

# Fundus Autofluorescence

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## Fundus Autofluorescence

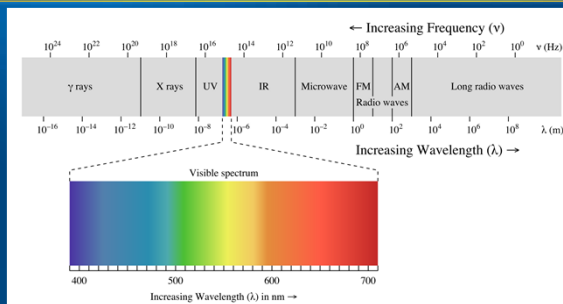
*"Fundus autofluorescence imaging is used to record fluorescence that may occur naturally in ocular structures or as a byproduct of a disease process. This technique allows the topographic mapping of lipofuscin distribution in the RPE. The intensity portrayed by FAF corresponds to the accumulation of lipofuscin, which increases with aging, RPE cell dysfunction or an abnormal metabolic load on the RPE."*

*-Saudi Journal of Ophthalmology*

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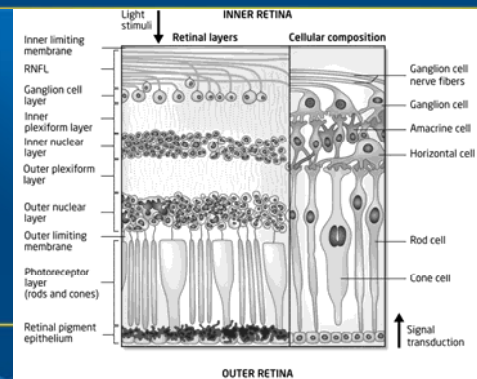
## Visible Spectrum



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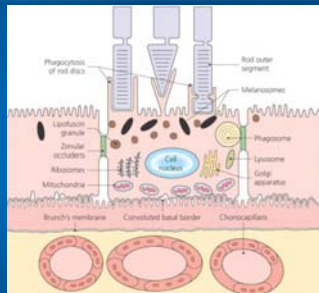
## The RPE



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## The RPE

- Retinal Pigment Epithelium – single layer of cells which separates the choroid from the neurosensory retina
- Five main roles, one of which is phagocytosis of the outer segment
  - Allows renewal process necessary to maintain photoreceptors (rods and cones) functioning normally

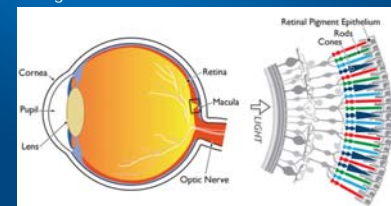


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## The RPE

- Aging can result in incomplete or partial breakdown of these segments which causes an accumulation of lipofuscin.
- Excessive lipofuscin is associated with visual loss.
- This accumulation is evident in ocular disease even before the visual cycle begins to degrade.

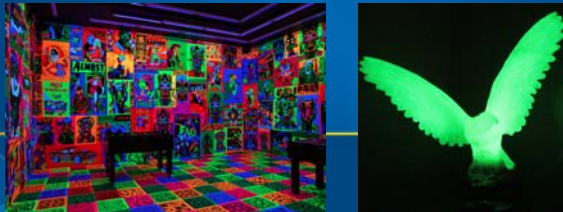


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## Fluorescence and Phosphorescence

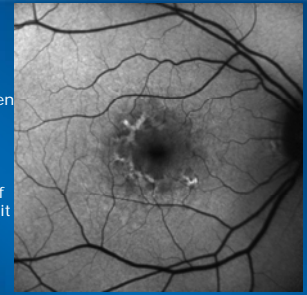
- Fluorescence – emission of light by a substance that has absorbed light.
  - Immediate release of light.
  - emitted light has longer wavelength
  - electron excited to a higher energy state and relaxes to ground state
- Phosphorescence – emission of light lasts after excitation has stopped
  - electrons cannot relax into ground state therefore emission lasts after excitation



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## FLAUM EYE INSTITUTE Fundus Autofluorescence

- Autofluorescence – fluorescence occurs naturally, without a dye
- Fluorescence occurs when molecules get excited by a specific light and then emit light at a longer wavelength which makes them glow
- Fundus autofluorescence works by shining a specific color/wavelength of light into the eye and capturing how it naturally fluoresces without dye
- The major source of fundus autofluorescence is this lipofuscin.

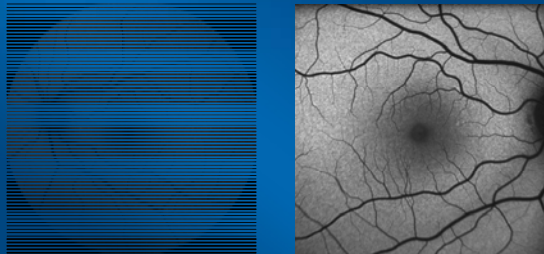


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## FLAUM EYE INSTITUTE Healthy Eyes



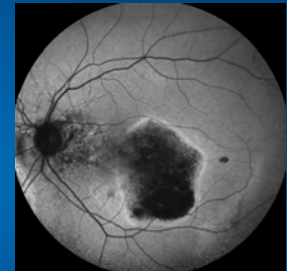
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## FLAUM EYE INSTITUTE Lipofuscin and FAF

- The RPE ages or becomes more damaged which means there is more lipofuscin
- Lipofuscin properties fluoresces at about 500nm to 750nm
- Intensity of signal emitted relates to the health level of the RPE based upon what is normal.
  - Hyperfluorescence (white/really bright areas) means that the RPE cells are failing
  - Hypofluorescence (black/really dark areas) means that the RPE is dead/ atrophy



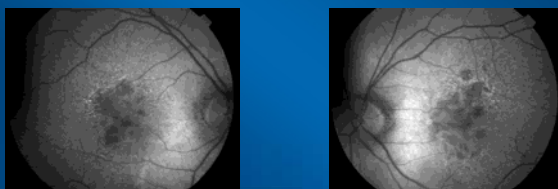
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## FLAUM EYE INSTITUTE Types – Fundus Camera

- Fundus camera uses white light
- Wideband exciter filter and wideband barrier filter
- Only a single image
- Limiting factors such as ocular media opacities
- The first fundus camera technique used filters around 580nm for the exciter and a barrier filter centered at 695nm.

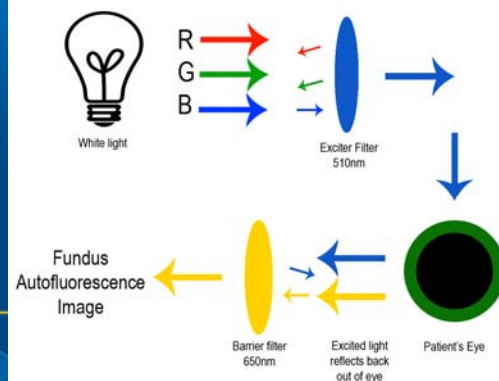


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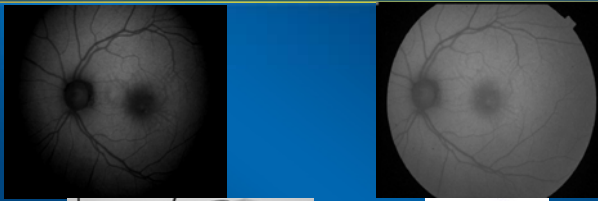


## FLAUM EYE INSTITUTE Fundus Camera Light



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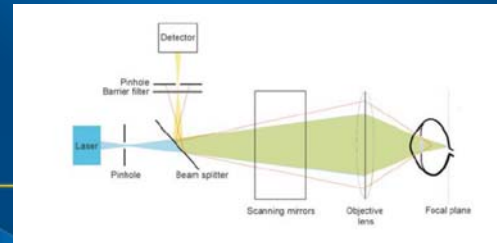
## FLAUM EYE INSTITUTE Types – Fundus Camera



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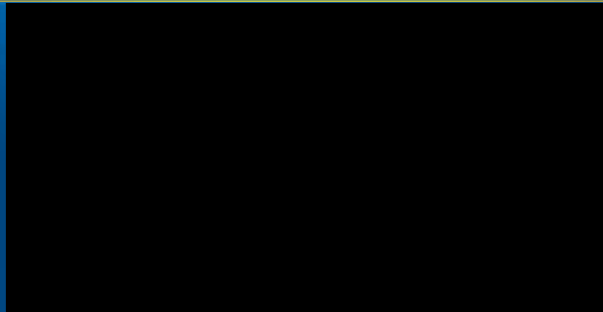
## FLAUM EYE INSTITUTE Types - cSLO

- Confocal Scanning Laser Ophthalmoscope
- Laser beam of specific wavelength on the retina to excite it
- Images “built” to reduce background noise
  - Acquire images at nearly a video rate and add the frames together
  - Provides higher-quality if there is minimal movement by patient.
- Two main types Heidelberg Retina Angiograph and the Optos



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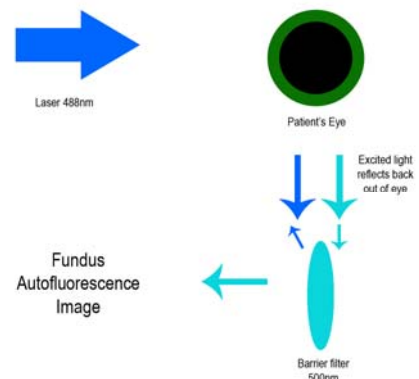
## FLAUM EYE INSTITUTE Types - cSLO



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<http://www.heidelbergengineering.com/us/products/spectralsis-models/tel@URMEDICINEocal-scanning-laser-ophthalmoscopy/>



## FLAUM EYE INSTITUTE cSLO Light



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## FLAUM EYE INSTITUTE Heidelberg Spectralis HRA

- Heidelberg Spectralis HRA
  - Fundus Autofluorescence imaging
    - Blue laser 488nm and 500nm barrier filter
- More than just FAF Images!
  - Indocyanine green angiography / Infrared Autofluorescence
    - Diode laser at 790nm and a barrier filter at 830nm
  - Infrared reflectance imaging
    - Second diode laser at 820nm
  - Blue or “red-free” reflectance imaging
    - Blue laser 488nm and no barrier filter
  - Fluorescein Angiography
    - Same filters as FAF, only difference is that dye is introduced

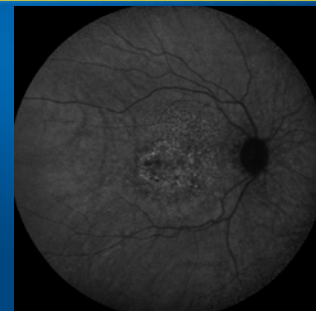


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## FLAUM EYE INSTITUTE ICGA/IRFAF



Laser at 790nm  
Barrier filter at 830nm

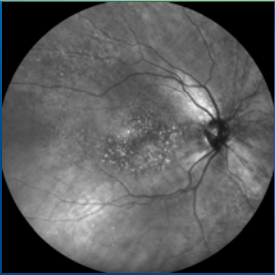
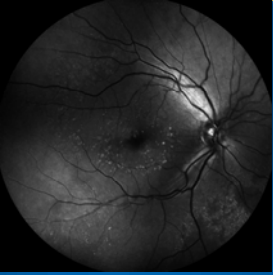
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## Infrared and Blue Reflectance





Infrared Reflectance  
Laser at 820nm

Red-Free Blue Reflectance  
Laser at 488nm

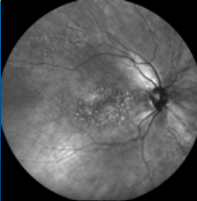
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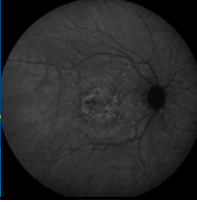


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## Options



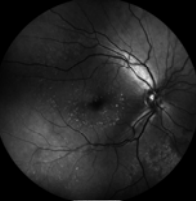
Infrared



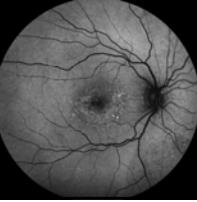
Red-Free



ICGA/IRAF




FAF



FAF

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## Optos Ultra-widefield

- Optos California Ultra-Widefield
- Fundus Autofluorescence
  - Green laser at 532nm
- Other imaging modalities:
  - Indocyanine Green Angiography
    - Infra-red 802nm
  - Red-Free
    - Green laser at 532nm
  - Fluorescein Angiography
    - Blue laser at 488nm



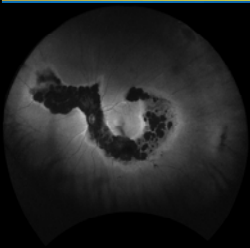
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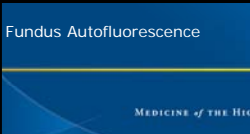


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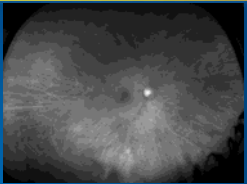
## Options




Choroidal/ICG



Fundus Autofluorescence




Red- Free



Red- Free

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## Examples



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## Dry AMD

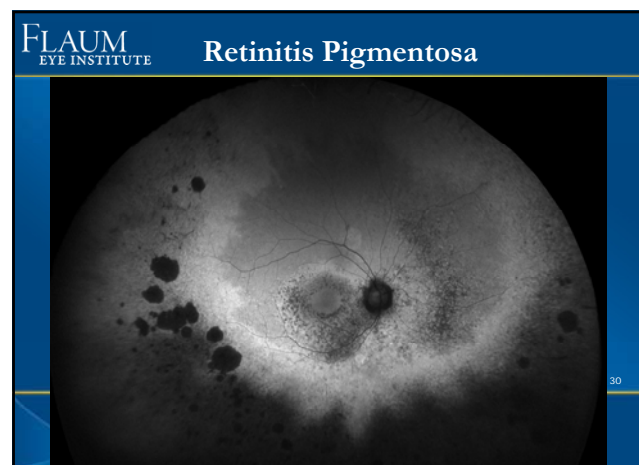
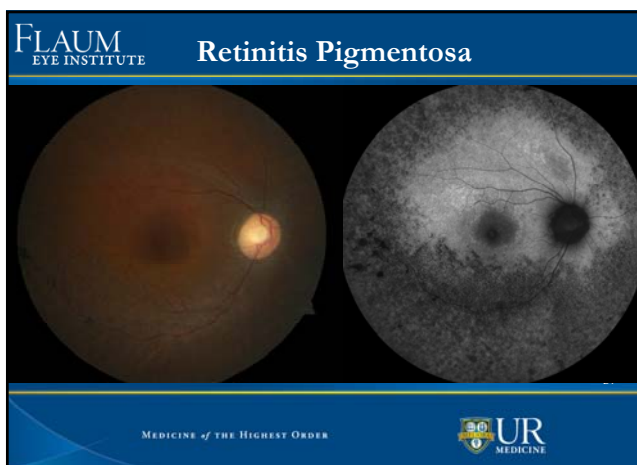
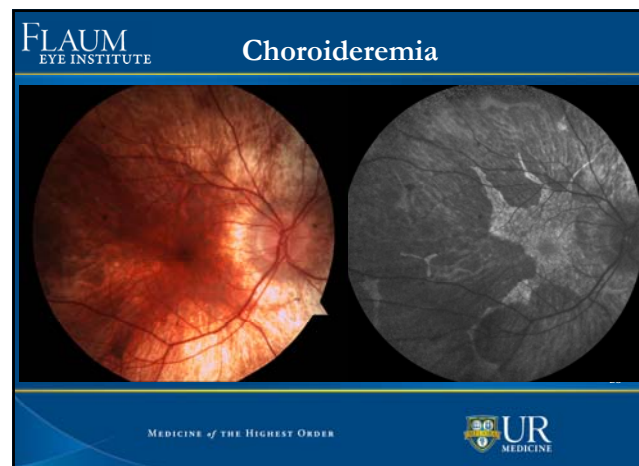
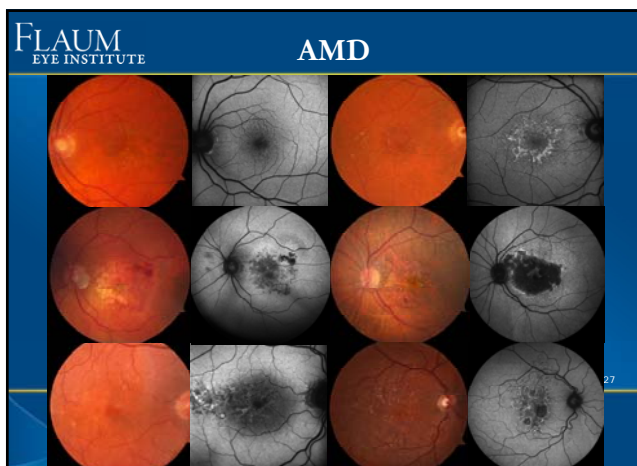
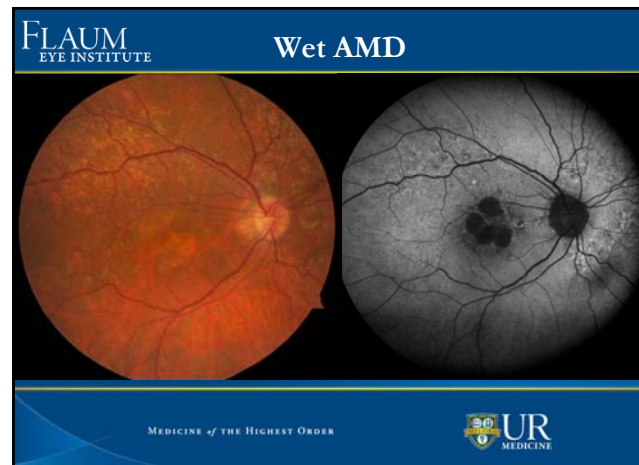
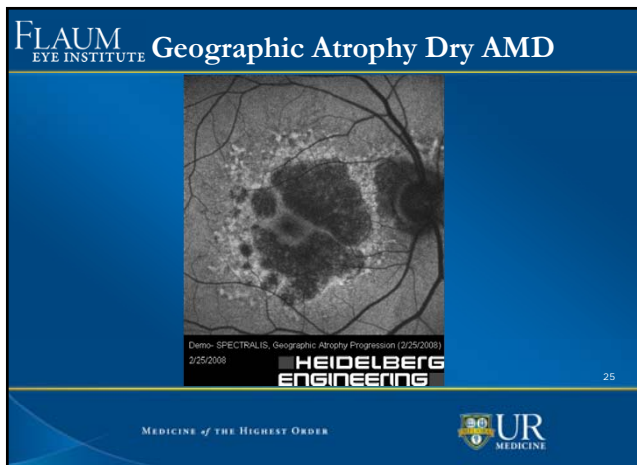



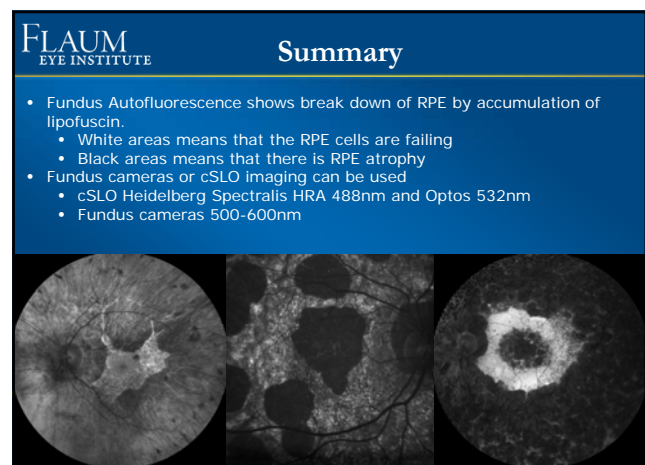
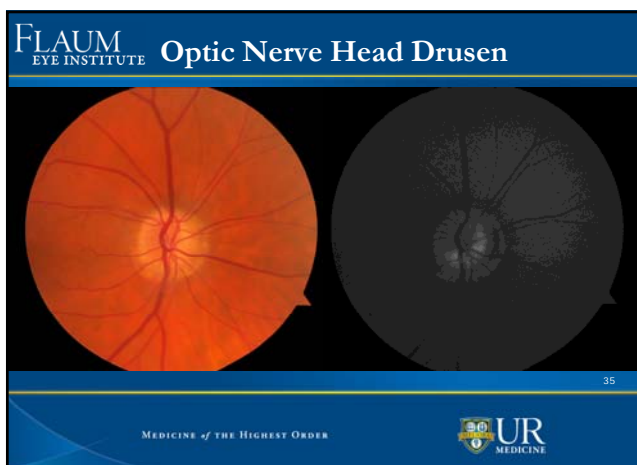
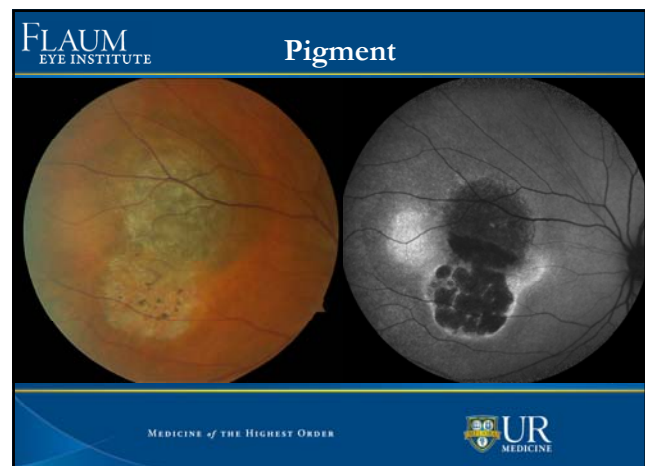
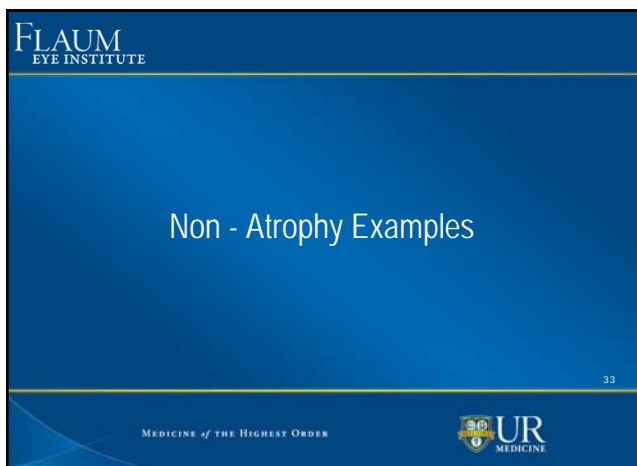
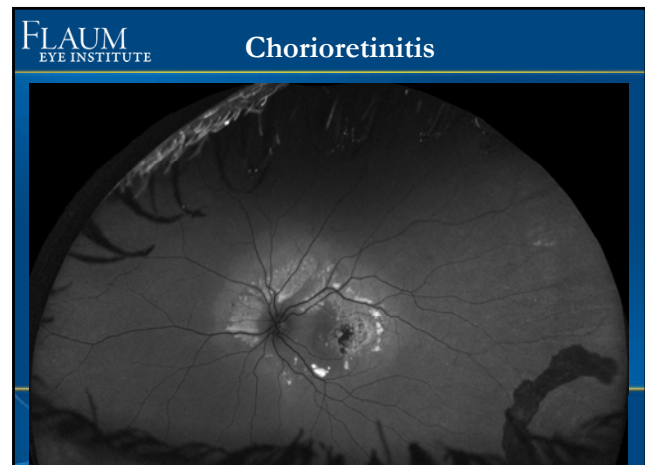
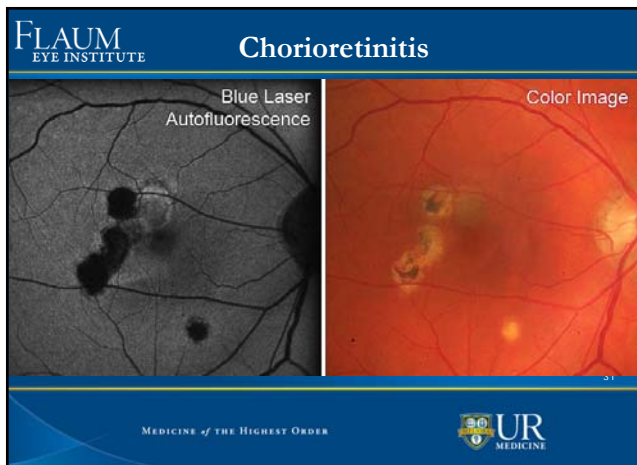
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**Credit**

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  - <http://www.comtecmed.com/cophy/2011/Uploads/assets/von%20strachwitz.pdf>

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- Diagrams
  - *Clinical Atlas of Ophthalmology*, Spalton
  - <http://www.closerlookatstemcells.org/stem-cells-and-medicine/macular-degeneration>
  - <https://en.wikipedia.org/wiki/Light>
  - <http://www.neurology.org/content/80/1/47/F1.large.jpg>
- Images courtesy of Flaum Eye Institute Diagnostic Team and Imagers
  - Brittany Bateman
  - Brittany Richardson, COA OCT-C CRA
  - Kassandra Mundt
  - Patricia Artman, COA
  - Rachel Hollar, OCT-C CRA
  - Taylor Pannell, OCT-C CRA

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**Thank you**

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