

# Five Ways an Ophthalmic Tech Can Save Someone's Life in Ophthalmology

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Start with a philosophical question & two really quick cases. Why are you here... because you believe as we all do that you can....?

# Five easy exam errors

Check pupil in light & dark (Don't use abbreviation "PERRLA": Pupils, equal, round, reactive, light & accommodation)

Letting only tech only check pupil

Not taking "Blurred disc margins" seriously Misusing the term "papilledema"

Writing "Dysconjugate gaze" or "EOMI"

Thinking "optic atrophy" is a diagnosis

The ways that an ophthalmic technician can save a life....

Avoid sole use of PERRLA for the pupil exam

Savino's rule - if there is a problem with the lid, motility, or pupil all three areas must be evaluated and documented

Don't use EOMI as the only assessment of motility

Don't order the same scan on all patients; and

Signs or symptoms are not diagnoses.



DRIVE THRU **LAWYER**



The big five

Refractive/cataract surgery (missed endophthalmitis )

Diabetic retinopathy

Glaucoma

Delayed diagnosis of brain tumor

Retinal detachment

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he plaintiff that she was not candidate for this procedure. Defendant's failure to do so was to be below the appropriate level of care.

atter: In my experience, a good attorney will sway the jury by arguing that the doctor could have done something more for the benefit of the patient. The jury will not believe that a physician would have done something deliberately and willfully harmful to the patient. During the course of the plaintiff's attorney will plant in the jury's mind a seed of doubt that, if the attorney had just been a little more aggressive, the plaintiff could have prevented the injury from suffering loss of vision and other damages.

(Courtesy of Ophthalmic Mutual Insurance Company)

Specialty	Average Settlement Amount (Approximate)
Cataract (43)	\$90,000
LASIK (26)	\$100,000
Retina (15)	\$120,000
Oculoplastics (13)	\$140,000
Glaucoma (9)	\$170,000
Neuro Opth (2)	\$310,000
Peds (10)	\$320,000
Cornea (5)	\$40,000
Fir Dx (10)	\$240,000
Gen (40)	\$160,000
Surgicenter (2)	\$180,000

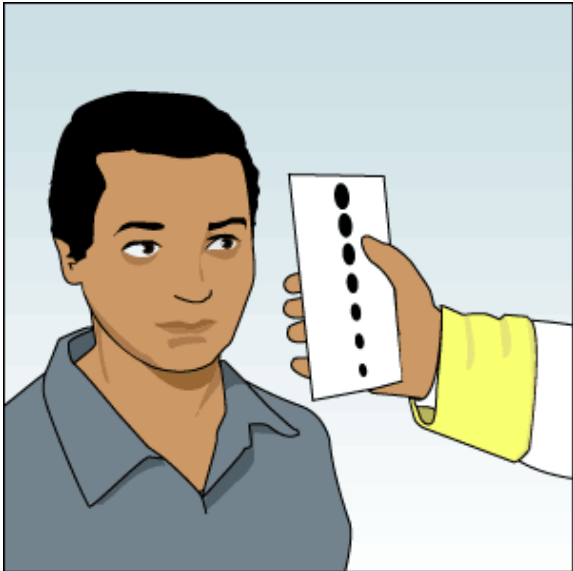
Figure 2. This chart shows OMIC's average malpractice settlement payment per specialty from 2001 to 2004.

# Here's a pearl don't use "PERRLA"

Pupils equal, round, reactive to light and accommodation (PERRLA)

Pupils can be equal, round & reactive to light and accommodation and have a HORNER syndrome

PERRLA only checks PNS pathway



Checking pupils in ambient light easily misses Horner syndrome (“PERRLA” ≠ NORMAL)





Apraclonidine test (inferior image) confirmed suspected diagnosis of Horner syndrome. González Martín -Moro et al. Horner Syndrome, a New Complication. J Oral Maxillofac Surg 2009.



LIGHT

DARK

AFTER APRACLONIDINE

BEFORE APRACLONIDINE

Horner syndrome

Anisocoria is greater in the dark

If pupil tested in light only then easy to miss subtle anisocoria:  
PERRLA can miss Horner pupil

Ptosis is always mild in HS (12% no ptosis)

Book Horner does not look like real world sometime



**ANISOCORIA IS LESS IN THE LIGHT!**

# How is your tech checking the pupil?

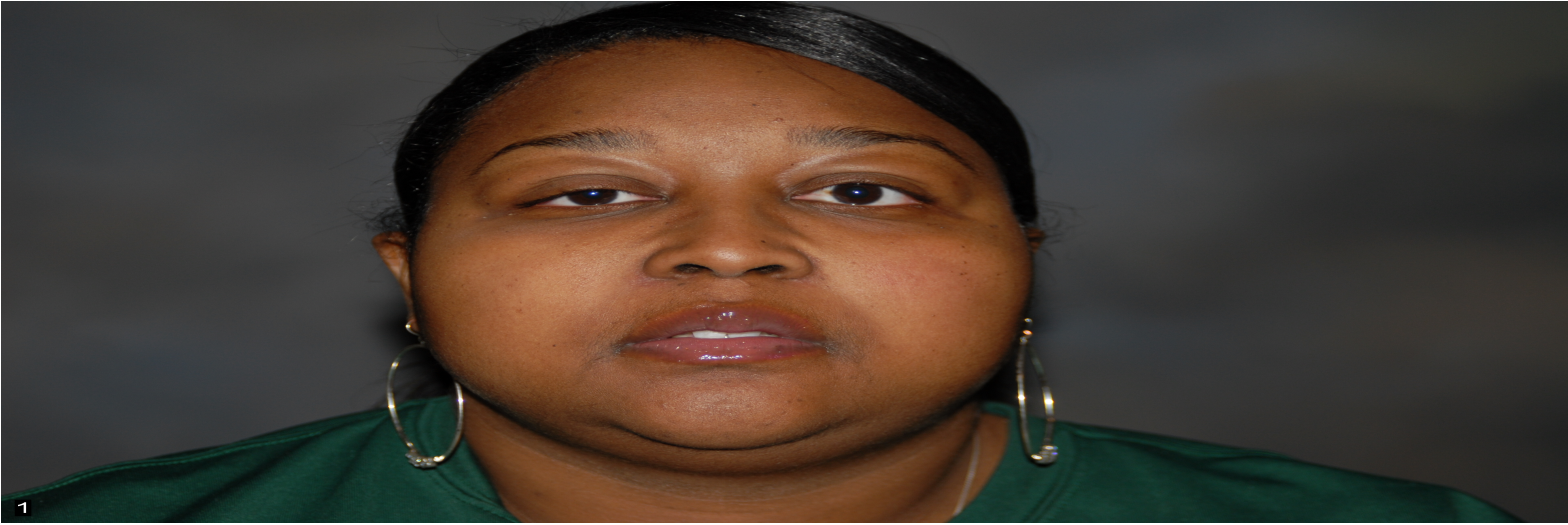
Behavior change(s)

If the chief complaint is diplopia or ptosis , let me check the pupil before dilation

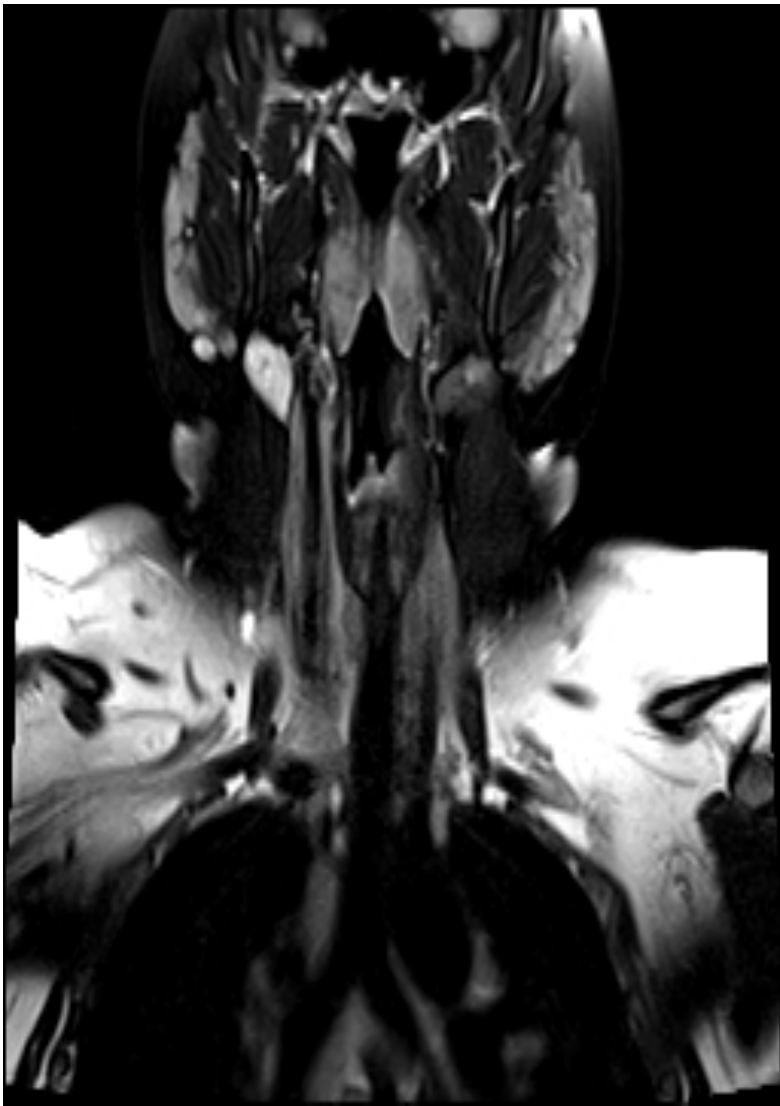
If you have to lift a droopy eyelid before putting in the drops come & get me

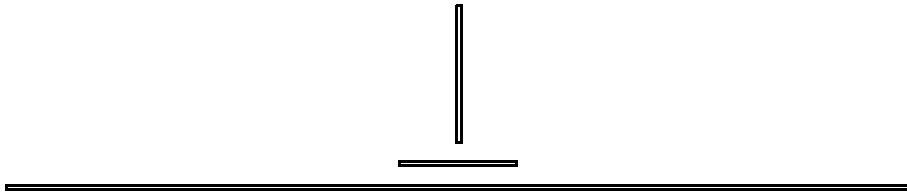
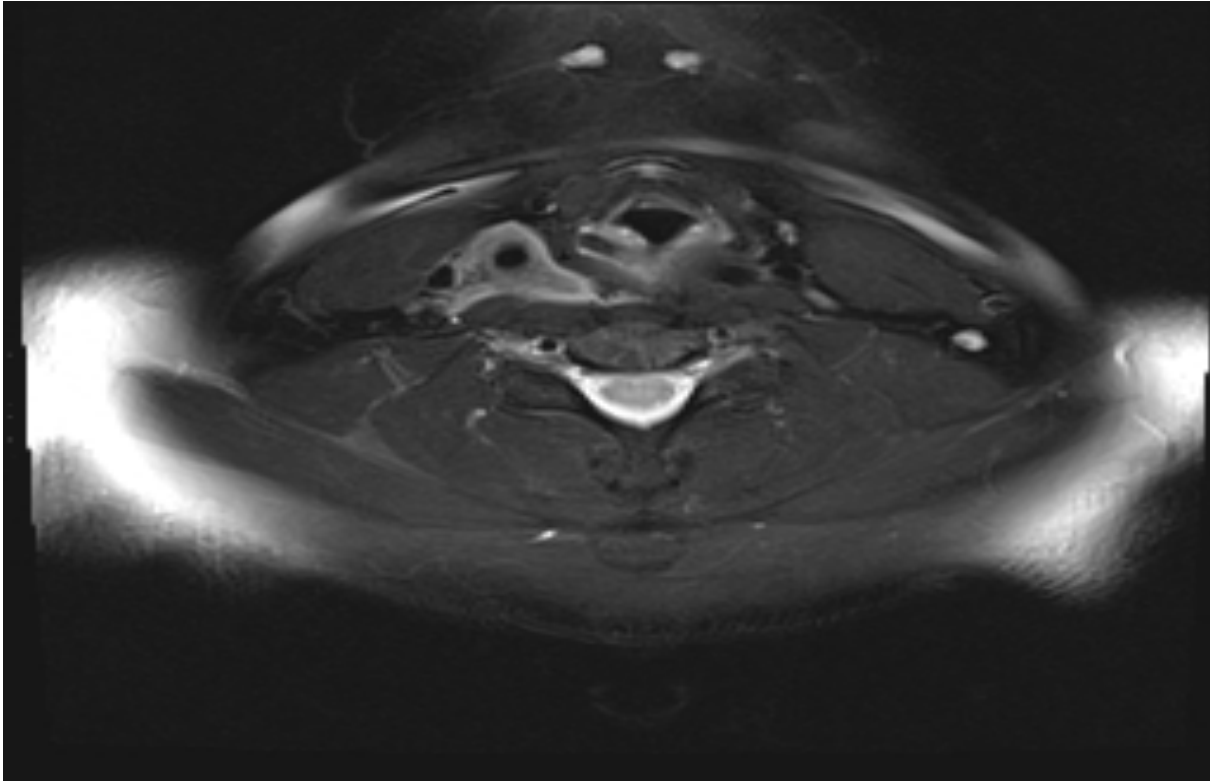
If you have a question about an afferent pupillary defect (??RAPD) come & let me check it too

Horner syndrome RE "Normal MRI head"



Imaging head alone is false sense of security: Pericarotid biopsy proven sarcoid







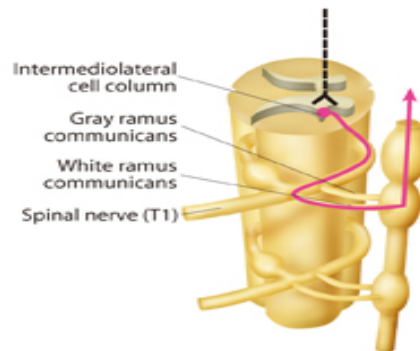
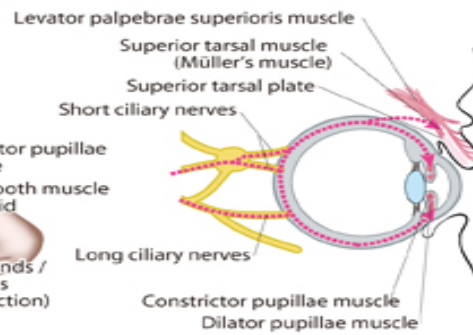
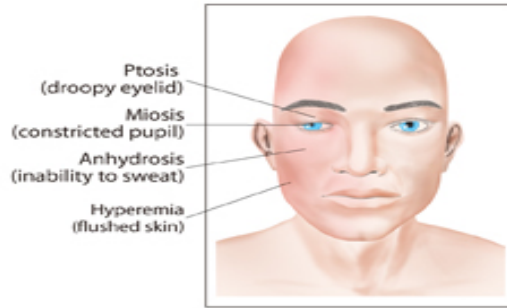
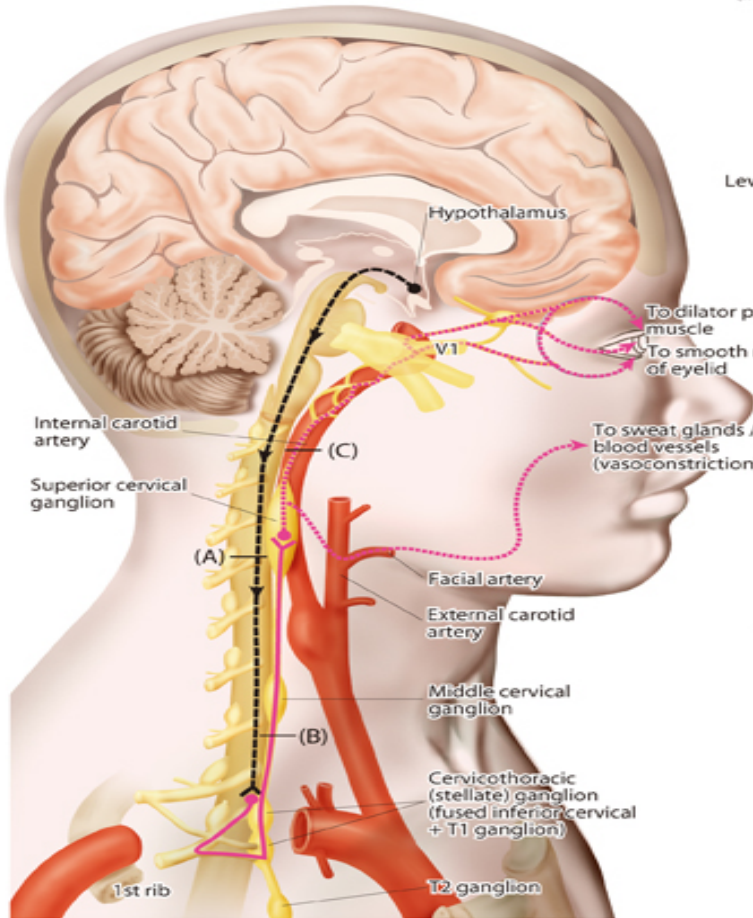
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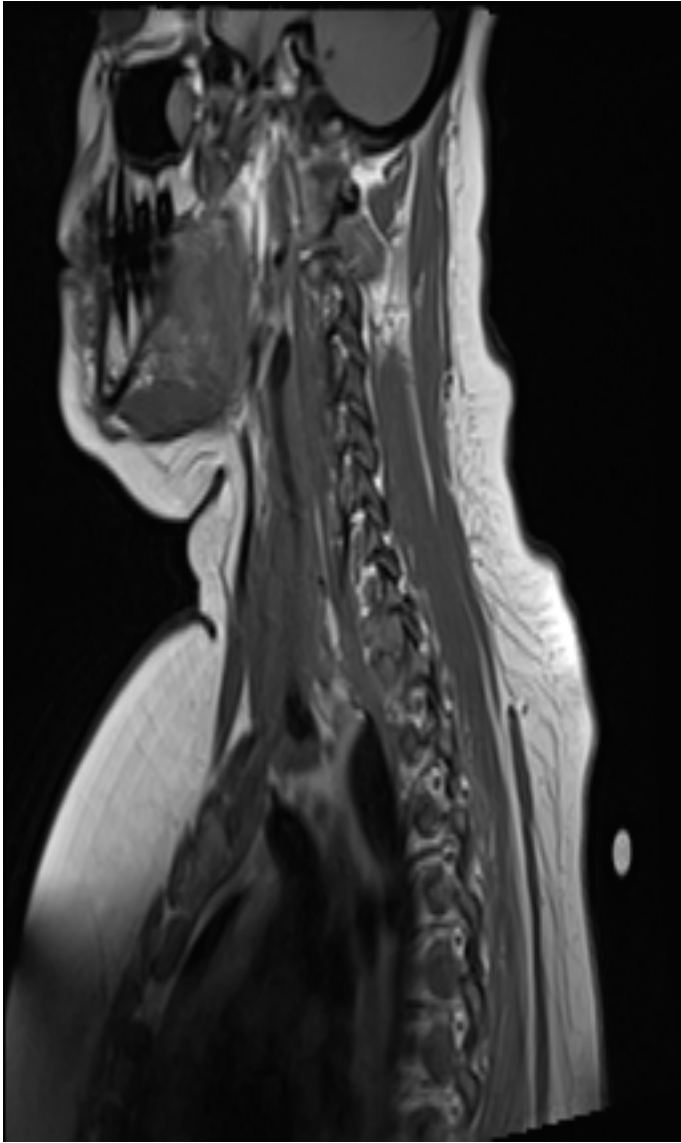
Life threatening cause of Horner syndrome  
= Carotid dissection is extracranial in NECK  
Crescent sign

## Horner's Syndrome

Horner's syndrome results when the cervical sympathetic pathway from the hypothalamus is interrupted. The lesion may be central (A), preganglionic (B), or postganglionic (C) in origin; it may be primary or secondary to another disorder. Symptoms may include ptosis, miosis, anhidrosis (lack of sweating) and/or hyperemia.

- Descending tract from hypothalamus
- Preganglionic tract
- Postganglionic tract





MRI head to neck T2 level

You can image the entire pathway  
with one MRI scan

You could do many overlapping expensive studies

MRI head

MRI neck

CT neck

CXR with apical views

CT chest

Or....you could do one scan (MR head to apex of  
lung (T2 level in chest))

Sagittal & parasagittal imaging on the SIDE of the  
lesion

A Horner protocol MRI

## “PERRLA” failure #2

There is no assessment of the relative afferent pupillary defect (RAPD) in PERRLA

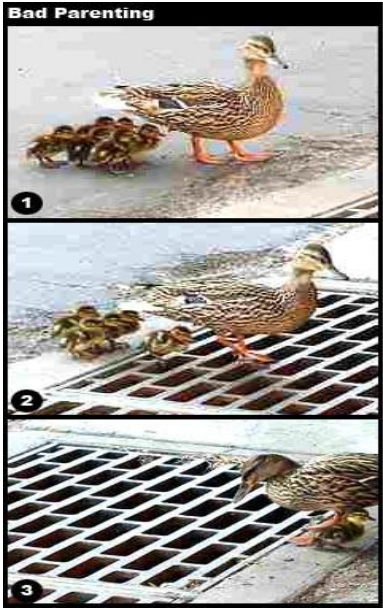
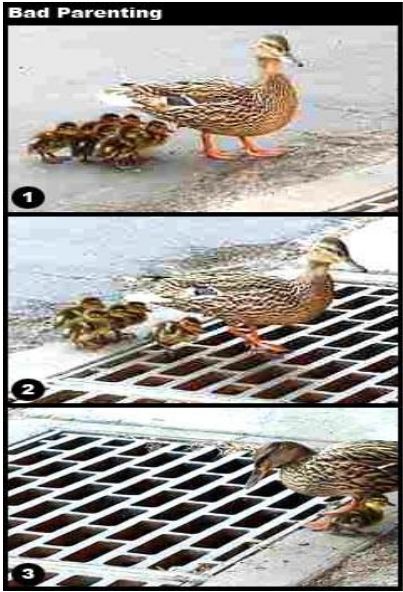
Proper format

OD: Dark 5 mm → Light 3 mm No RAPD

OS: Dark 6 mm → Light 3 mm



Who's fault is it if the resident/fellow/technician doesn't check the pupil properly?



Bad Parenting



Uh-Oh

The behavior change

You: Strike PERRLA from your lexicon & your encounter forms, check tough ones personally

Your tech: Don't use PERRLA, call the doctor for the tough ones

“Blurred disc margin” is a worthless description of what you already know

Does NOT differentiate pseudopapilledema from true papilledema (Both have “blurred disc margins”)

Is disc margin blurred because of something above (peripapillary nerve layer) or below (deeper like drusen?)





LOOK HERE

Obscuration of peripapillary

NFL (blurred VESSELS)

# Never underestimate optic disc edema: When to call neuro-op!

“Next available” optic disc edema

Unilateral nonarteritic AION

Unilateral optic neuritis

Unilateral “neuroretinitis”

Not “next available” (pick up the phone)

Arteritic AION

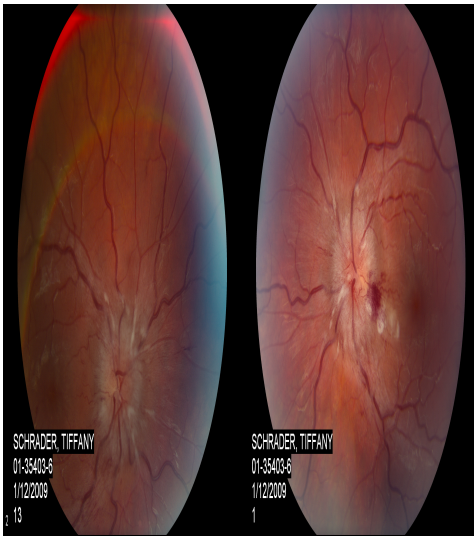
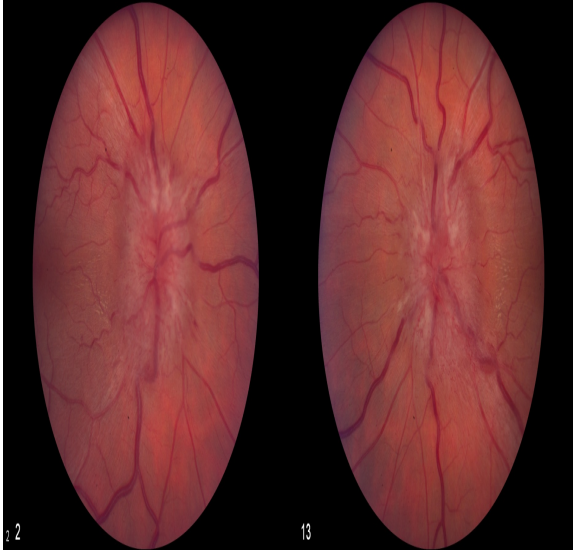
Bilateral optic disc edema (including “neuroretinitis”)

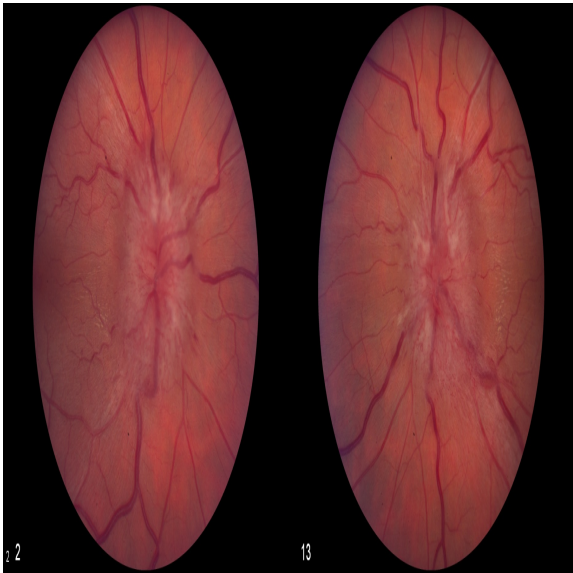
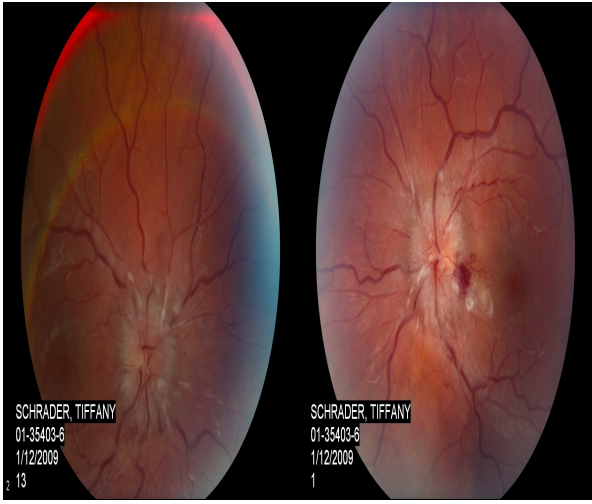
Severe visual loss with disc edema

Chronic atrophic papilledema

Optic disc edema in elderly (rule out giant cell)

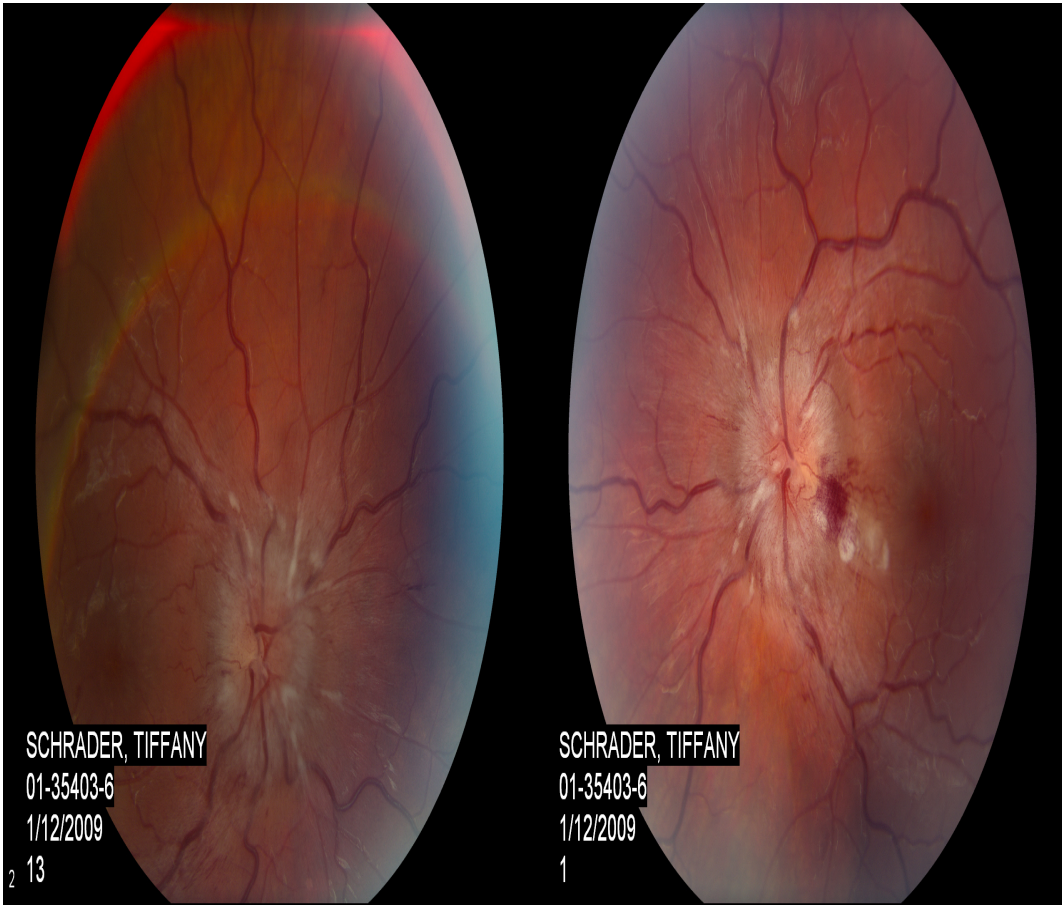
# Which is tumor & which is pseudotumor cerebri?





Don't use "blurred disc margin"

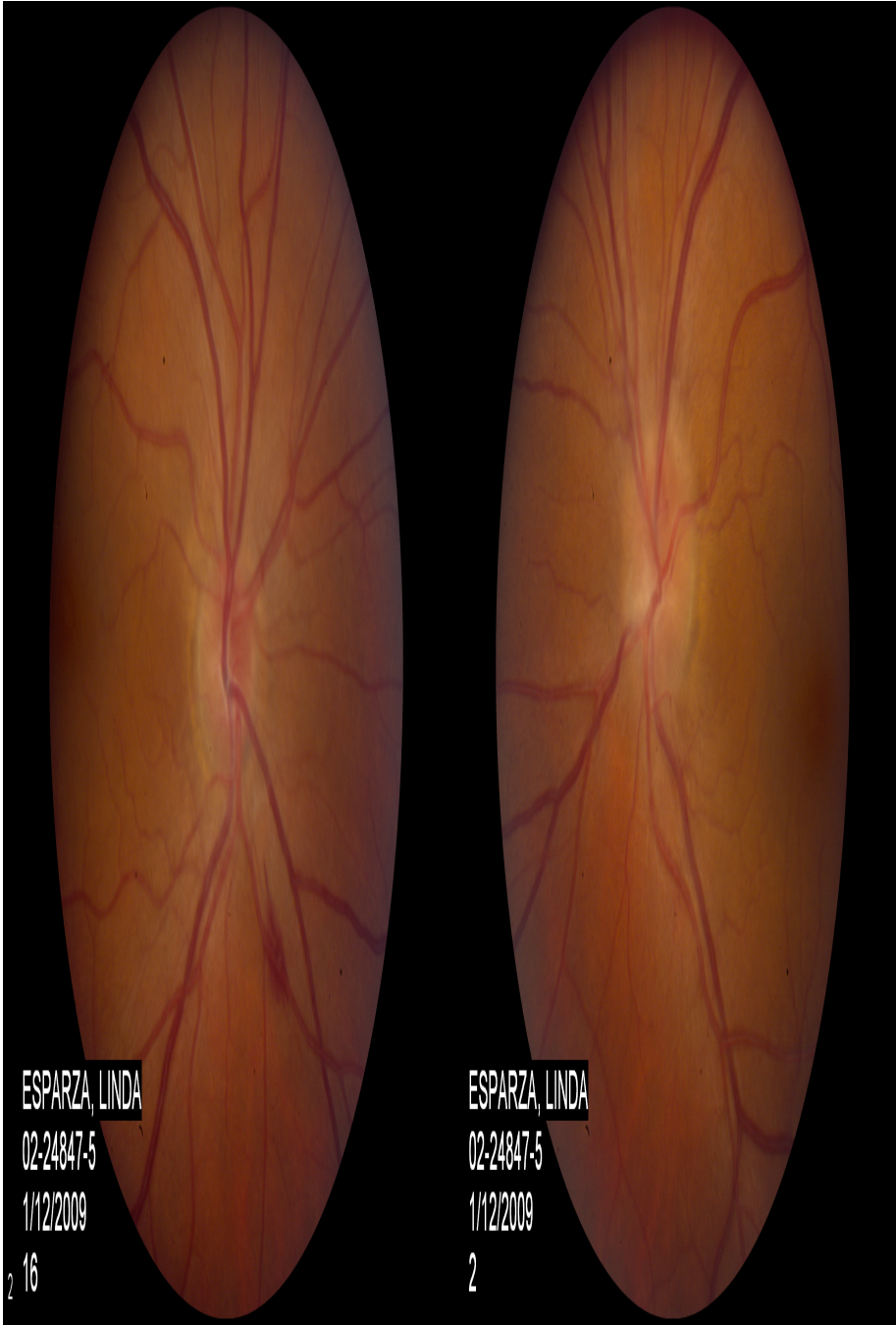


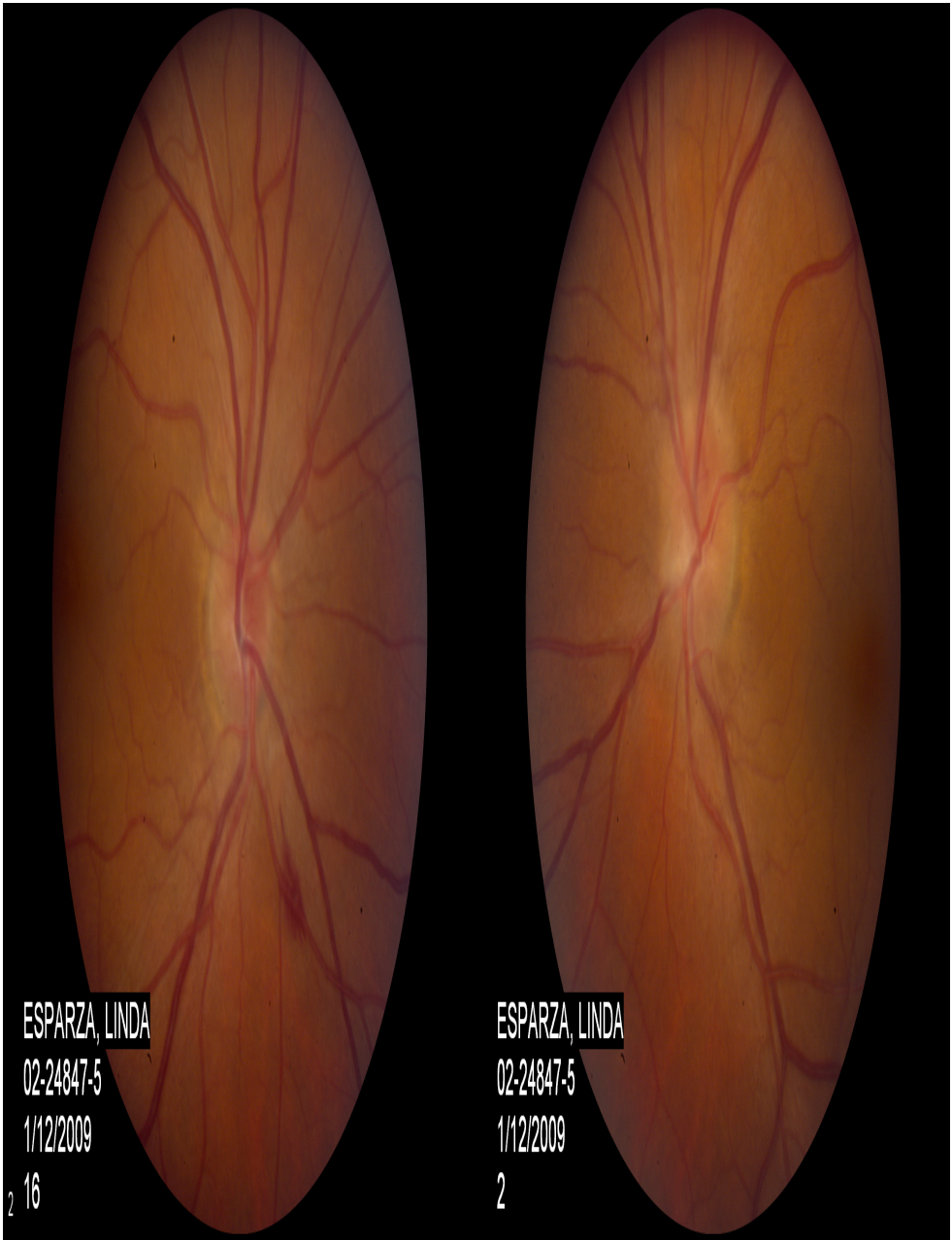


NOT Disc drusen

Disc drusen

Little edema





ESPARZA, LINDA

02-24847-5

1/12/2009

2 16

ESPARZA, LINDA

02-24847-5

1/12/2009

2



The behavior change

MD: Don't write "? papilledema " or use word " papilledema " (increased ICP) when you mean optic disc edema

There are no Little signs in Neuro -ophthalmology (little RAPD, little papilledema have same significance as big!!)

Tech: Don't let your doctor write " papilledema " for optic neuritis, NAION, pseudopapilledema , funny discs, etc.

# Don't use "EOMI" as your sole documentation of motility exam

"EOMI"= Extraocular muscles intact

Primary position deviation will be missed if no cover-uncover testing performed

Small incomitant deviation will be missed if cover-uncover test not performed in diagnostic positions of gaze

Sixth nerve palsy can be “EOMI”

Need cover-uncover test in diagnostic positions of gaze to find small esotropia

Ductions-versions can be normal in patients with ocular motor cranial neuropathies

Book sixth=complete abduction deficit: 50 ET!

Real world 6 th = small incomitant ET in right gaze only



Behavior change

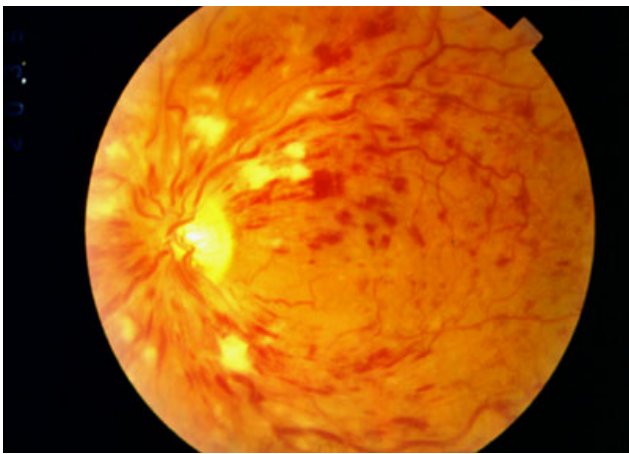
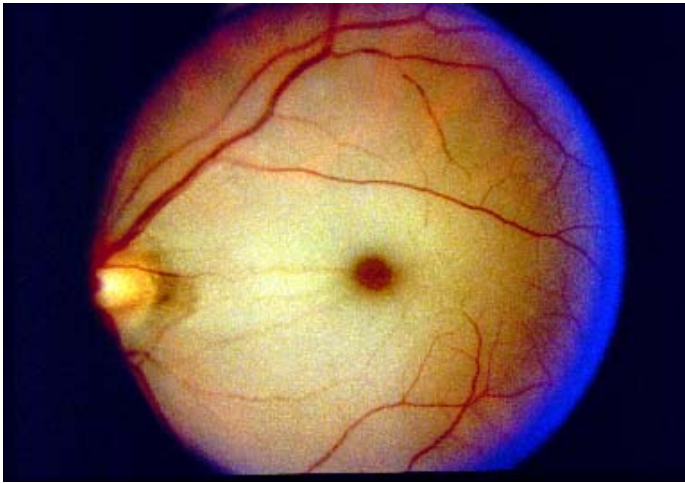
You: Do cover/uncover test for patients with diplopia & test in diagnostic positions of gaze

Don't use symptom (i.e., " diplopia ") or sign (i.e., hypertropia ) as diagnosis or impression

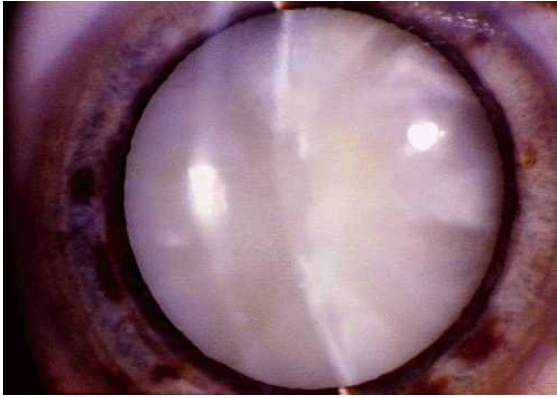
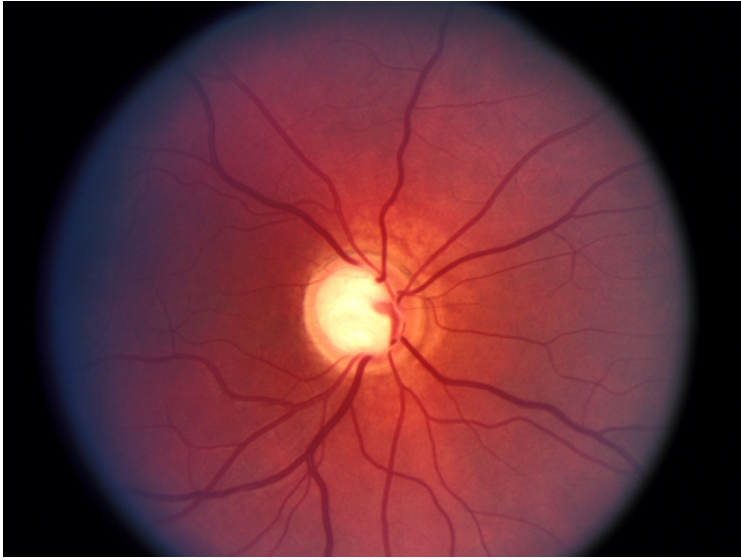
Your tech: Tell eyeMD if diplopia is problem, don't let people leave clinic without a diagnosis

Diplopia is NOT a diagnosis

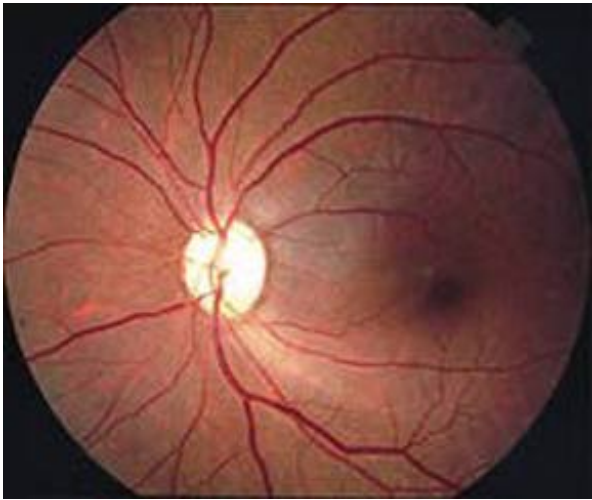
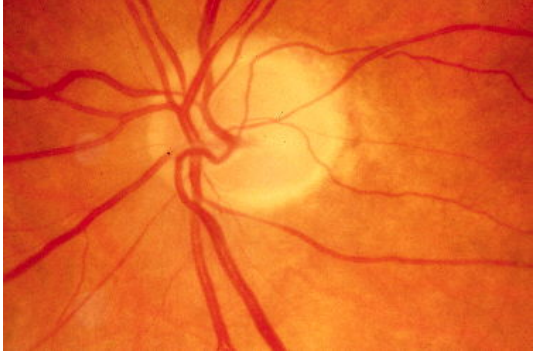
Most of your diagnoses are obvious!  
(Augenblick)

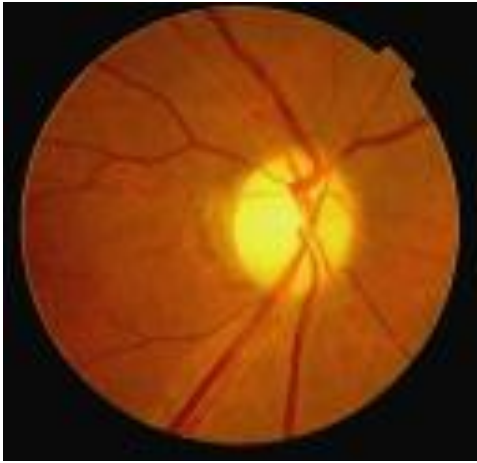






Optic atrophy is **NOT** a diagnosis!  
Not Augenblick!







Is this nerve pale? Mild pallor?  
Temporal pallor? Optic atrophy?



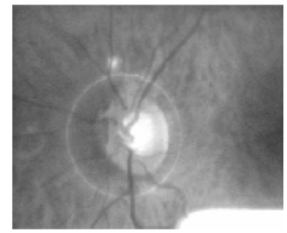
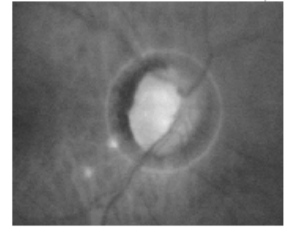
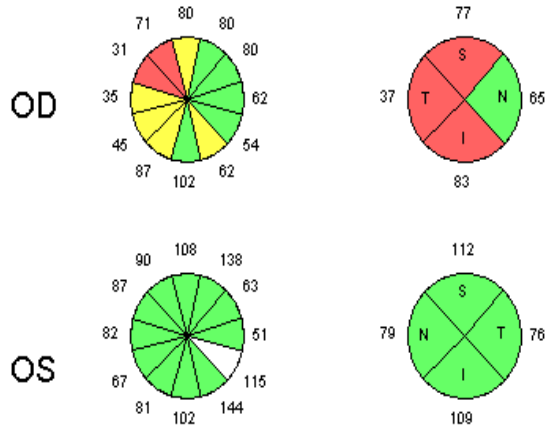
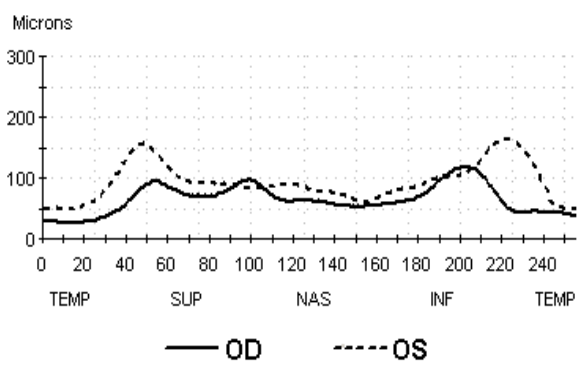
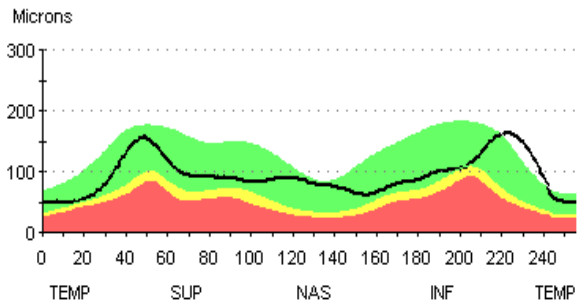
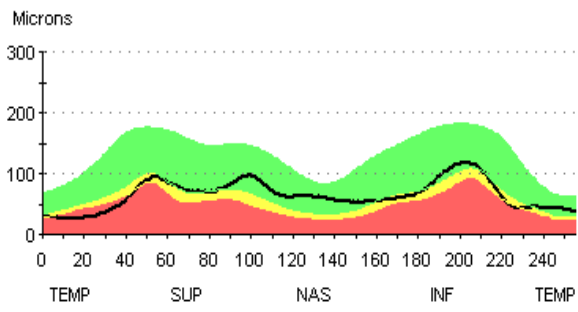
Look for clinical signs of optic neuropathy  
(RAPD, visual field, fellow eye, OCT)





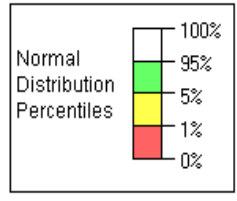
Determination of Pallor vs No Pallor

### RNFL THICKNESS AVERAGE ANALYSIS



Patient/Scan Information	
TEVIS	DENNIS
DOB:	02/09/1947, ID: 98-44624-2, Male
ScanType	Fast RNFL Thickness (3.4)
ScanDate	11/13/2003
ScanLength	10.87

	OD (N=3)	OS (N=3)	OD-OS
lmax/smmax	1.23	1.05	0.18
smmax/lmax	0.81	0.95	-0.14
smmax/tavg	2.59	2.04	0.55
lmax/tavg	3.20	2.15	1.05
smmax/navg	1.47	1.98	-0.51
Max-Min	91.00	114.00	-23.00
smmax	96.00	156.00	-60.00
lmax	118.00	164.00	-46.00
savg	77.00	112.00	-35.00
lavg	83.00	109.00	-26.00
Avg.Thickness	65.61	94.00	-28.38



OCT can see better than me

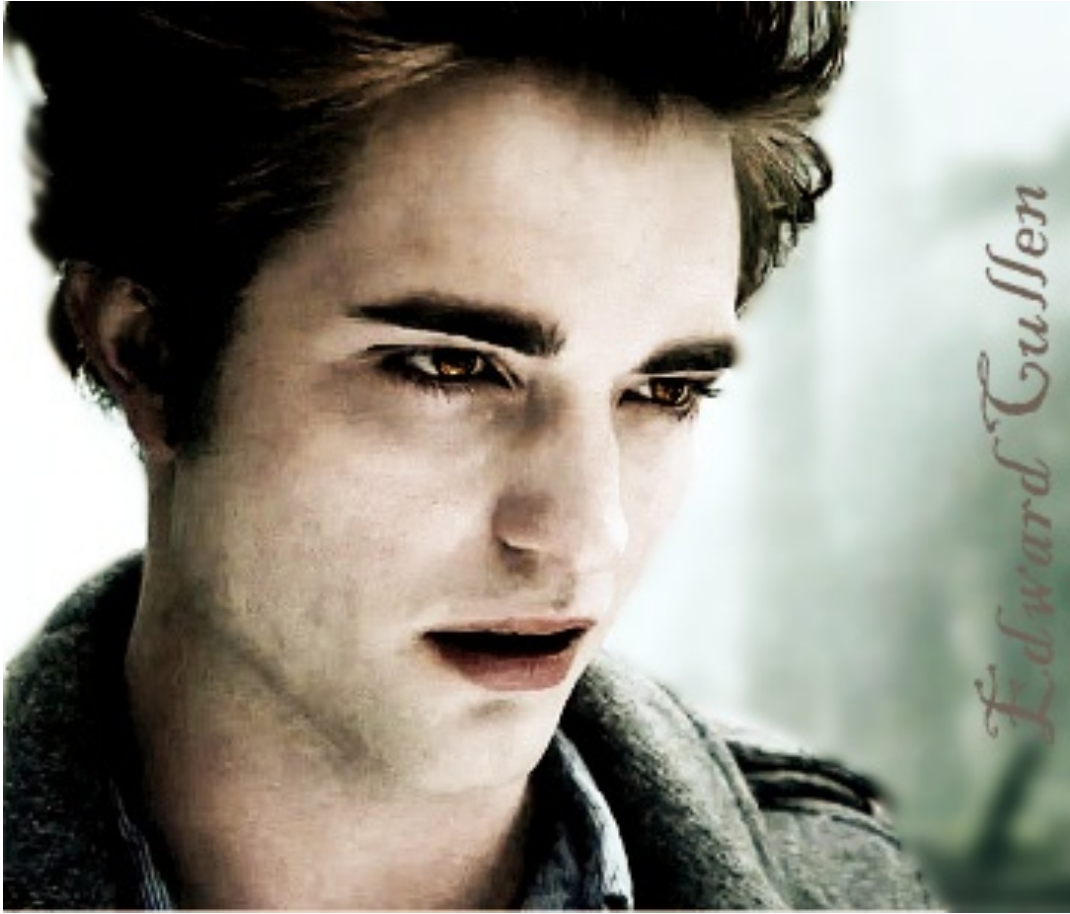


Am I pale?





Am I pale?

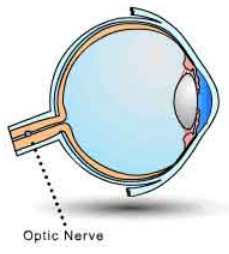
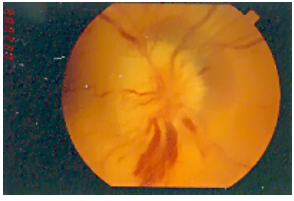




# Common things are common

Is it old AION

Is it old optic neuritis?







Dad's rules of DDX

If it sounds like a duck, looks like a duck, & acts like a duck then it's a #@! Duck

Is it old AION?

Disc edema

Vasculopath

Older patient

Static course

Is it old ON?

Younger

Recovered

MS history



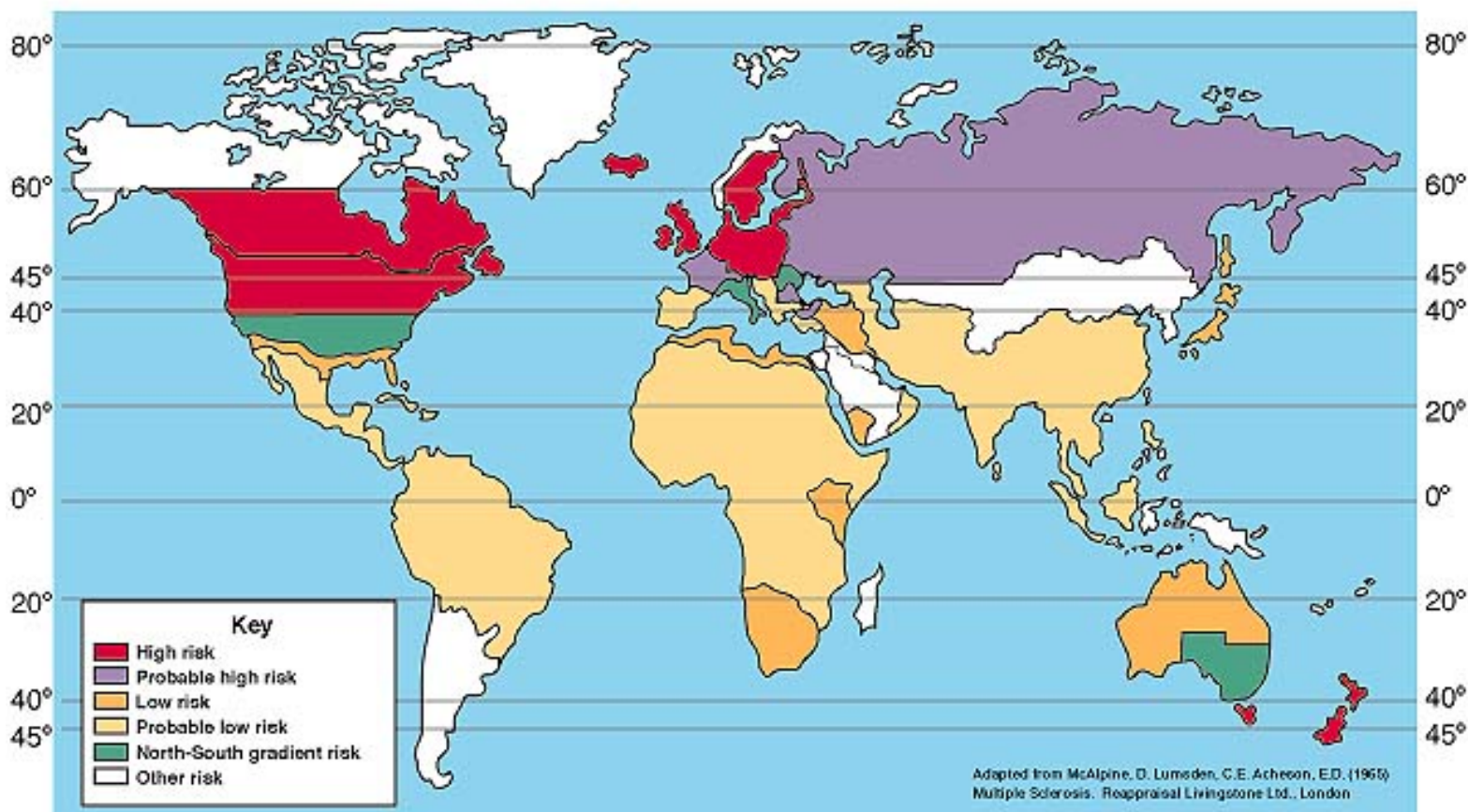




Uncommon presentations of  
common diseases are **COMMON**

# MS: Old optic neuritis?

## World Distribution of Multiple Sclerosis



If not AION or ON then more history & exam

Bilateral progressive central-cecocentral scotoma  
=> B12/folate/Leber's hereditary optic  
neuropathy/ethambutol toxicity

Chronic progressive optic neuropathy =>  
compressive lesion (get formal fields)

Bitemporal hemianopsia: Chiasmal

Homonymous hemianopsia: Optic tract

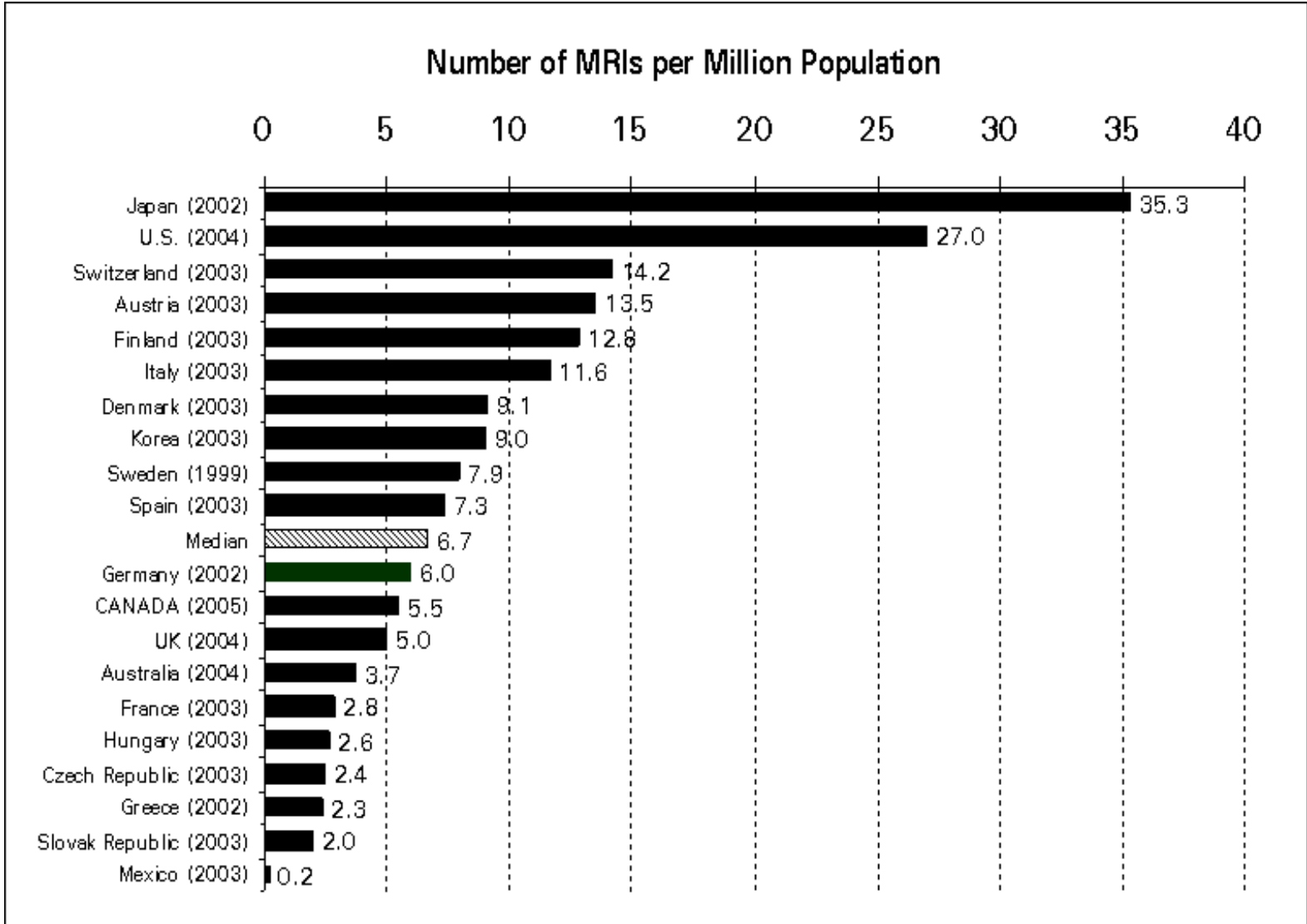
Uveitis (old or new) Sarcoid, syphilis

# If history & exam come up short then image unexplained optic atrophy

MRI head/orbit fat suppression and gadolinium (optic nerve protocol)

If suspicion low for compression or cost is an issue in your part of the world you might choose observation (if static , old NAION then no imaging) or CT scan with contrast

Optic atrophy can always be a tumor!



China and India - 1 MRI per 1 million population

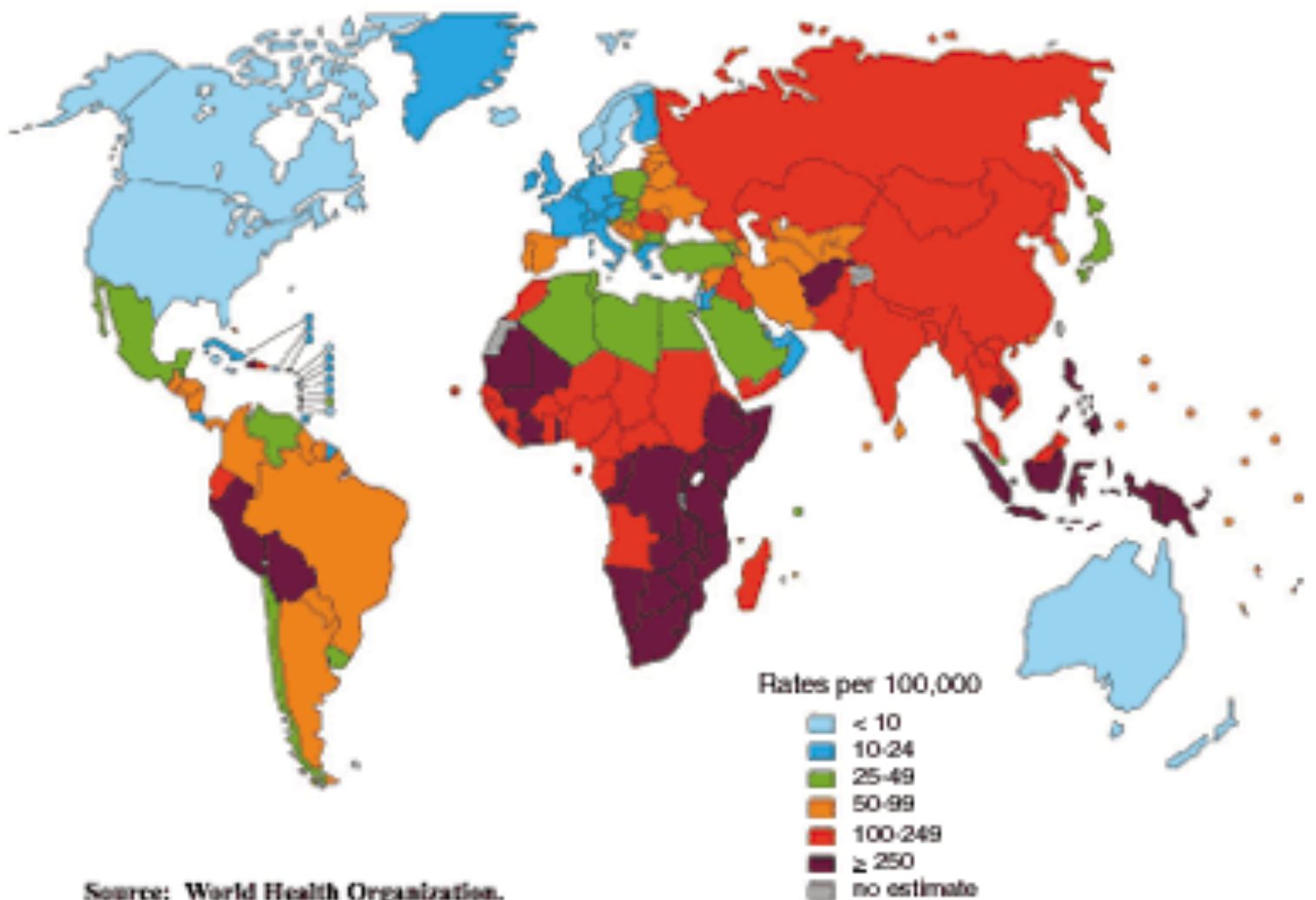
Directed evaluation vs. shotgun





## Tuberculosis in India

### Estimated Rates of New Cases of Tuberculosis, 1997



## Syphilis worldwide



Round up the usual suspects



Compressive

Ischemic

Demyelinating

Infectious

Inflammatory

Toxic-nutritional

# Optic atrophy is NOT a diagnosis

Impression equals diagnosis = most of your clinic day!

Cataract

CRVO

CRAO

RD

Optic atrophy is NOT a diagnosis

Could be a compressive lesion

Image if unexplained optic atrophy (don't write "? Mild")

If not imaging document WHY (e.g., "I believe that this is old NAION and I am following this patient")

Document RATIONALE for decision making

Behavior change

You: Don't use "optic atrophy" as a diagnosis

Your techs: Don't let people leave clinic with a photo or OCT or chart that says "optic atrophy" and has NO etiologic diagnosis

The ways that an ophthalmic technician can save a life....

Avoid sole use of PERRLA for the pupil exam

Savino's rule - if there is a problem with the lid, motility, or pupil all three areas must be evaluated and documented

Don't use EOMI as the only assessment of motility

Don't order the same scan on all patients; and

Signs or symptoms are not diagnoses.

Five behavior changes TODAY

Check pupil in light & dark (not “PERRLA”)

Don’t let technician be only pupil exam for tough ones

Avoid “Blurred disc margins” & take the finding seriously (i.e., “? papilledema ”)

Don’t use vague motility terms like “ dysconjugate gaze” or “ extraocular muscles intact (“EOMI”)

Remember: “optic atrophy”, “ diplopia ”, “ esotropia ” are not diagnoses (PS: Neither are “ ptosis ” or “blurred vision” or “unexplained visual loss”)

There is a difference between data  
and information

DATA

28 17 26 80 81

INFORMATION

(281) 726-8081

If you have questions call me  
or email me

[AGLee@tmhs.org](mailto:AGLee@tmhs.org)



End with a philosophical question & two really quick cases. Why are you here... because you believe as we all do that you can....?

Chief complaint: NONE

73-year-old WF

Chief complaint: NONE now (2010)

PMH: Paraneoplastic optic neuropathy, recovered

CXR: Small cell carcinoma of lung

Resected, chemotherapy, radiation in 1997

Published: Luiz JE , Lee AG , Keltner JL, Thirkill CE, Lai EC. Paraneoplastic optic neuropathy and autoantibody production in small-cell carcinoma of the lung. *J Neuroophthalmol.* 1998;18:178–181.

Follow up 2010

Pt: “You don’t remember me do you Dr. Lee?”

Me: “Well,...I um....sure...maybe”

Pt: “I had lung cancer & you found it thru my eye”

Me: “Really”

Pt: “Yeah, you wrote it up in a journal”

Me: “Oh, yeah, sure, now I remember. How are you, why are you coming today?”

Pt: “I just wanted to tell you that I was still alive and it is been 14 years, so thanks.”

## Longest known survivor

Long\_Term\_Survivor\_of\_Paraneoplastic\_Optic.17.pdf - Adobe Reader  
File Edit View Document Tools Window Help  
2 / 5 151% Find

# Long-Term Survivor of Paraneoplastic Optic Neuropathy

**S**mall cell lung cancer carries a very poor long-term prognosis. In a survey performed at the Mayo Clinic from 1997 to 2003, the 5-year survival rate was only 9% (1). In addition, to our knowledge, the longest published survival duration for paraneoplastic optic neuropathy secondary to small cell lung cancer has been 8 years (2). We wish to provide an update on a patient previously reported by one of us (A.G.L.) in this Journal in 1998 (3) who returned 14 years later without evidence of tumor recurrence and believed to be in clinical remission. The earlier detection of the tumor from her neuro-ophthalmologic examination followed by timely systemic treatment may have contributed to her favorable outcome. To the best of our knowledge, she is the longest survivor of paraneoplastic optic neuropathy secondary to small cell lung cancer. At the time of her diagnosis, she underwent surgery, chemotherapy, and radiation therapy and was believed to be in remission at the last follow-up.

The patient, a 73-year-old white woman, was last seen in the neuro-ophthalmology clinic on July 20, 2010. She was complaining of blurred vision in the left eye that had worsened since sustaining a fall on March 1, 2010. She was seen by her neurologist who obtained a brain MRI that showed no focal lesions.

in March 2010 showed no evidence of recurrent or metastatic disease. The patient returned to The Methodist Hospital after 10 years of follow-up to specifically report on her progress and survival from small cell carcinoma of the lung.

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Iowa City, Iowa*  
*Department of Ophthalmology, UTMB-Galveston  
Galveston, Texas*

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His name is Andrew....

33-year-old WM

Transient dizziness, blurry vision, followed by loss of consciousness after watching bungee jumpers at Iowa St. Fair

On regaining consciousness, bilateral ptosis, exotropia: Noncontrast cranial CT in ER was “normal” MRI with contrast : “normal”

About to be discharged

MRI head negative

Course

Top of the basilar syndrome

Intravascular tPA

Locked in syndrome

Recovered slowly

Rehab, walked out of hospital

Writing a book about his experience  
called “One Fine Day”....

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### PACEMAKER: Winter 2006-07

## 'Just happy to be here'

**34-year-old stroke patient achieves a remarkable recovery with help from the experts at UI**



Like many people, Bill Pitzen used to associate strokes with age.

Today, Pitzen knows better. Having experienced two strokes of his own at age 33, he realizes that age has little to do with one's risk for stroke.

The first event occurred last year during a seemingly routine day at his former place of employment, McLeod USA in Cedar Rapids, Iowa.

The initial symptoms—dizziness, hot flashes, slurred speech, and double vision—rendered him unresponsive as paramedics rushed him to a nearby hospital for evaluation.

Fortunately, he woke up and was able to return home after MRI and CT scans failed to detect the cause of his symptoms. Still, doctors were concerned and referred him to University of Iowa Hospitals and Clinics for further evaluation by UI neuroscientists.

Two days later, during a clinic visit, UI neuro-ophthalmologist Andrew Lee, MD, diagnosed Pitzen with a significant stroke. A repeat MRI scan also showed a developing blockage in his brain, putting him at high risk for a second stroke.

Unfortunately, Pitzen suffered the second stroke and collapsed just as he was being admitted for treatment.

Pitzen now faced the distinct possibility of losing movement on his left side and the ability to feed himself. To counter these risks, UI neuroscientists quickly formulated a plan that they hoped would lead to a better outcome.

Time was the enemy. The sooner doctors could intervene, the better. A highly specialized endovascular procedure to free the

#### Grand Slam

Stroke survivor Bill Pitzen celebrates by running the bases at a "Homerun for Life" event sponsored by the Cedar Rapids Kernels baseball club.

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A Neuro-ophthalmologic Emergency - Windows Internet Explorer

http://www.ophsource.org/periodicals/ophtha/article/S0161-6420(06)00576-8/abstract


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A Neuro-ophthalmologic Emergency

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Volume 113, Issue 8, Pages 1477-1477.e2, August 2006

## A Neuro-ophthalmologic Emergency

[Sandeep Randhawa, MD](#), [Vinay A. Shah, MD](#), [Shereen Chang, MD](#), [Harold P. Adams Jr, MD](#), [Andrew G. Lee, MD](#)  
Iowa City, Iowa

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Years later...

Receive a phone call from this patient

“ Hey, Dr. Lee...you don't remember me probably but I had a stroke at age 33 and you helped me at Iowa ”

Me: “Sure, I remember you ”

“ I was just calling to let you know that I went back to college, I got married, and now I have a new baby, his name is Andrew ”

Me: “ That is so great, congratulations ”

“ No, Dr. Lee you don't understand...his name is ANDREW!”

ONE PERSON  
CAN MAKE A  
DIFFERENCE,  
AND EVERYONE  
SHOULD TRY

-JOHN F. KENNEDY-

Thank you for your time & attention



BCM

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Medical College



Thanks for your attention

