



KING'S COLLEGE LONDON
OPHTHALMOLOGY SOCIETY

Teaching Day March 2013

Gibran F Butt MSc

- 0) Introduction
- 1) History
- 2) Ocular cranial nerves
- 3) Fundoscopy

Intro...



WHAT DO THEY WANT!!!!!!



Can you get enough information?

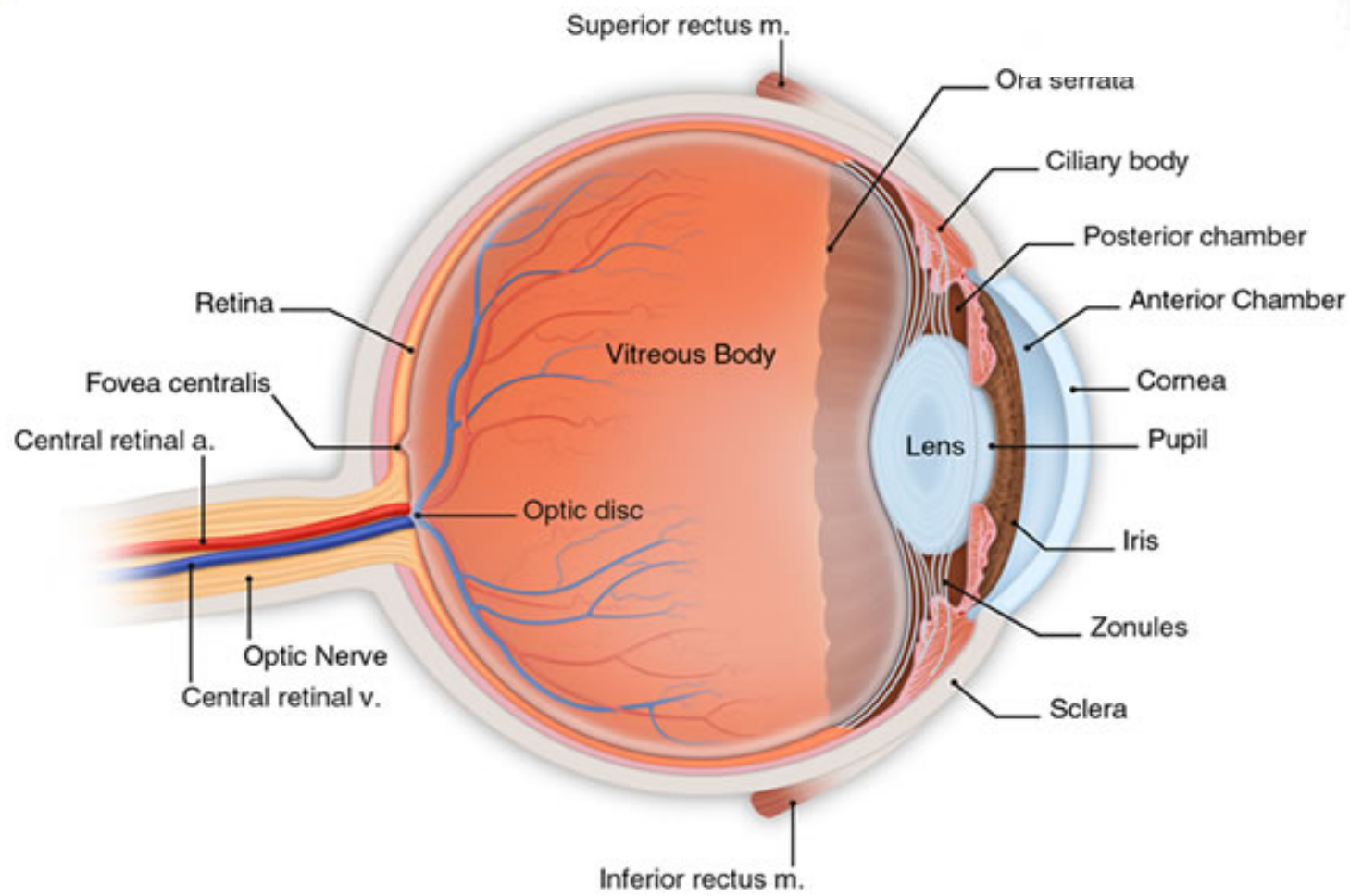
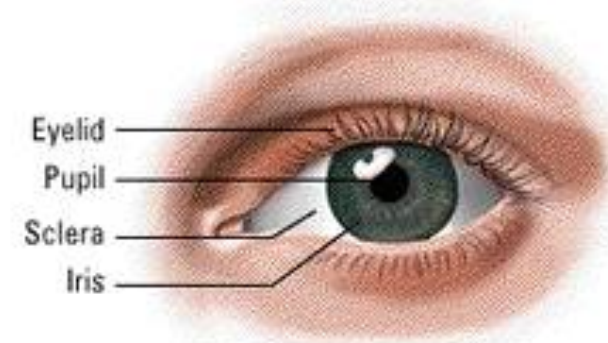
Are you nice?



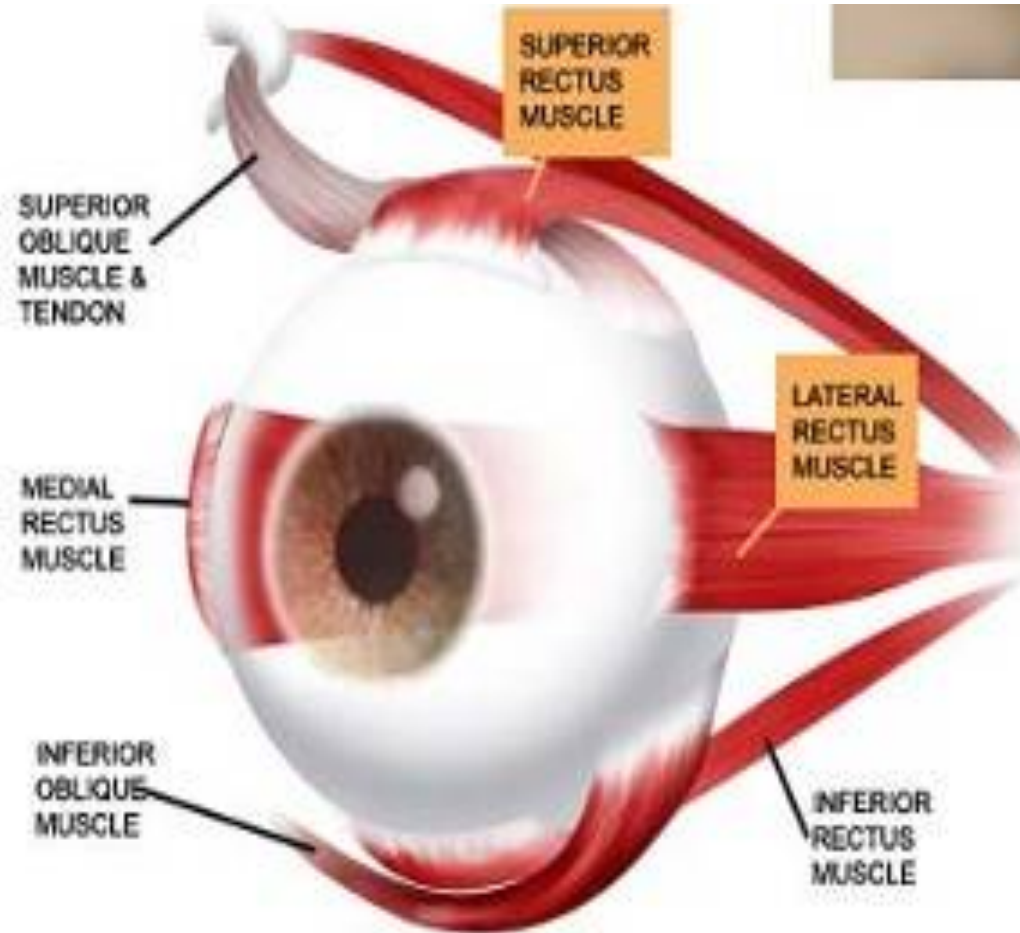
Candidate Num and Name.					
	CLEAR PASS	PASS	BORDERLINE	FAIL	CLEAR FAIL
KNOWLEDGE (Interpretation of findings)	<ul style="list-style-type: none"> Highlights appropriate positive & negative findings Understands implications & significance of findings for patient Correct differentials Good clinical reasoning 	<ul style="list-style-type: none"> Highlights some positive & negative findings Understands implications of findings (interprets findings to clarify diagnosis / management) Correct differentials Some clinical reasoning 	<ul style="list-style-type: none"> Highlights positive findings but not negative findings Does not adequately eliminate alternative diagnosis Incorrect differentials or Errors in clinical reasoning 	<ul style="list-style-type: none"> Going through the motions - does not highlight relevant positive or negative findings Examination cannot eliminate alternative diagnosis Incorrect differentials Poor / wrong clinical reasoning 	<ul style="list-style-type: none"> Highlights irrelevant findings; lacks expected knowledge Examination cannot eliminate alternative diagnosis Incorrect differentials No clinical reasoning
SKILLS (Performance of examination)	<ul style="list-style-type: none"> Well-structured / systematic Fluent Correctly identifies all clinical signs 	<ul style="list-style-type: none"> Good structure A little rushed / slowed but overall acceptable Correctly identifies most of the clinical signs 	<ul style="list-style-type: none"> Erratic structure Hurried approach that hinders performance Misses important/ too many minor signs 	<ul style="list-style-type: none"> Poor structure Slow uncertain approach Misses important signs / Describes non-existent findings 	<ul style="list-style-type: none"> No structure / system Marked hesitancy / haste Describes incorrect / non-existent signs with confidence
ATTITUDE (Interaction with patient)	<ul style="list-style-type: none"> Full greeting & introduction Explains examination fully and asks permission Mindful of rapport, patients comfort & dignity Patient feels at ease 	<ul style="list-style-type: none"> Good greeting & introduction Explains examination Some thought to rapport, patients comfort & dignity Patient feels some ease 	<ul style="list-style-type: none"> Incomplete greeting & introduction Some explanation Inconsistent rapport & attentions to patient's comfort & dignity Patient not at ease 	<ul style="list-style-type: none"> Poor approach on a number of levels No explanation No attention to patient comfort & dignity Patient feels uncomfortable 	<ul style="list-style-type: none"> Lack of interaction Brusque approach Candidate is flustered and halts performance Lack of civility / inappropriate comments inc. flippancy

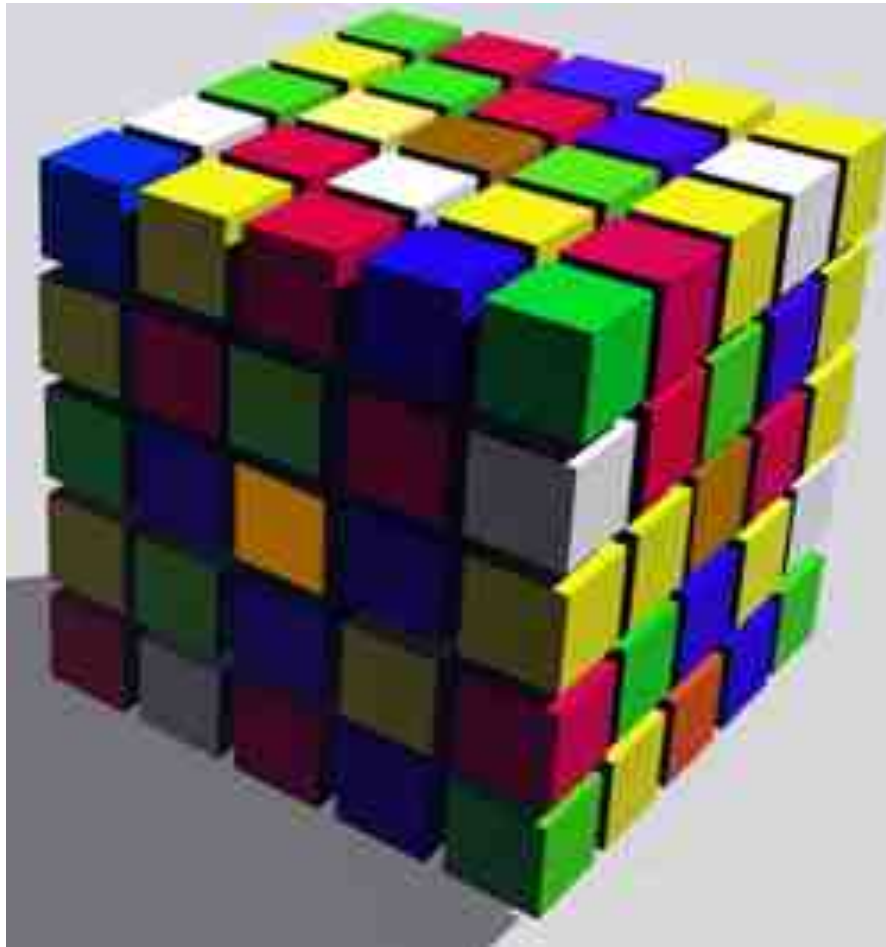
What could the candidate do to improve their performance?

Anatomy recap



Anatomy Recap





1) Eye History



- **PC/HPC:**

UNI/BI-lateral

Red eye

Pain

Visual disturbance/loss

- **Ocular hx**

Refractive error/
contact lens/ trauma/
foreign bodies

- **PMH**

C/V risk factors

Surgical history

- **FH**

Visual conditions
(retinitis pigmentosa)

Mypopia

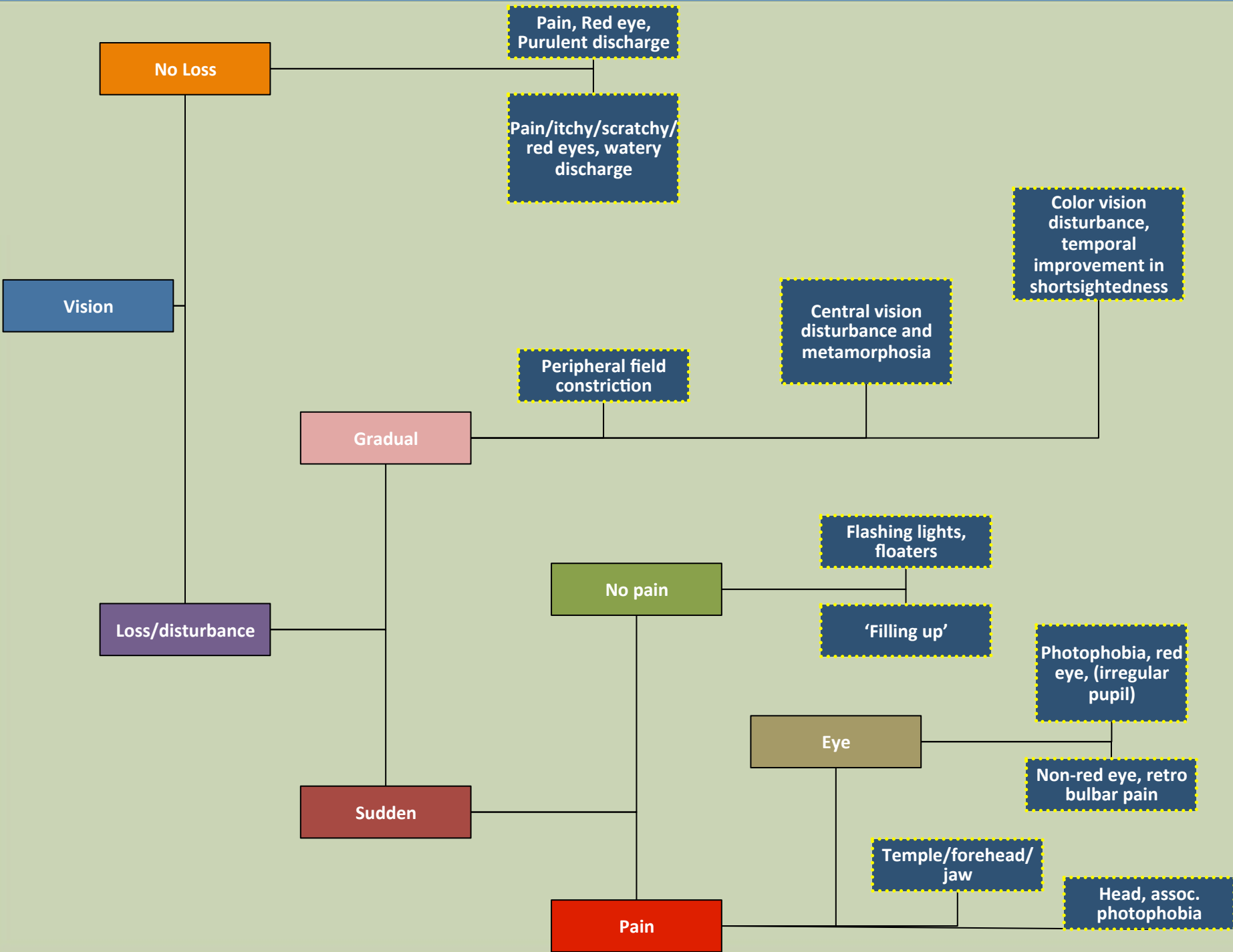
- **SH**

- **DH**

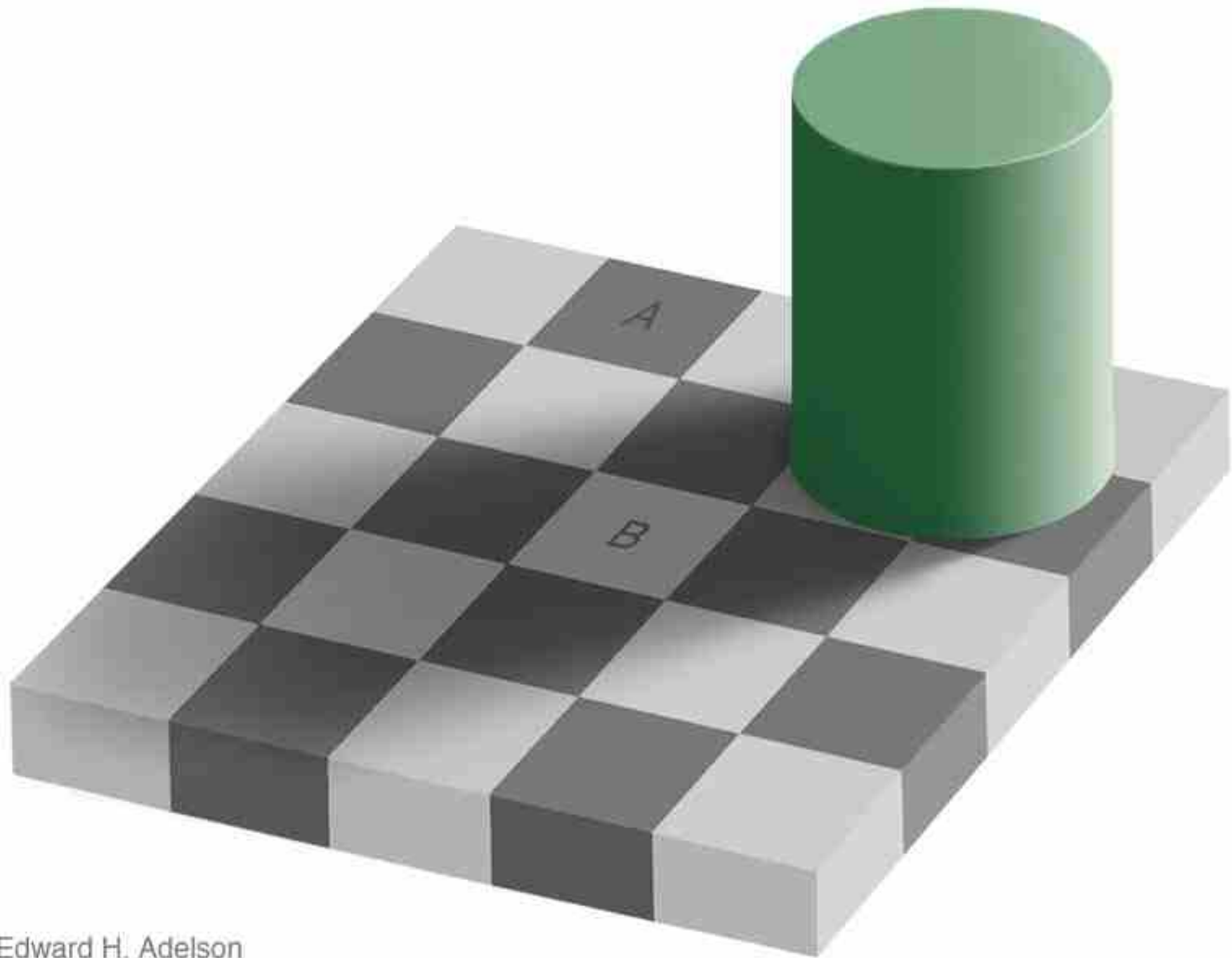
Classic histories



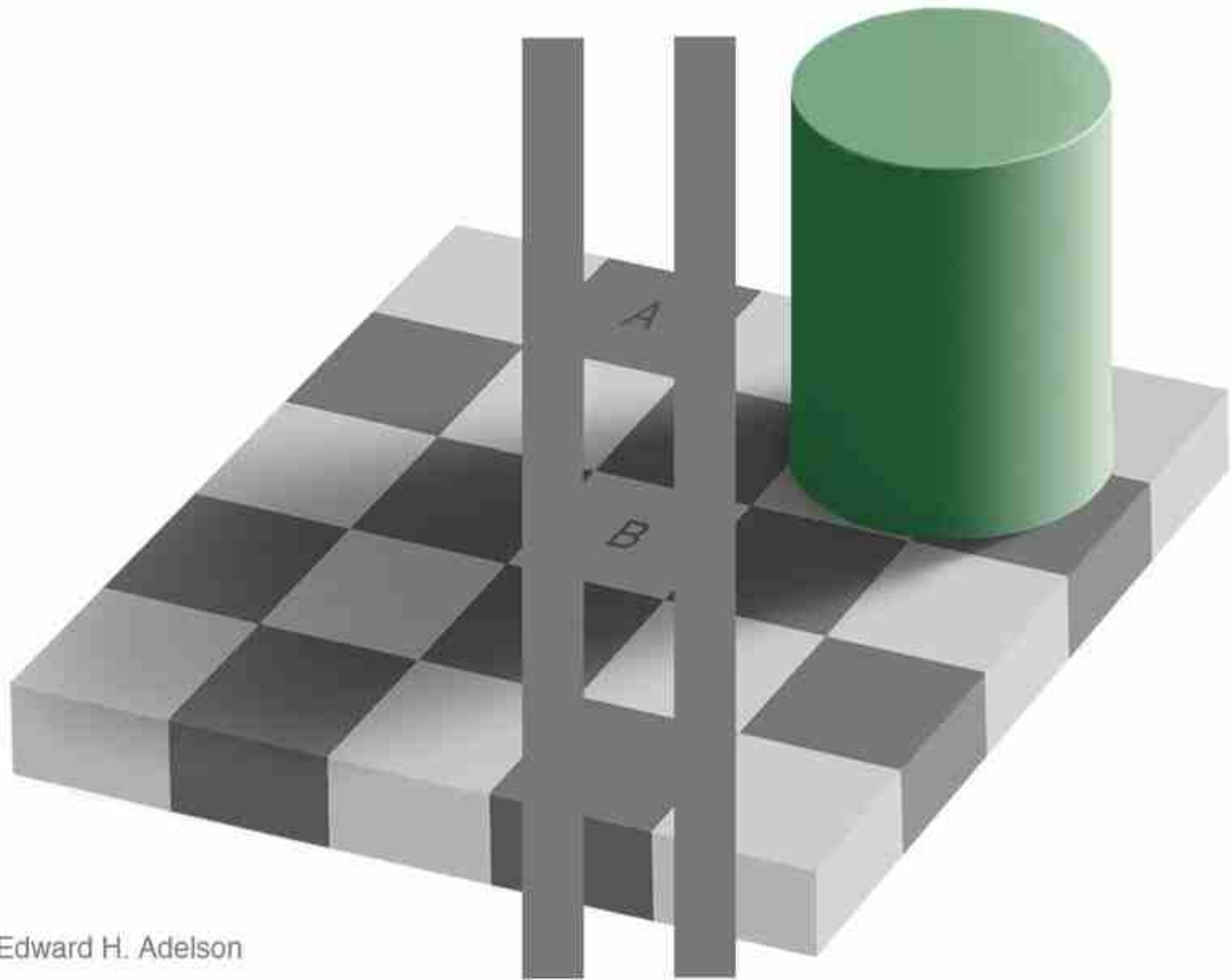
- Keratitis (bacterial/viral/atopic)
- Conjunctivitis
- MS
- Glaucoma (acute or chronic)
- Giant cell arteritis
- Cataract
- Age-related Macular Degeneration



- Diabetes
- HTN
- Previous focal neurology
- Polymyalgia rheumatica
- Atopy



Edward H. Adelson



2) Ocular Cranial Nerves



- CN: 2,3,4 & 6
- 2.1) CN2: Visual acuity (+ defects)
- 2.2) CN2/visual system: fields (+ defects)
- 2.3) CN2: Reflexes (+ defects)
- 2.4) CN3,4,6: Movement (+ defects)
- 2.5) To complete the examination I would like to perform direct ophthalmoscopy, take a full history, examine intraocular pressure....

2.1) Visual acuity: far vs. near



Distance between the patient and the eye chart

Distance at which the letter can be read by a person with normal acuity

- Ask what their vision is like (need to assess based on hand movements, number of fingers or snellen?)
- Ask about glasses
- Place patient at 20 ft from Snellen chart
- OD then OS
- VA is line in which $> \frac{1}{2}$ letters are read

??? 20/28 +2 ???

OD - ocular dexter, OS - ocular sinister

E

1 20/200

F P

2 20/100

T O Z

3 20/70

L P E D

4 20/50

P E C F D

5 20/40

E D F C Z P

6 20/30

F E L O P Z D

7 20/25

D E F P O T E C

8 20/20

L E F O D P C T

9

F D P L T C E O

10

P E Z O L C F T D

11

2.1) Visual acuity: near



- Test of **function**
- (may get away with mentioning it)
- Use a card if available (at ~40 cm)
- Record (or at least remember!)

Near Visual Acuity Test						Chart 1			
MODIFIED ETDRS WITH SLOAN LETTERS						Snellen Distance Equivalent			
For testing at 40 cm (16 inches)						Diopters of Add for 1 M			
Letter Size (metric)						at 40 cm	at 20 cm	at 40 cm	at 20 cm
8.0 MM	N	C	K	Z	O	20/400	20D	20/800	40D
6.4 MM	R	H	S	D	K	20/300	15D	20/600	30D
5.0 MM	D	O	V	H	R	20/250	12D	20/500	25D
4.0 MM	C	Z	R	H	S	20/200	10D	20/400	20D
3.2 MM	O	N	H	R	C	20/150	8D	20/300	15D
2.5 MM	D	K	S	N	V	20/125	6D	20/250	12D
2.0 MM	Z	S	O	K	N	20/100	5D	20/200	10D
1.6 MM	C	K	D	N	R	20/80	4D	20/160	8D
1.25 MM	S	R	Z	K	D	20/60	3D	20/125	6D
1.0 MM	H	Z	O	V	C	20/50	2.5D	20/100	5D
.8 MM	N	C	D	O	K	20/40		20/80	4D
.6 MM	V	H	C	R	O	20/30		20/60	3D
.5 MM	S	R	Z	K	D	20/25		20/50	2.5D
.4 MM	O	N	H	R	C	20/20		20/40	
.3 MM	D	K	S	N	V	20/15		20/30	

Instructions: the 40cm test distance requires a maximum add of +2.50. If the patient cannot see the top line, move test distance to 20cm with a maximum add of +5.00. (Similarly if a 10cm test distance is required, the maximum add is +10.00)

Record test distance and letter size from the left column. Examples: 40/4M, 20/4M

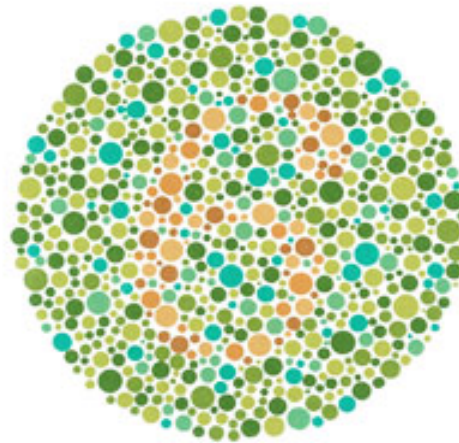
The columns on the right provide: reference to Snellen distance equivalent for two test distances; diopters of add for 1M print size for two test distances.

2.1) Ishihara plates

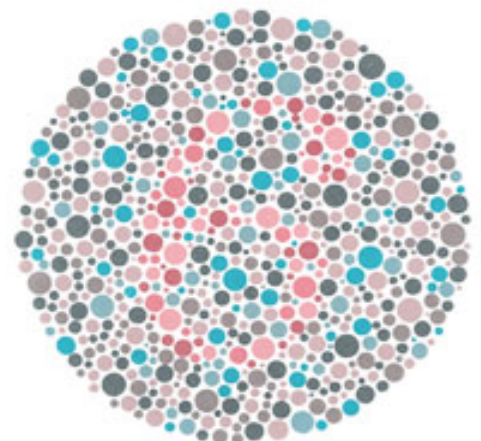
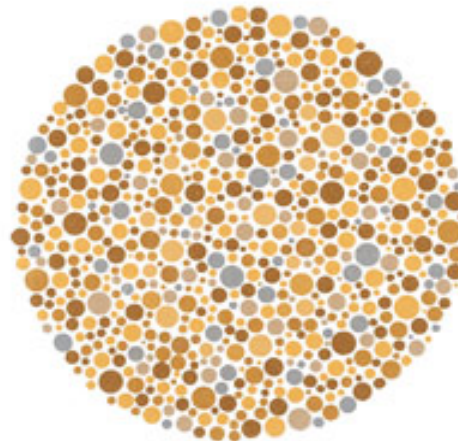
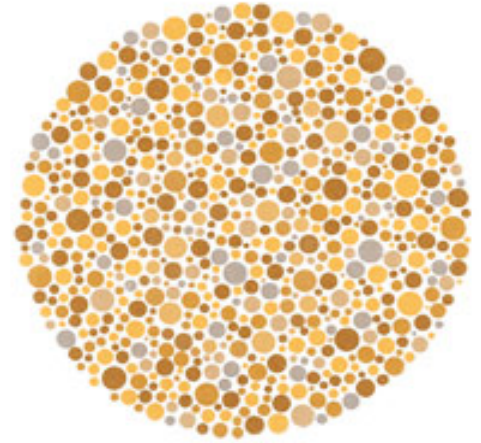
- Mention it – may be prompted to use the book and stopped and asked to explain use.

So:

- Check pt can read
- Open on control page
- Proceed with examination making sure to note which pages pt has difficulty with



Normal



2.2) Visual fields

Confrontation

- Same level, same eye.
- Red Pin to measure color saturation

Visual inattention

Blind spot defect

2.3) Reflexes: Pupillary Examination

- **Inspect**
- **Direct**
 - Constriction of ipsilateral eye
- **Consensual**
 - Constriction of contralateral eye
- **Swinging light test - RAPD**
- **Accommodation**
- **Defects:**
 - Fixed dilated – Tonic Holmes Aides pupil
 - Pinpoint – Horner's syndrome
 - RAPD – optic neuritis
 - AR Pupil

2.4) CN 3,4,6: Eye movements



A. Esotropia



B. Exotropia



C. Hypertropia



D. Hypotropia



- **Inspection:** Ptosis, squint, ask about double vision
- **Eye movements** – any double vision?
- Where is the (neuromuscular) lesion?

How to work out which muscle is affected



1. Which eye is affected
2. Which movement is affected
3. Which muscle(s) is responsible
4. Which nerve is responsible

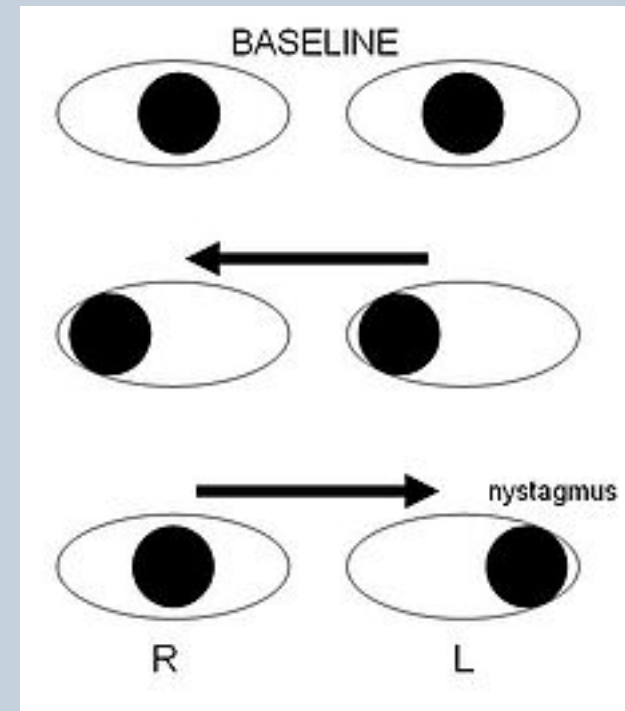
Important defect:

Internuclear ophthalmoplegia:-

Nystagmus and impaired abduction on lateral gaze.

Lesion is in MEDIAL LONGITUDINAL FASCICULUS on side of nystagmus

May be feature of Multiple Sclerosis



3) Fundoscopy



- Turn on the scope
- Set to 0
- Red reflex (leukocoria/darkened)
- Look in the eye
- Locate a blood vessel and follow back to the disc
- Inspect quadrants
- Inspect macula by getting them to look directly into the light

Fundus signs

- Gross observations
- Specific Structures:
 - Vessels: arteries and veins
 - ONH
 - Retina
 - Macula

Likely 3rd year Diagnoses



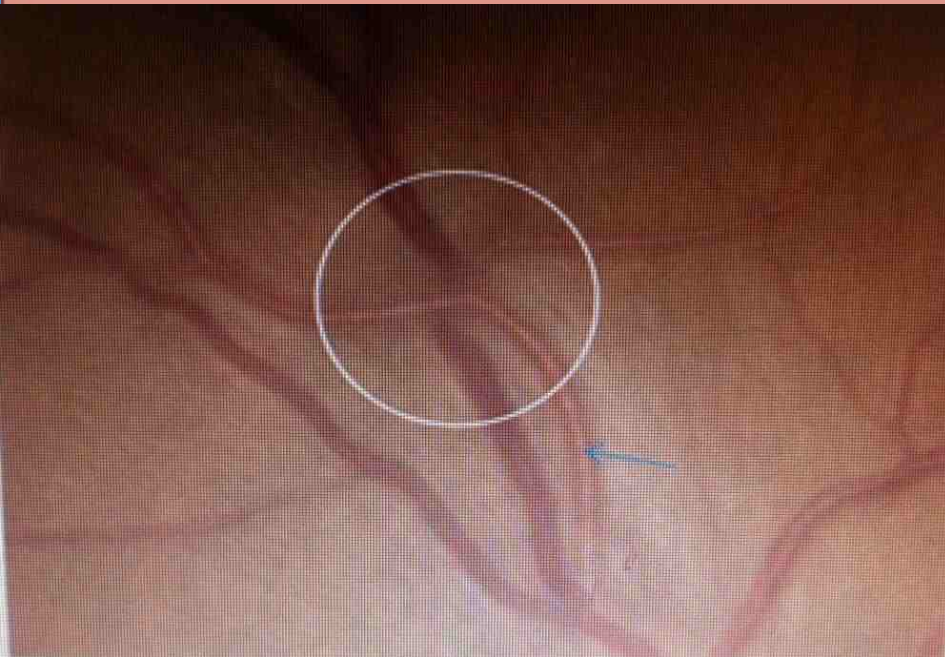
- Diabetic Retinopathy
- HTN retinopathy
- Macular Degeneration
- Retinitis pigmentosa

Others to consider?

- Macular hole
- Epiretinal membrane

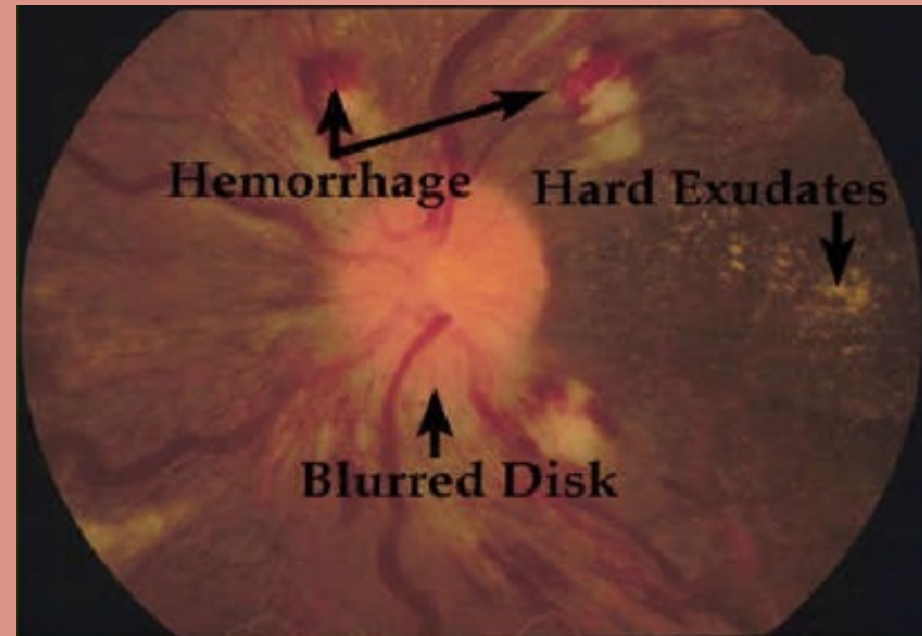
HTN

LOF



I) SILVER/COPPER WIRING

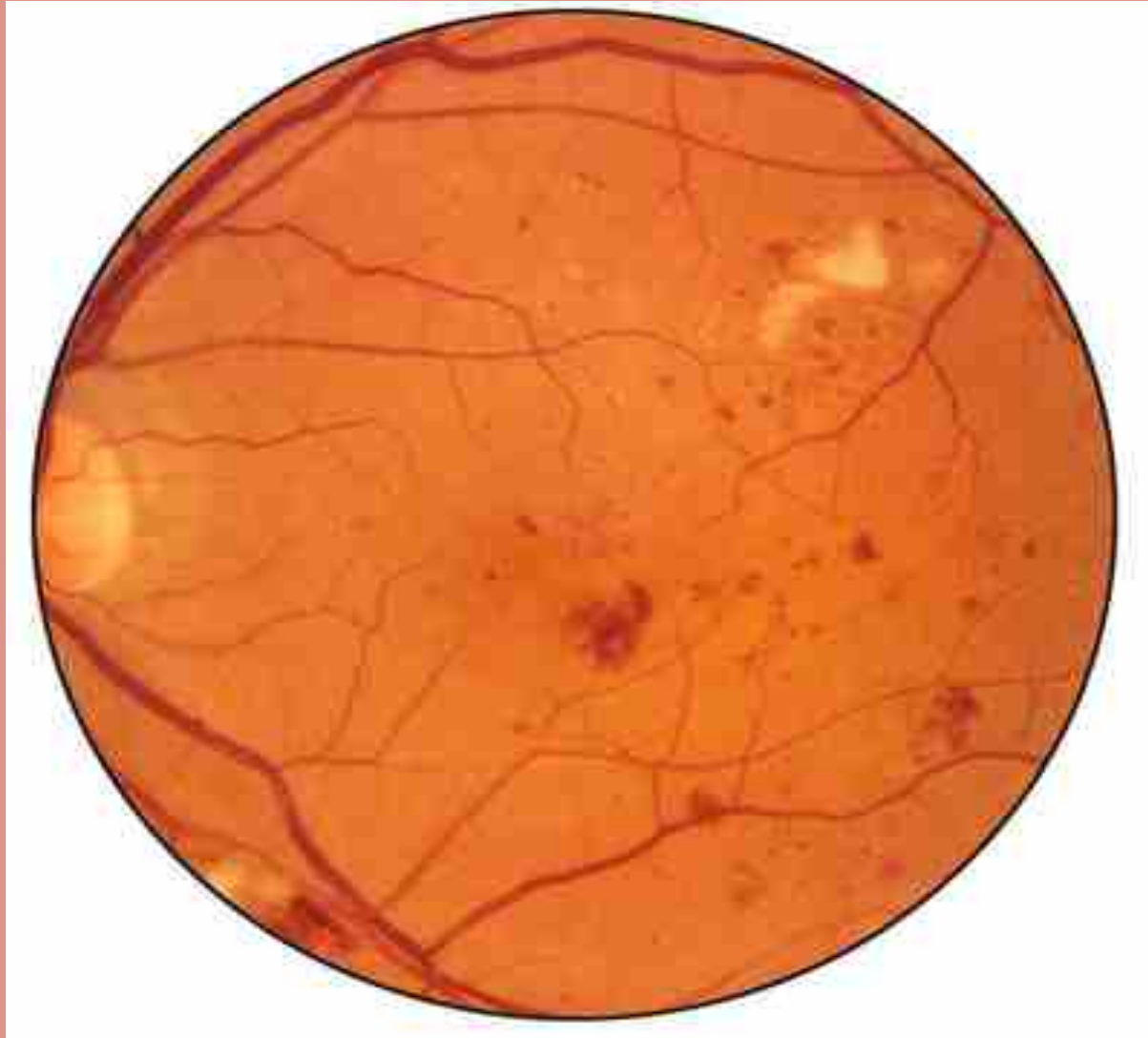
II) AV NIPPING



III) Cotton wool spots and flame hemorrhages

IV) PAPILLODEMA

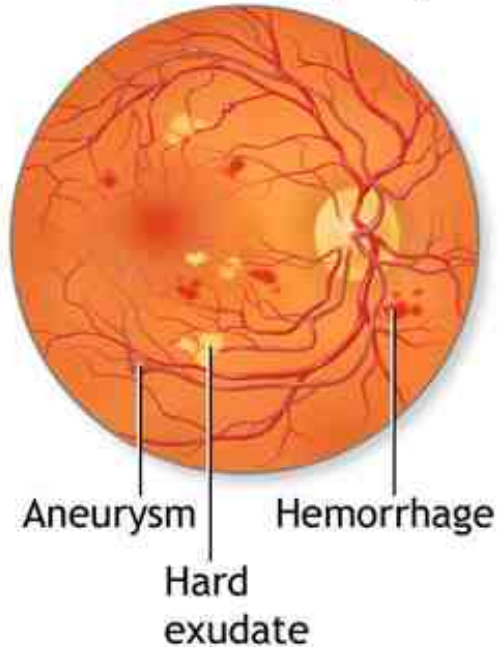
Diabetic retinopathy



Diabetic retinopathy



Non-proliferative
diabetic retinopathy



Proliferative
diabetic retinopathy



 ADAM.



Laser Photocoagulation

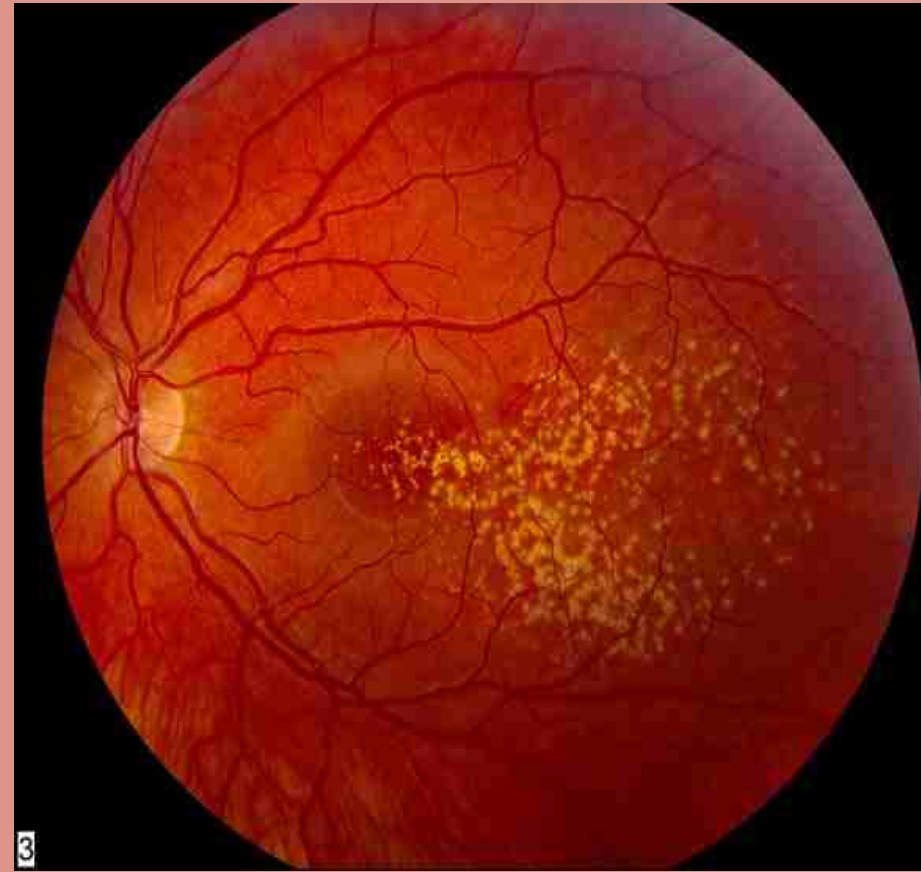
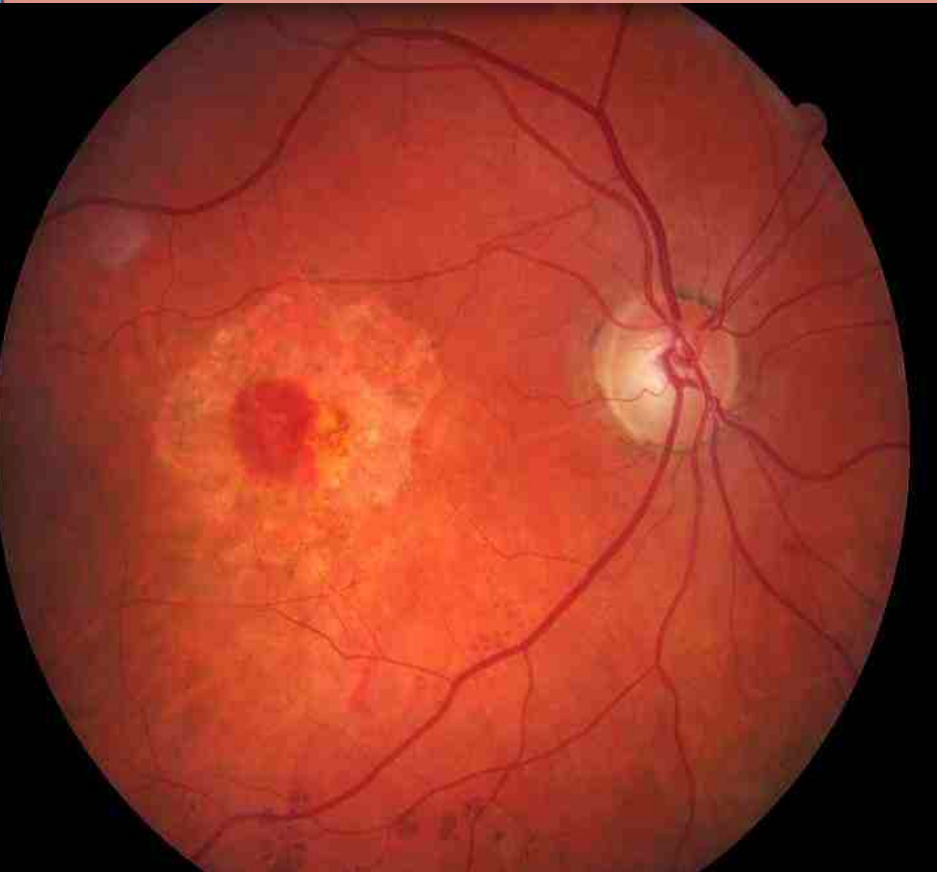
Macular Degeneration



Wet

V.S

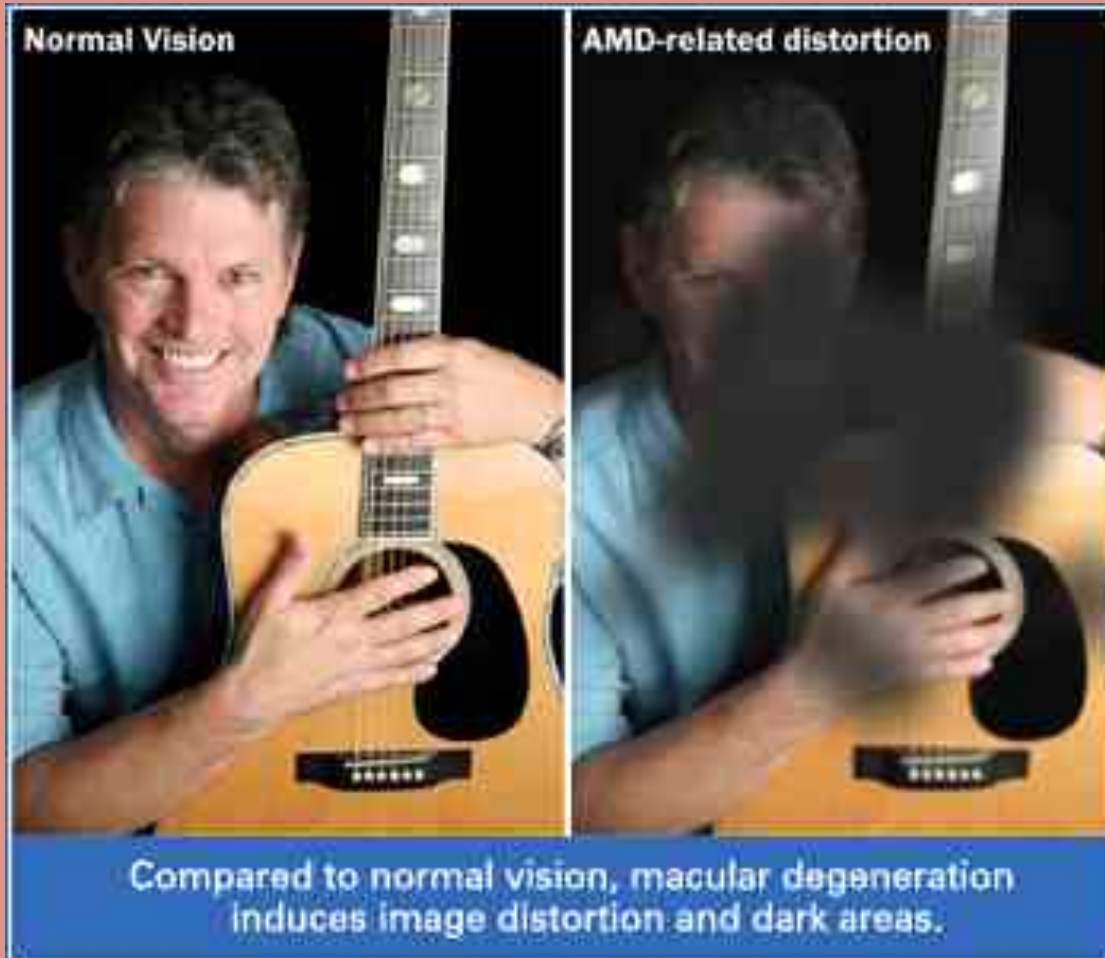
Dry



- Exudative changes/ Macular edema/ atrophy

- Soft exudates / Drusen

Macular Degeneration



Retinitis Pigmentosa



-Bone
Spicule
deposits

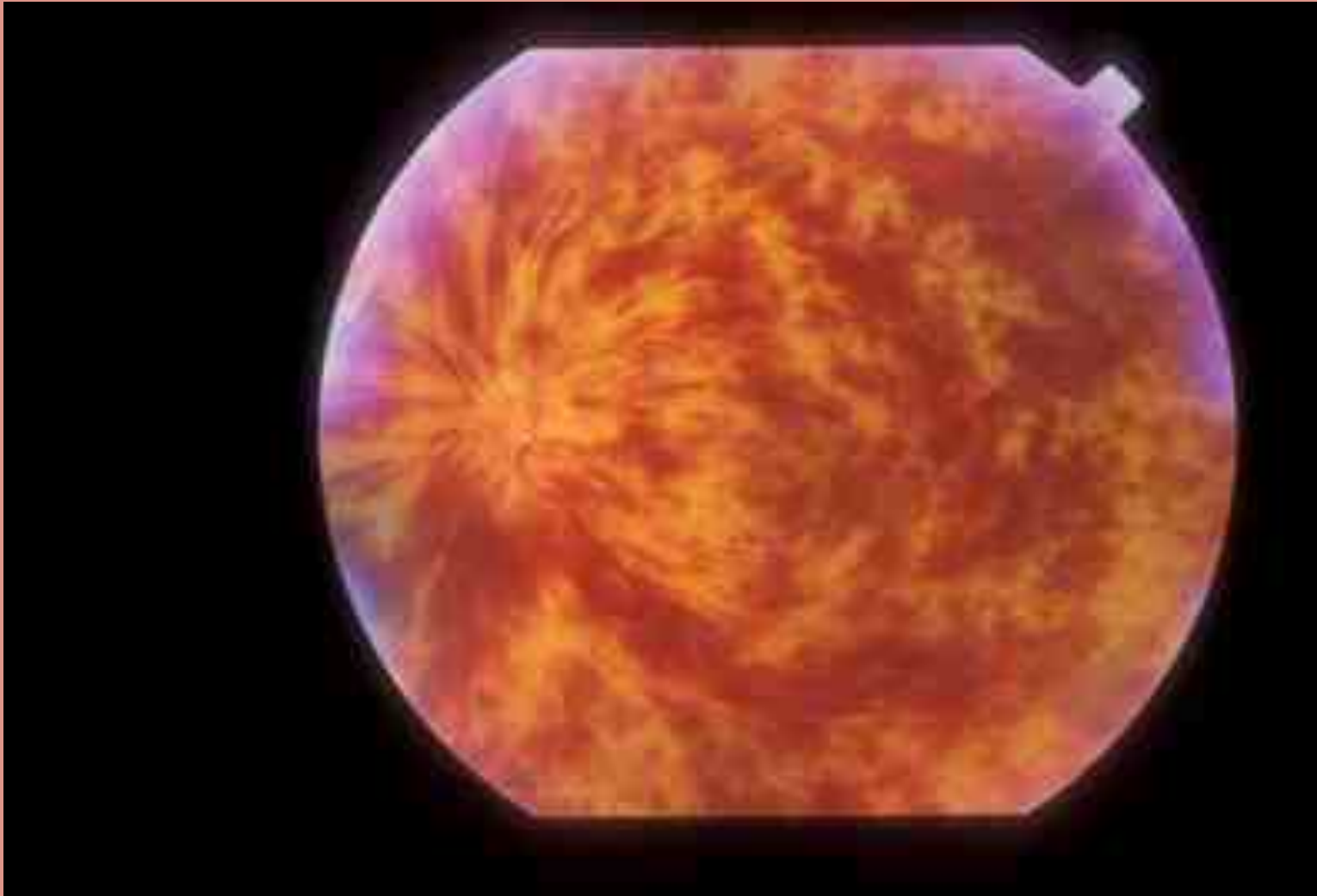
-Cherry red
Macula

-Waxy pale
optic disc

Chronic Glaucoma



Central Vein Occlusion



1. SENILE MACULA DEGENERATION. The disc appears normal but there is unusual pigmentation at the macula. The patient also has drusen which are asymptomatic nodules occurring in the choroid.
2. CENTRAL RETINAL VEIN OCCLUSION. Typical "stormy sunset" appearance with engorged veins with haemorrhages alongside them.
3. HYPERTENSIVE RETINOPATHY. The retinal arteries have become narrow and tortuous. In more advanced cases haemorrhages and "star burst" exudates occur together with papilloedema.
4. PAPILLOEDEMA. The disc is swollen and the disc margin has disappeared. The veins are congested.
5. DISC CUPPING. Here the degree of cupping is mild but suggestive of glaucoma. As the condition progresses the optic disc becomes pale and the cup wider and deeper.
6. OPTIC ATROPHY. The optic disc is pale and the condition is associated with gradual loss of vision. It may be secondary to a number of conditions including glaucoma, retinal damage, ischaemia and poisoning.
7. MILD BACKGROUND DIABETIC RETINOPATHY. Haemorrhages and microaneurysms can be seen.
8. BACKGROUND DIABETIC RETINOPATHY. There are areas of hard exudates and some evidence of macula involvement.
9. PREPROLIFERATIVE DIABETIC RETINOPATHY with haemorrhages, microaneurysms and hard and soft exudates.
10. PREPOLIFERATIVE DIABETIC RETINOPATHY with extensive haemorrhages and exudate formation and a preretinal (subhyaloid) haemorrhage.
11. PROLIFERATIVE DIABETIC RETINOPATHY with new vessel formation.
12. DIABETIC RETINOPATHY recently treated with laser photocoagulation.

Good Luck!!!!

- Sources:

Picture credits to google etc...

www.uninteforsight.com

www.labordeeyegroup.com

<http://www.health.state.mn.us/divs/fh/mch/webcourse/vision/mod4a.cfm>

<http://web1.ncoptometry.org/nonpro.aspx>