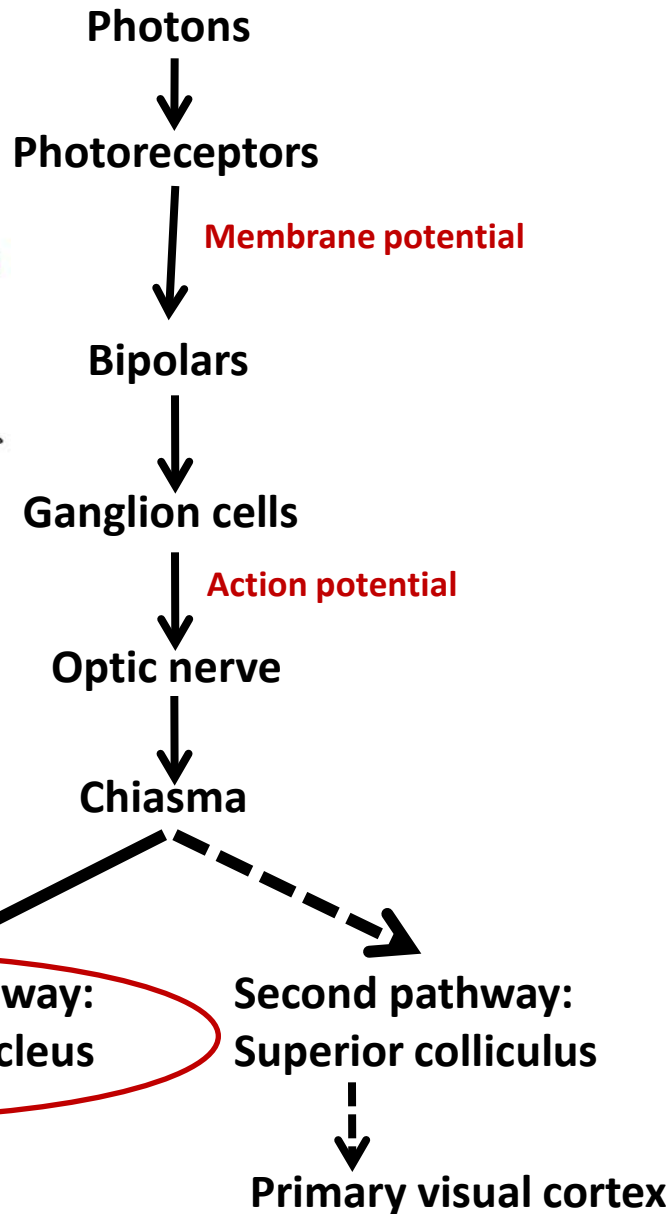
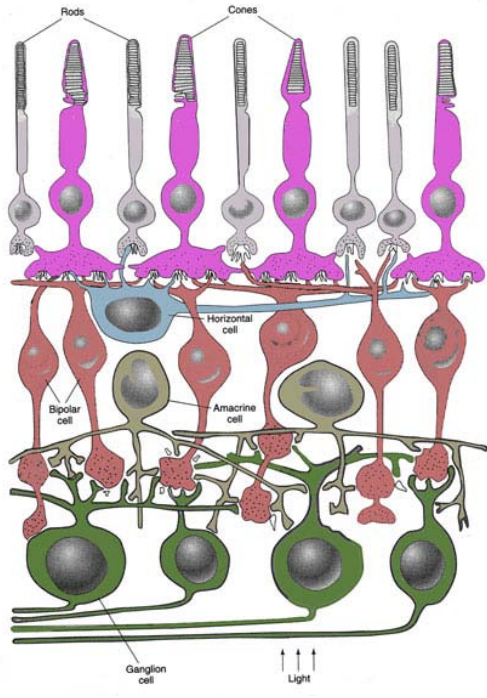
# The fovea is optimized for highest spatial resolution

„private line“ between cones,  
bipolars and ganglion cells

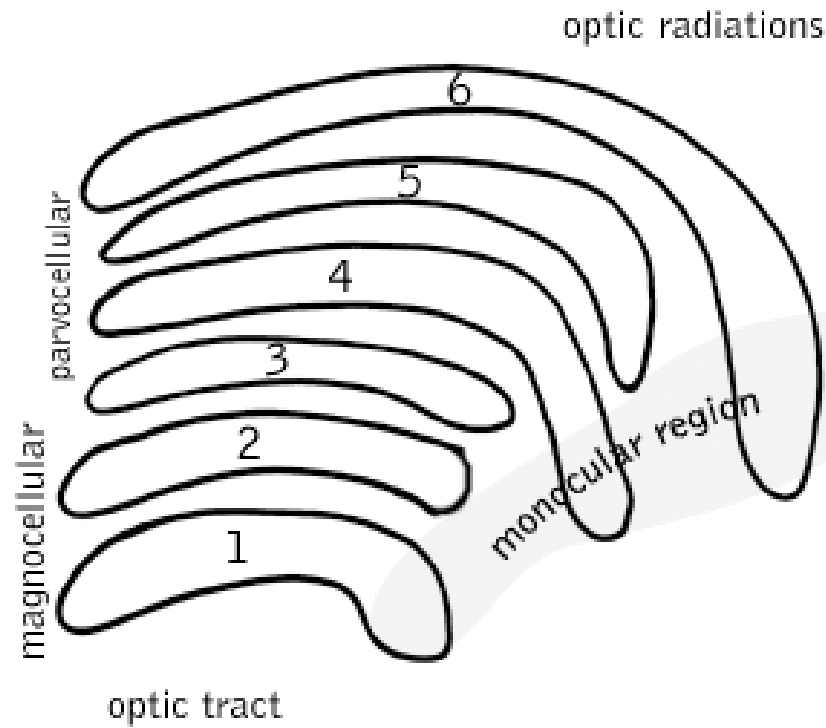


# **Color perception**

# The Retina and the Brain: Visual Information Processing



# The LGN is a distinctively layered structure





# Correspondence between ganglion cells and LGN cells

Retinal ganglion cells	LGN cells	Type of information
Parasol ganglion cells	Layers 1&2: Magnocellular cells	perception of form, <b>movement</b> , depth, and brightness
Midget ganglion cells	Layers 3-6: Parvocellular cells	perception of <b>color</b>
Small bistratified ganglion cells	In between layers 1-6: Koniocellular cells	perception of <b>color</b>

# Interactive Stroop Effect Experiment

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Red	Green	Blue	Yellow	Pink
Orange	Blue	Green	Brown	Black
Green	Yellow	Pink	Red	Orange
Brown	Red	Black	Blue	Yellow
Black	Orange	Green	Brown	Red

# Interactive Stroop Effect Experiment

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Blue

Pink

Black

Red

Brown

Brown

Red

Blue

Green

Orange

Yellow

Blue

Red

Orange

Black

Brown

Red

Green

Black

Red

Red

Pink

Blue

Green

Black

**So, what does this all mean?**

It means that color perception is relative and not absolute. And, since color perception is relative, we are always subject to these effects. In other words, it's in our mind not our eye.