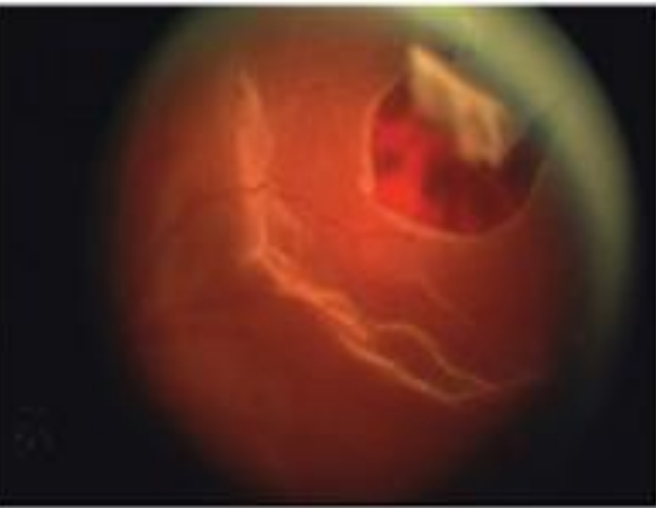
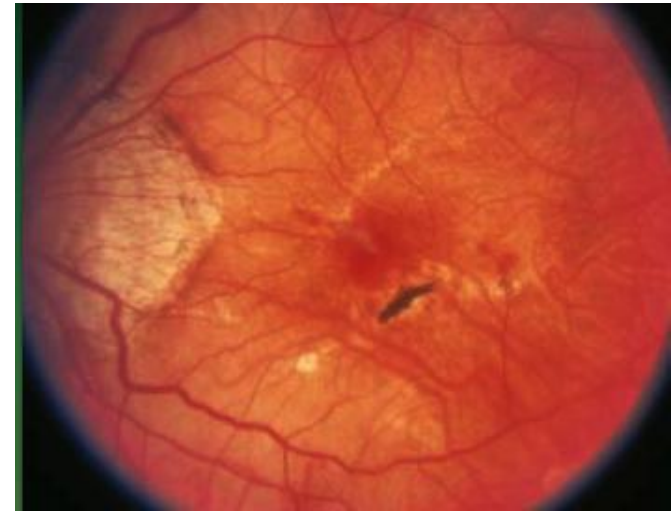


# PATHOLOGICAL CORRELATES OF MYOPIA



TEAR WITH DETACHMENT



FUCH'S SPOT

LIONEL KOWAL

ACBO 2009

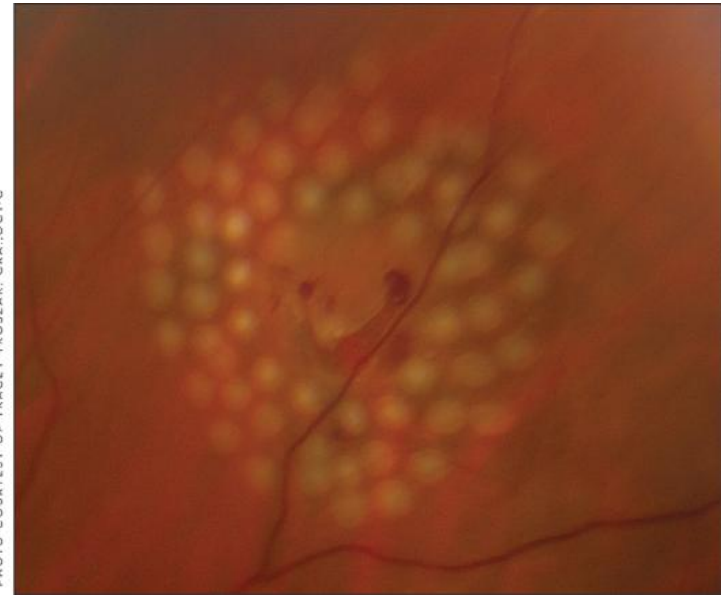
# OCULAR : ISSUES IN DAILY PRACTICE

- Risk of retinal detachment
- Foster- Fuch's spot *Refer early ?VEGF drugs*
- Disc changes / Staphyloma
- Symptomatic floaters
- Worrying signs / symptoms
- Difficulty of scleral indentation

## ***NON- OCULAR***

- *ORBITAL*
- *SYSTEMIC*

PHOTO COURTESY OF TRACEY TROSAK, CRA.,OCT.C



# OCULAR CHANGES IN MYOPIA

As axial length increases, there is an increasing disparity between surface area of retina  $4\pi r^2$  and volume  $\frac{4}{3}\pi r^3$

Pathology more common with increasing myopia esp > -6

# 5 fundus changes are associated with increased axial length of the eye.

- **Optic nerve crescent**

Not visually threatening – can be confusing

- **Chorio- retinal atrophy**

Common. Not visually threatening.

- **Central pigment spot (Fuchs's)**

Uncommon. Can be visually threatening.

- **Lacquer cracks**

Uncommon. Can be visually threatening.

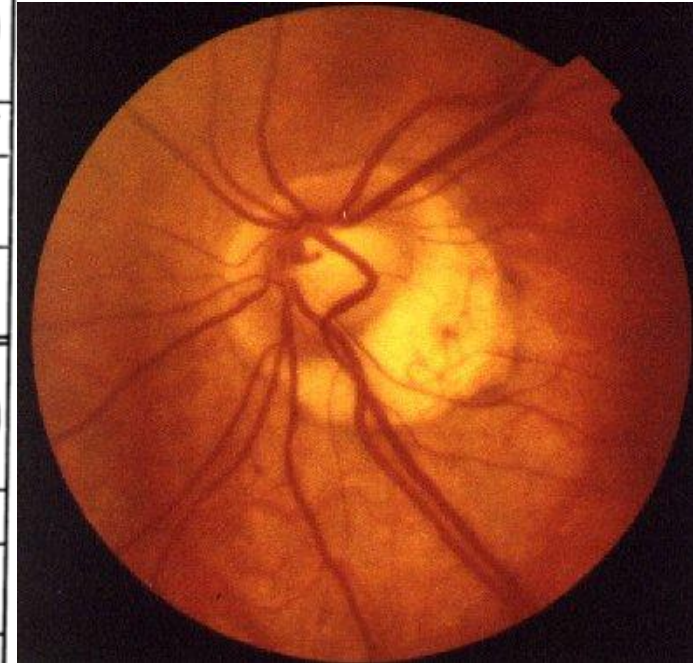
- **Posterior staphyloma**

Not visually threatening – can be confusing

# Myopic discs

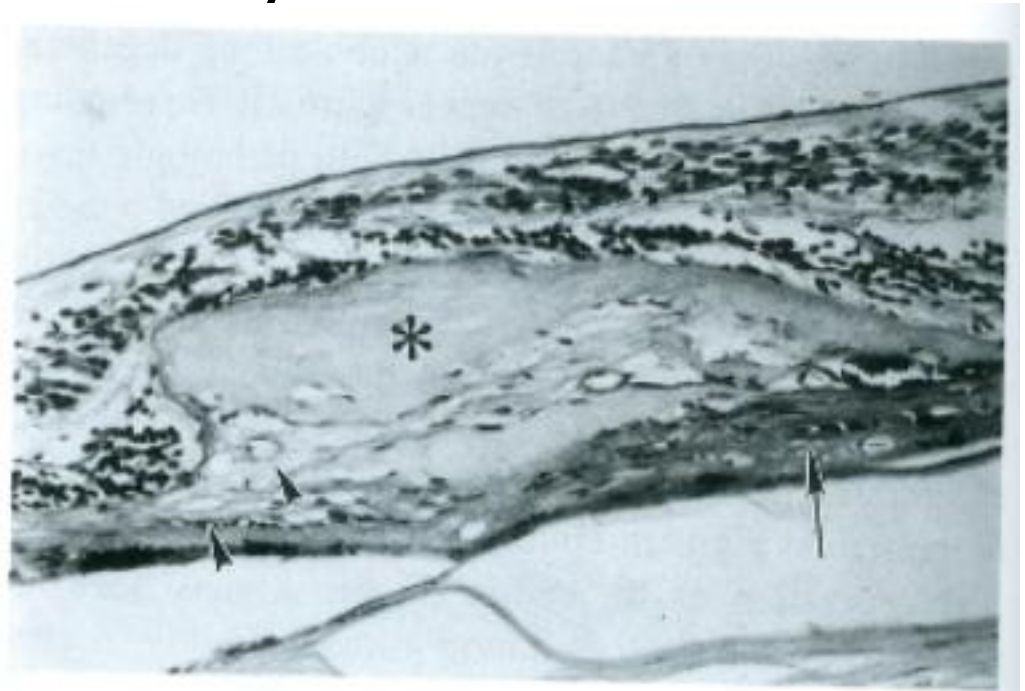
## 100% of all eyes >28.5mm

OPTIC NERVE CRESCENT: TYPE AND INCIDENCE						
APPEARANCE (LEFT EYE)						
NAME	TEMPORAL	ANNULAR	NASAL	TEMPORAL- ANNULAR	INFERIOR	TEMPORAL- INFERIOR
ALL CRESCENTS TOTAL 1032	62%	25%	3%	2.7%	2.5%	2.3%
CRESCENTS WITHOUT PERIPAPILLARY ATROPHY 841 TOTAL	71%	17%	2.9%	2.4%	2.7%	2.2%
APPEARANCE (LEFT EYE)						
NAME	NASAL- INFERIOR	TEMPORAL- INF.-NASAL	NASAL- ANNULAR	INFERIOR- ANNULAR	SUPERIOR	TEMPORAL- NASAL
ALL CRESCENTS TOTAL 1032	<1%	<1%	<1%	<1%	<1%	<1%
CRESCENTS WITHOUT PERIPAPILLARY ATROPHY 841 TOTAL	<1%	<1%	<1%	<1%	<1%	<1%



# Central Pigment Spot (Fuchs's)

- A black area of variable diameter at the macula occurs in ~5% of eyes  $\geq 26.5$  mm or more axial length



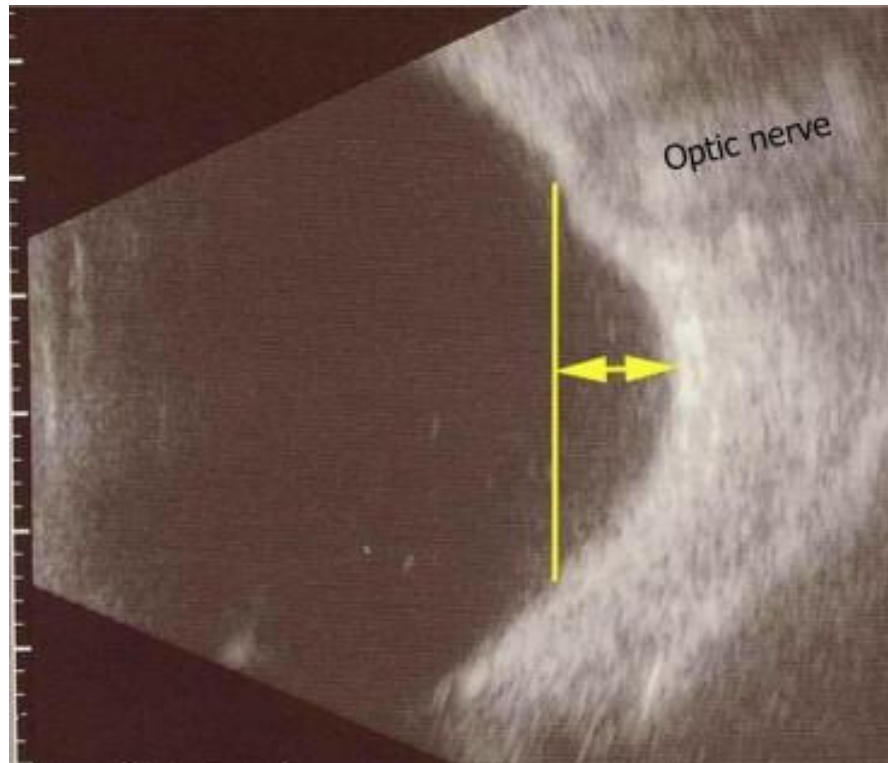
**Fig. 10.** Fuchs spot in this eye consists of a subretinal fibrovascular nodule (asterisk) with a defect in Bruch's membrane (arrow marks one margin of defect) with vessels (arrowheads) from the choroid (periodic acid-Schiff; original magnification  $\times 120$ ).

# Lacquer cracks





# Posterior staphyloma





# Classic pathology paper

## Grossinklaus & Green

- 308 eyes with pathological myopia
- 285 postmortem. 23 surgical...over 67 y!
- Myopic disc 38%
- Post staphyloma 35%
- Degenerative vitreous 35%
- Cobblestone 14%
- Myopic degeneration of retina 11%
- Retinal detachment 11%
- Retinal pits, holes, tears 8%
- SRNV 5%
- Lattice 5%
- Fuch's spot 3%
- Lacquer cracks 0.6%

### Surgical:

Degeneration after ret det  
2ary glaucoma  
Endophthalmitis  
Expulsive h'age  
Epithelial ingrowth  
Degen after cataract surgery  
Presumed tumour

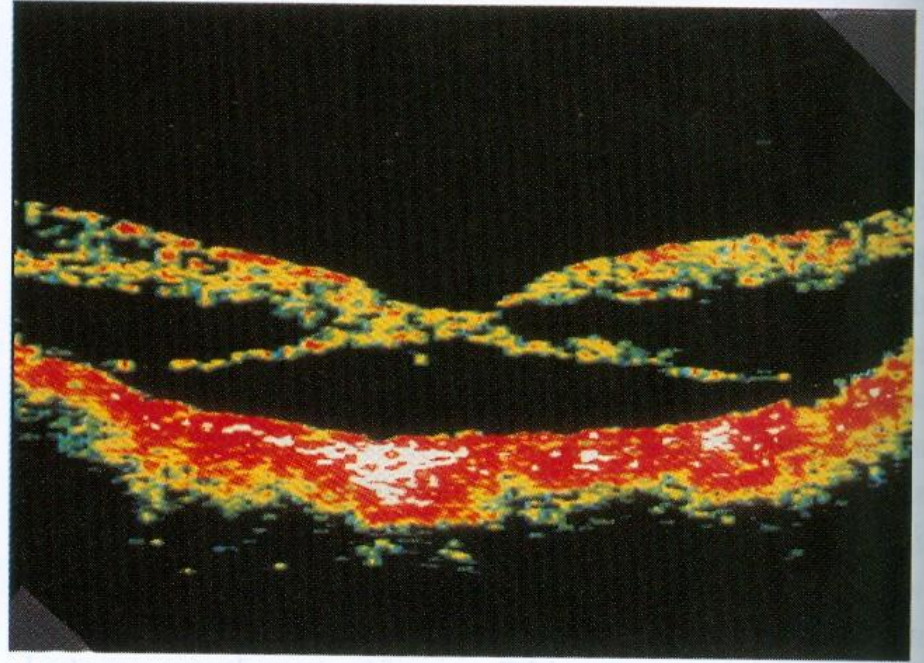
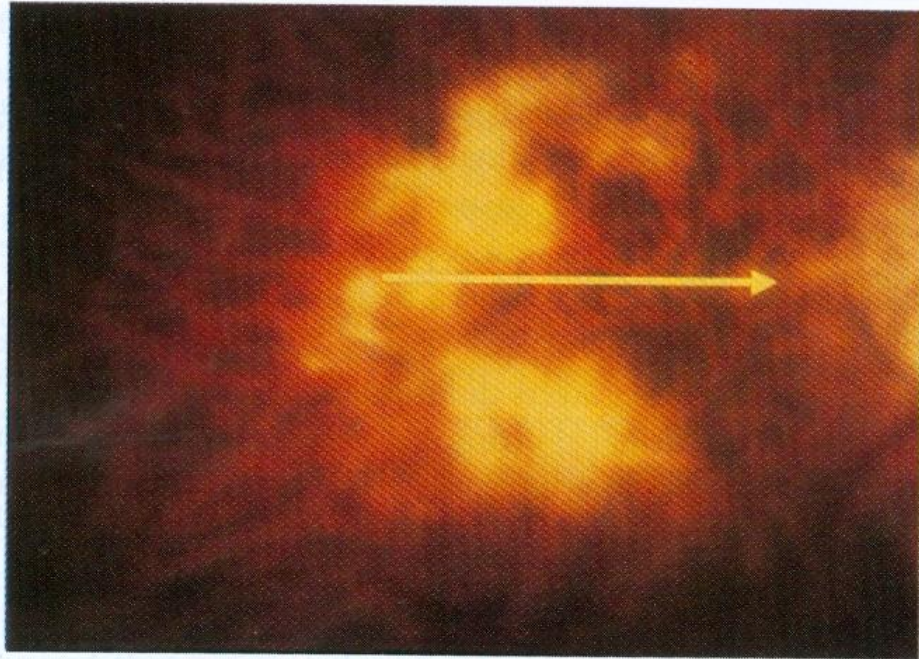
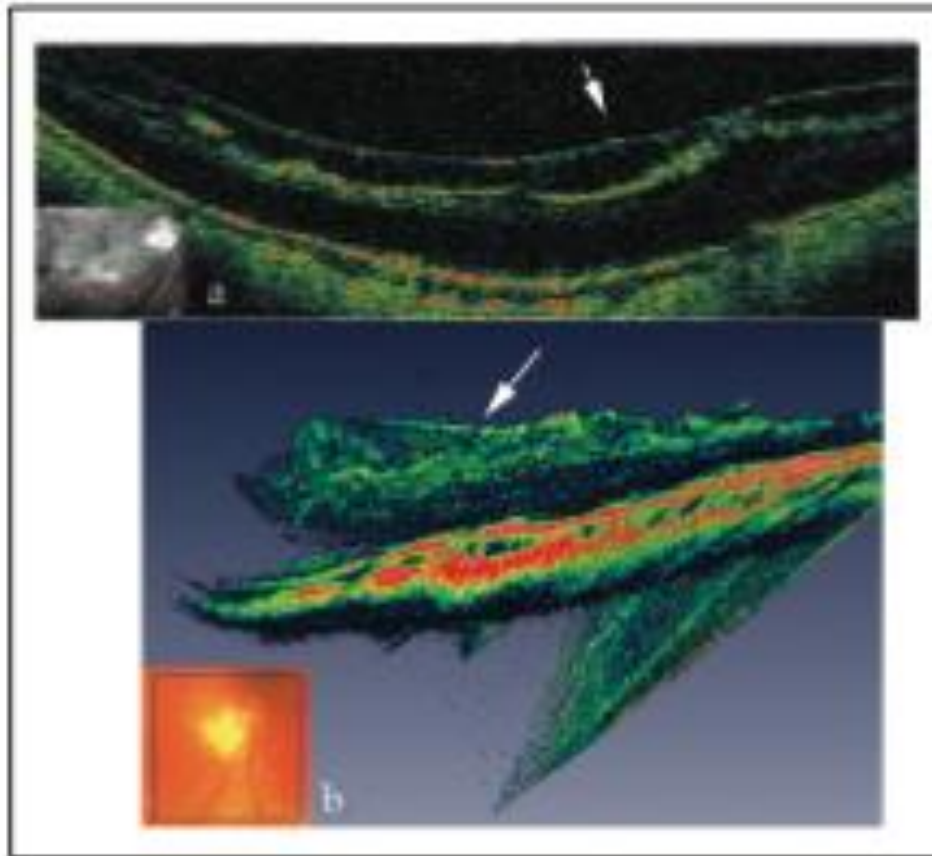


FIGURE 2. Right eye of a 41-year-old woman, with refraction of  $-17$  diopters. Best-corrected visual acuity of the right eye was 20/400, and the axial length was 25.7 mm. (Left) Scars from photocoagulation performed 10 years earlier are evident in the perifoveal area. The arrow indicates the area of the optical coherence tomographic scan. (Right) Optical coherence tomography shows a retinal detachment at the fovea. The overlying retina has retinoschisis in the perifoveal area.

# OCT redefining pathology

Retinoschisis



**Figure 2.** Case 2. Retinoschisis (arrow) is evident in the right eye at both (a) cross-sectional B-scan and (b) three-dimensional visualization.



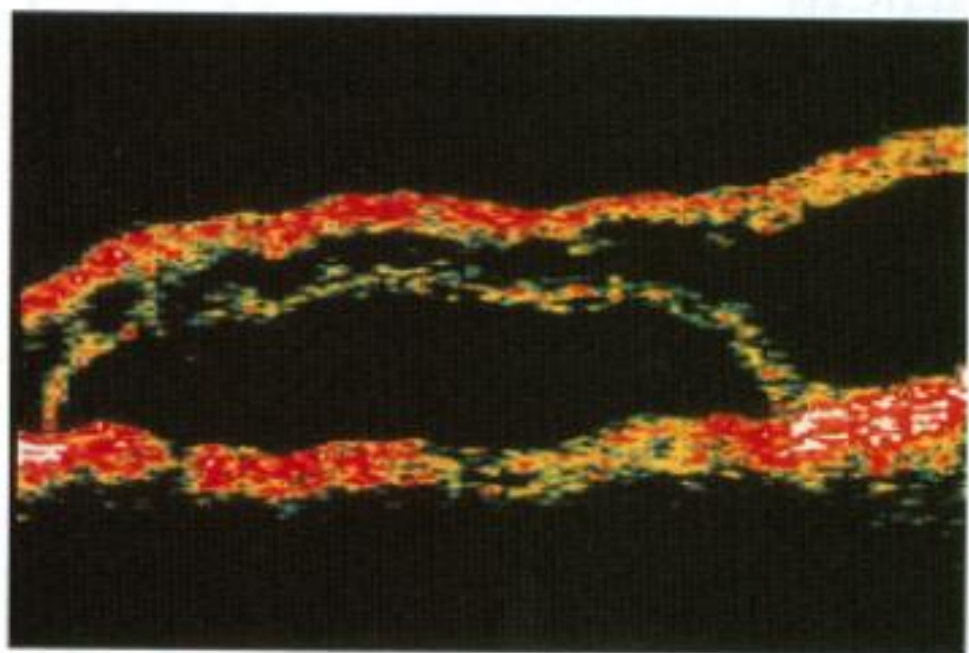
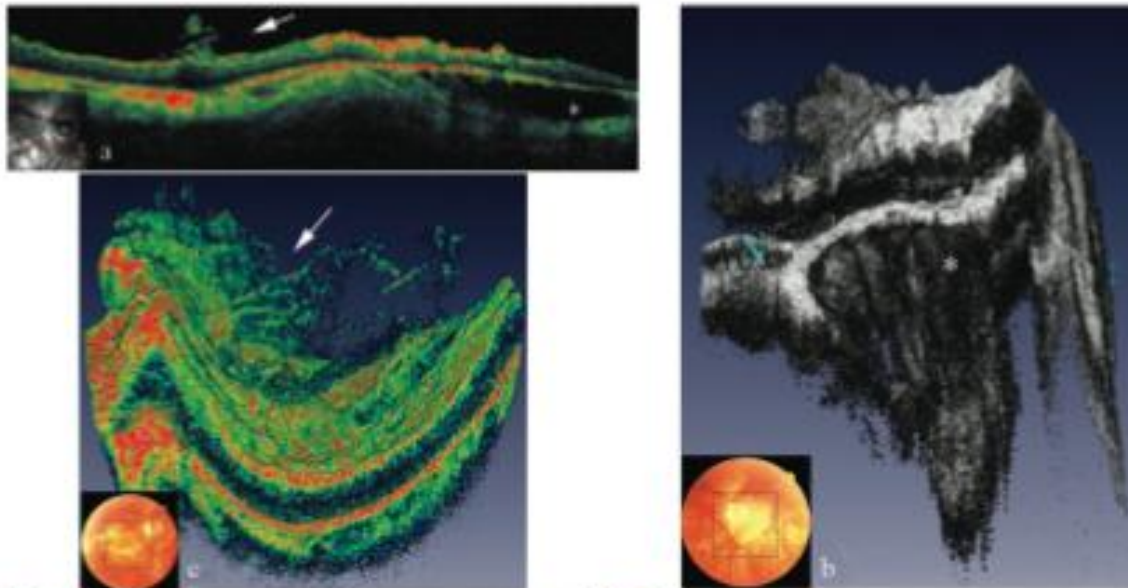
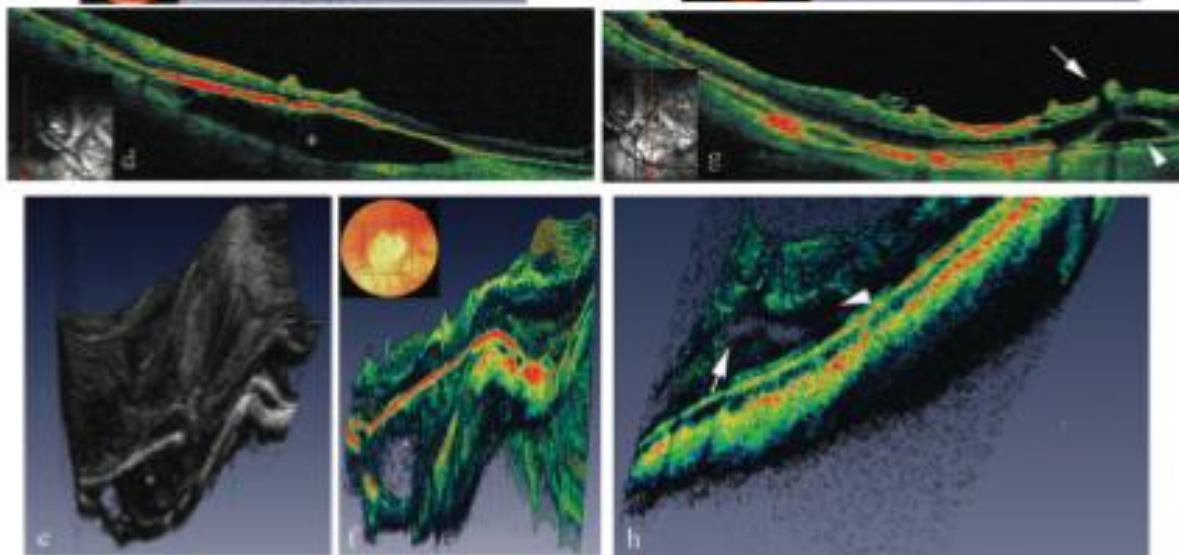


FIGURE 4. Right eye of a 70-year-old woman, with refraction of  $-26$  diopters. Best-corrected visual acuity of the right eye was  $20/400$ , and the axial length was  $28.5$  mm. (Left) The fundus has retinochoroidal atrophy within the staphyloma. The arrow indicates the area of the optical coherence tomographic scan. (Right) Optical coherence tomography shows a localized retinal detachment at the fovea. The detached retina has retinoschisis. A columnar structure bridges the inner and outer retinal layers.

Vitreo  
retinal  
traction



Peri  
Papillary  
detachment



**Figure 1.** Case 1, right eye. Peripapillary detachment of the pigment epithelium (asterisk) and vitreo-retinal traction (arrow) in a highly myopic eye, as imaged at both (a) cross-sectional B-scan and (b, c) three-dimensional spectral domain OCT (SD-OCT) views with different degrees of rotation. Case 1, left eye. Peripapillary detachment of the pigment epithelium (asterisk) in high myopia imaged with (d) cross-sectional B-scan and (e, f) three-dimensional SD-OCT visualization. An optically empty space is located beneath the retinal pigment epithelium (RPE). In the macular area, a small retinal detachment (arrowhead) and a cleavage of the neuroepithelium (arrow) are present at both (g) longitudinal OCT scan and (h) three-dimensional SD-OCT.

# Potential blinding disease in myopia

## CNVM:

- 5-10% of high myopes

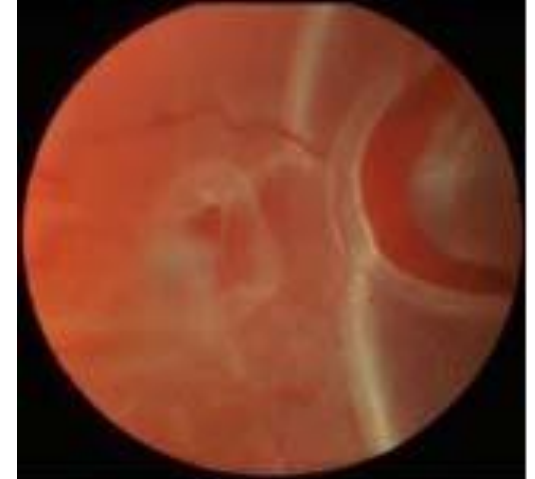
## Vitreoretinal:

- Floaters
- Earlier PVD
- Lattice
- Retinal tears/ detachment

# Daily issues

## Worrying signs / symptoms

- Risk of retinal detachment
- Foster- Fuch's spot    ?*VEGF* drugs
- Symptomatic floaters
- Difficulty of scleral indentation



Retinal Tear with  
Detachment

***Be quick to refer to a retina specialist***

***Delayed treatment can compromise outcome***



# Medico Legal obligations of an optometrist to detect retinal detachment & pre- detachment disease 1

- WCF in 20's
- Previous low myope : PRK. Re-treatment.
- Retina examined by 2 ophthalmologists associated with PRK
- Consulted optometrist ~2y later with decreased VA : diagnosed CSR, missed retinal detachment

# Medico Legal obligations 2

- One opinion: Optometrist NOT liable for missing ret det.
- NOT part of optometry competency

# ASSOCIATIONS OF HIGH MYOPIA IN CHILDREN

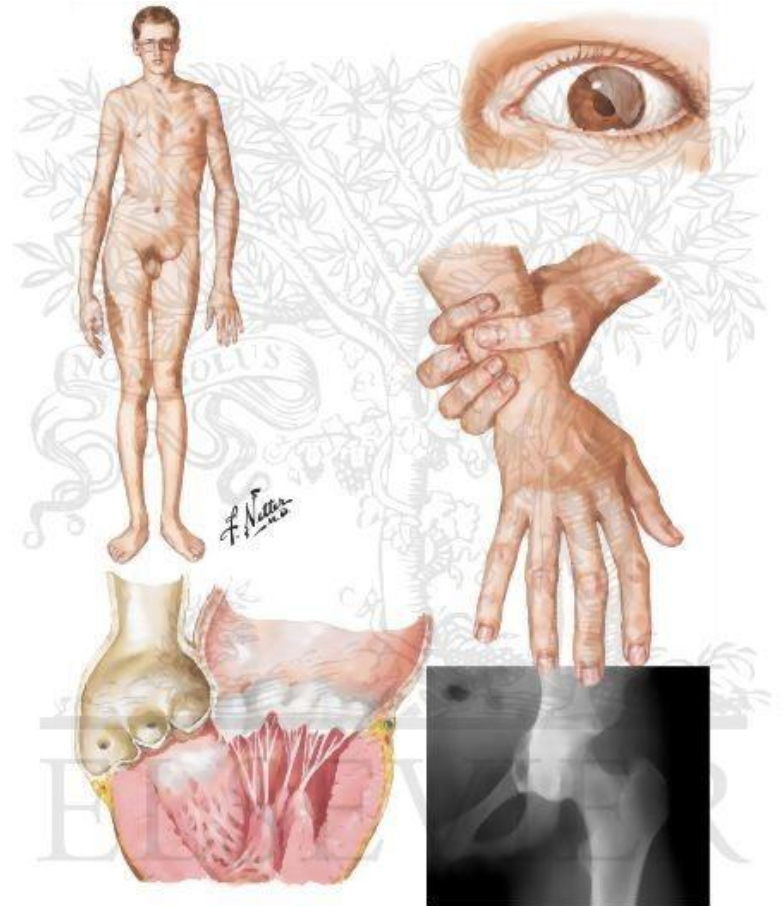
Marr...Ainsworth Eye, 2001 **Birmingham**

- N=112. <10yo.  $\geq -6$ DS.  $\geq 12$  mo followup
- ERG if reduced BCVA, nystagmus, ?night blindness
- M sl > F
- 58% Caucasian. 38% Asian [Birmingham 15%]
- Only 10 referred from optometry
- 89% bilateral high myope. 11%: unilateral
- 32% aniso  $\geq 2$ DS
- No spontaneous decrease in myopia

# Family history

## Marr...Ainsworth

- Eye problem 44%
- 29% myopia
- 10: **Marfan**, juvenile cataract, high myopia, Stickler synd, nyctalopia
- 4: Marfan or Stickler subsequently diagnosed in other family members



# ASSOCIATIONS

Marr...Ainsworth

- **8% 'simple high myopia'**
- **54% systemic association**
- **56% orthoptic problem [amblyopia, strab, nyst]**
- **34% ocular abnormality**

# Ocular abnormality

## Marr...Ainsworth

- Anisometropic amblyopia 32%
- Strabismus 18%
- Nystagmus 12%
- ROP stage  $\geq 3$ : 7%
- Retinal dystrophy 7% [cone dystrophy, CSNB, Stargardt's]
- Coloboma 5% + MGDAnomaly 1%
- Glaucoma / Oc Ht 3%
- Cataract 4%
- Subluxed lens 4%
- Albinism 2%
- Microphthalmos, aniridia, spherophakia, post lenticonus, persistent pupillary membrane, traumatic lenticonus....all 1%

# Systemic associations

Marr...Ainsworth

- Severe devptl delay 12%
- Extreme prematurity 10%
- Stickler's 8%
- Down's 5%
- Marfan's 5%



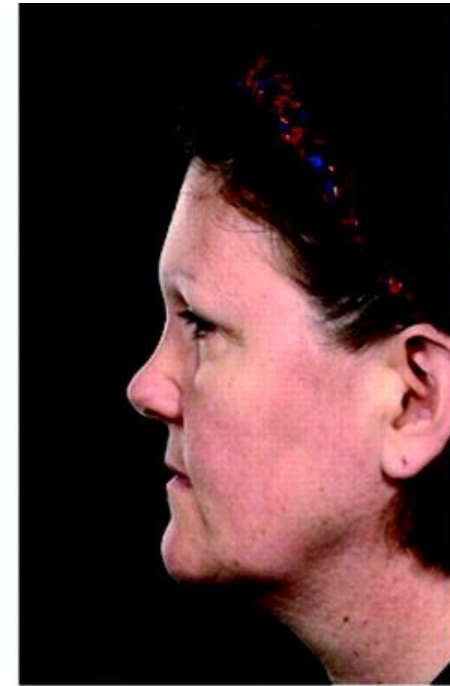
# Stickler's

## Eye

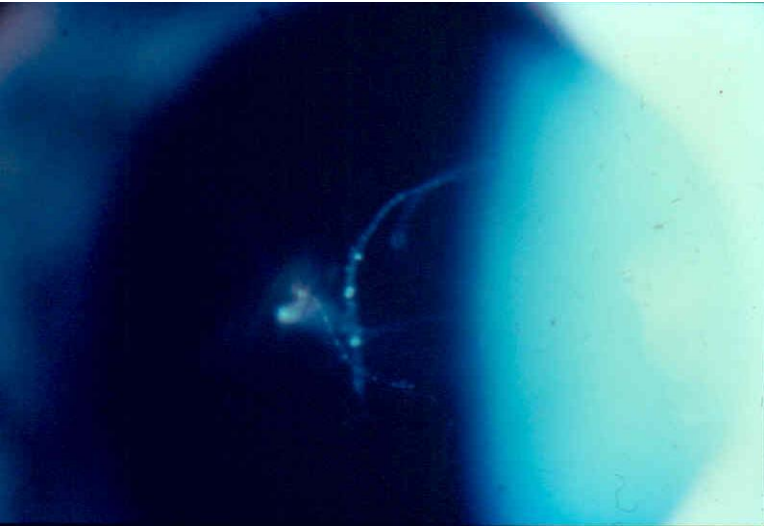
- **Myopia**
- **POAG**
- **Cataract**
- **Vitreoretinal changes**  
predisposing to ret det

## Face

- Midfacial flattening
- Small chin
- Cleft palate



# Stickler's

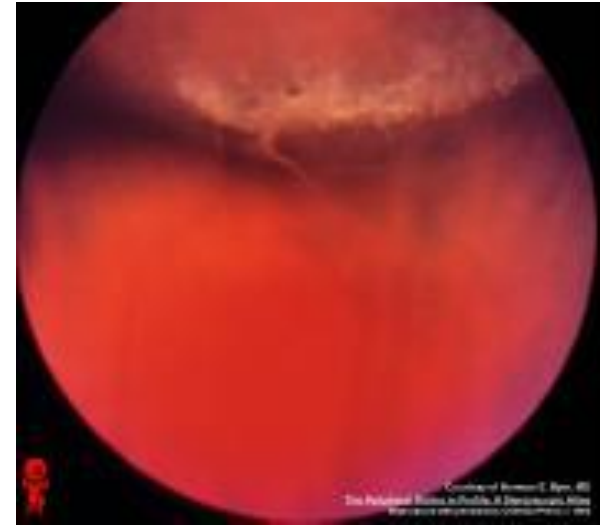


## SEMINAL

1. Empty vitreous

OR

2. Abn beaded vitreous



Lattice

'Snail track'

Increased V-R adhesion @  
edges

Retinal thinning

# Stickler's

- Very underdiagnosed
- Commonest cause of inherited retinal detachment
- Av age @ diagnosis: Child 4, Adult 32

# UNILATERAL HIGH MYOPIA

## Weiss BJO 2003

- N=48. mean age 7y [4m to 17y]
- Mean anisomyopia -  $9 \pm 4$  DS
- 16 ET, 11 XT
- 30% abn optic nerve [hypoplasia, myel nerve fibres, atrophy, coloboma]
- 21% abn CNS
- 12% abn lens
- 10% ROP
- 6% FH high myopia
- 6%: NO associated factors

# Myopia- associated esotropia

- 2 types:
- 1. Bielschowsky.

Antedates modern diagnostic tests

- 2. 'Heavy Eye'.

# *Case 1: Eye is turned IN & DOWN*

*Preoperative*

## *Looks 'Heavy' hence Heavy Eye*



*Postoperative (52 days after surgery)*



# *Definition of Progressive Esotropia Caused by High Myopia*

- *Presence of high myopia with an axial length sometimes greater than 30 mm.*
- *Abduction and elevation are limited*



# Case 2

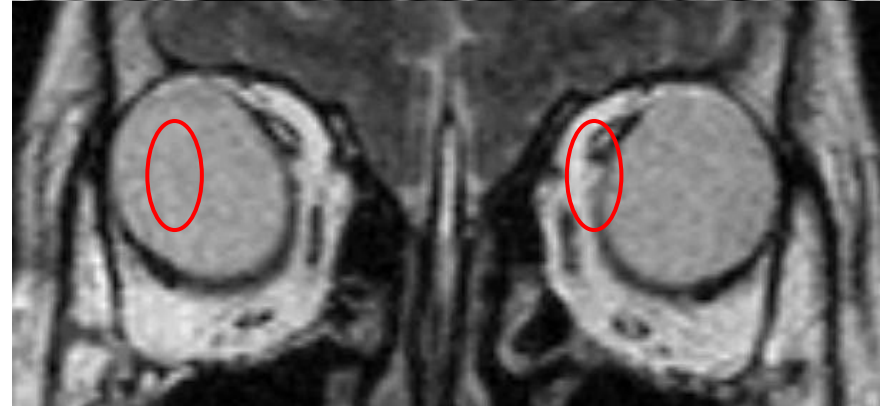
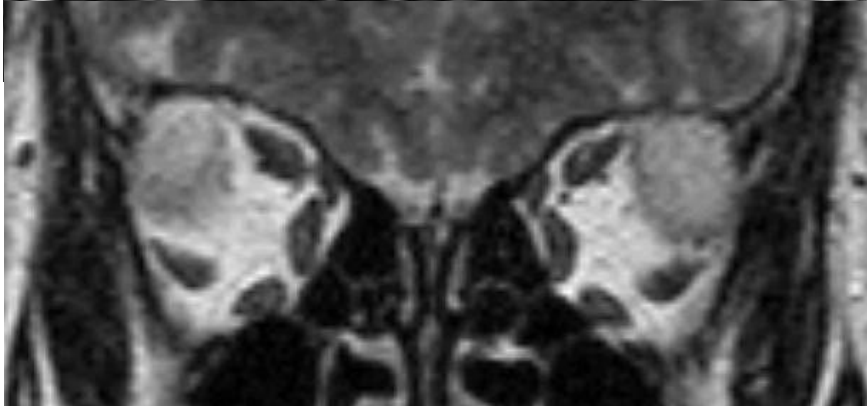
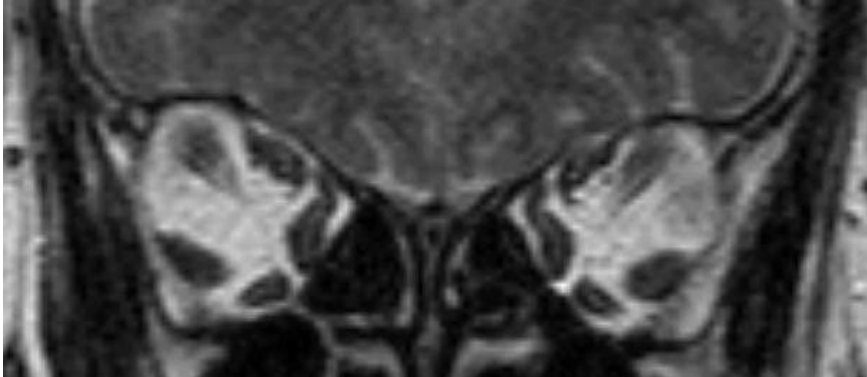
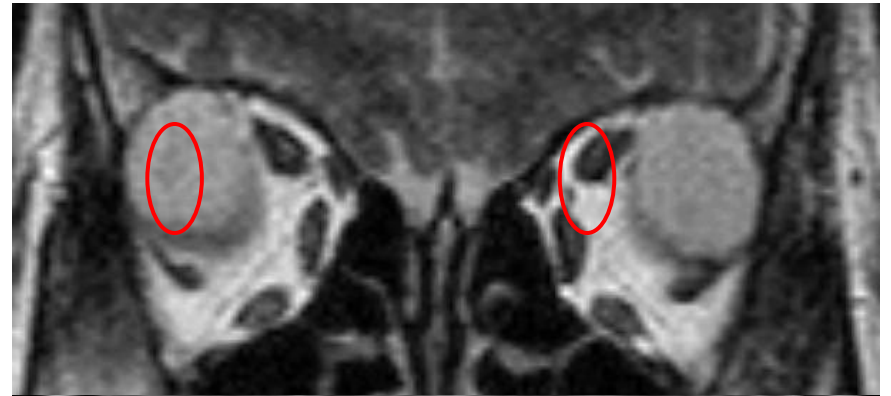
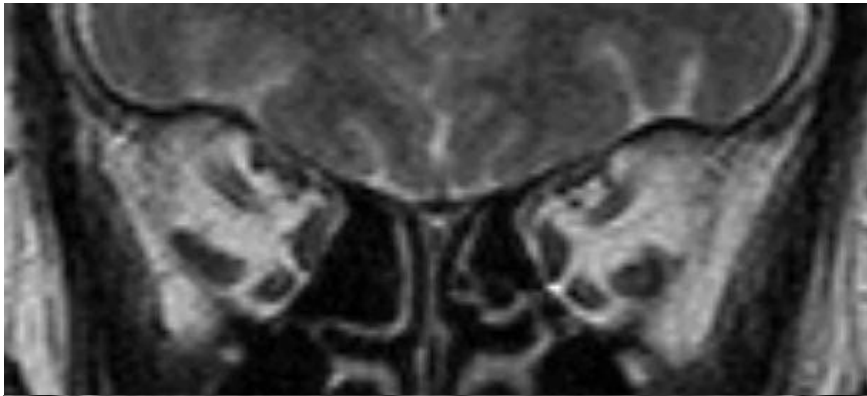
*Preoperative*



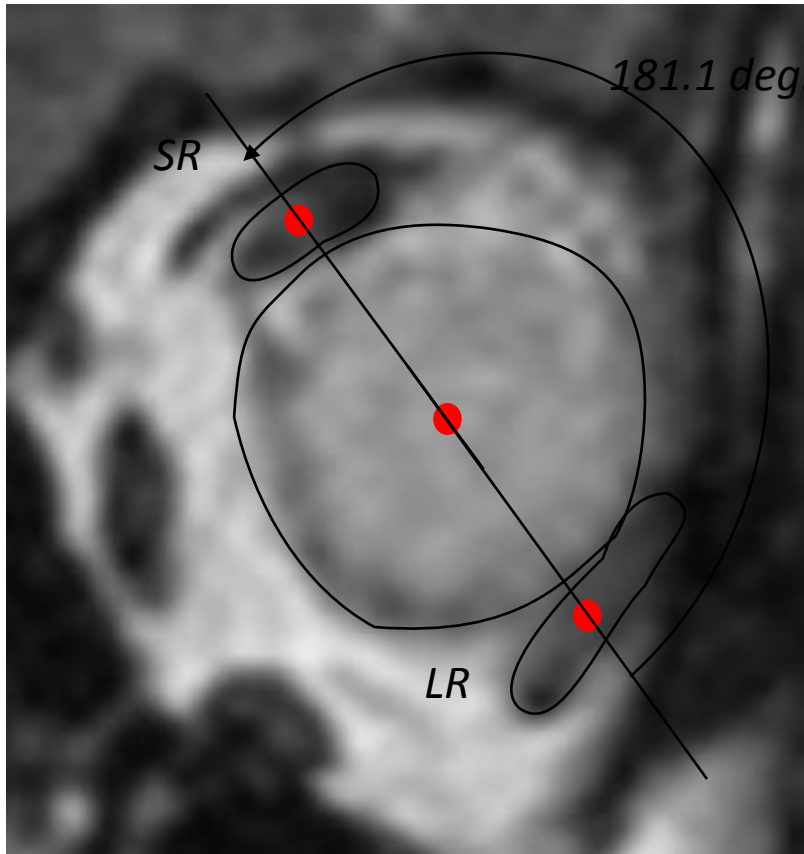
*Postoperative OS (69 days after surgery)*



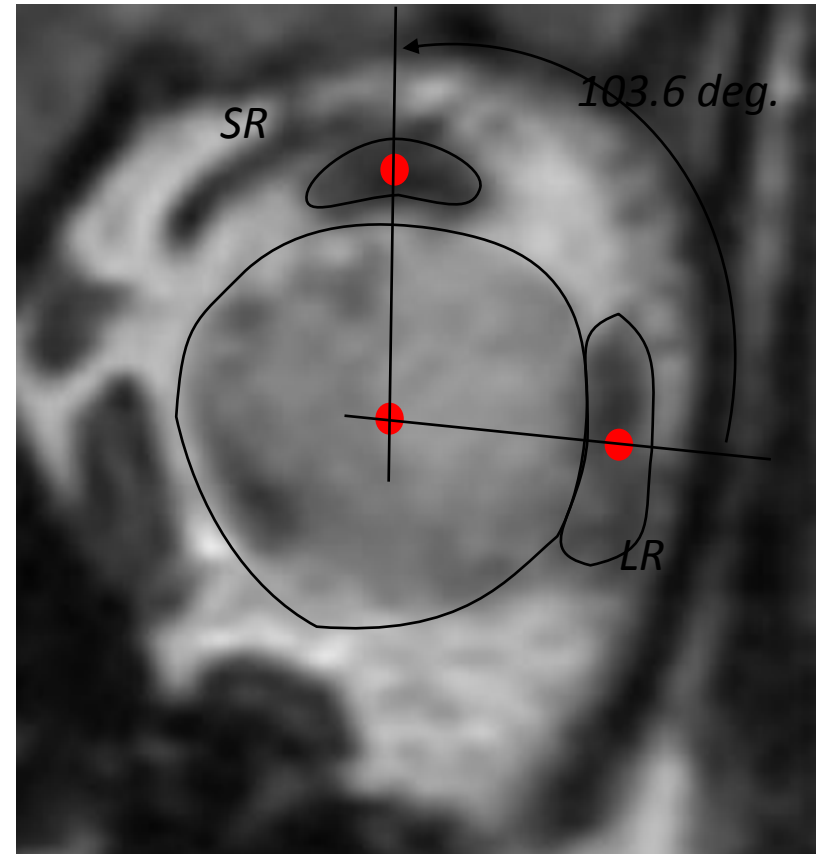
# *Coronal MRI Scans of Case 2*



# Measuring the Angle of Dislocation of the Eyeball



*Preoperative*

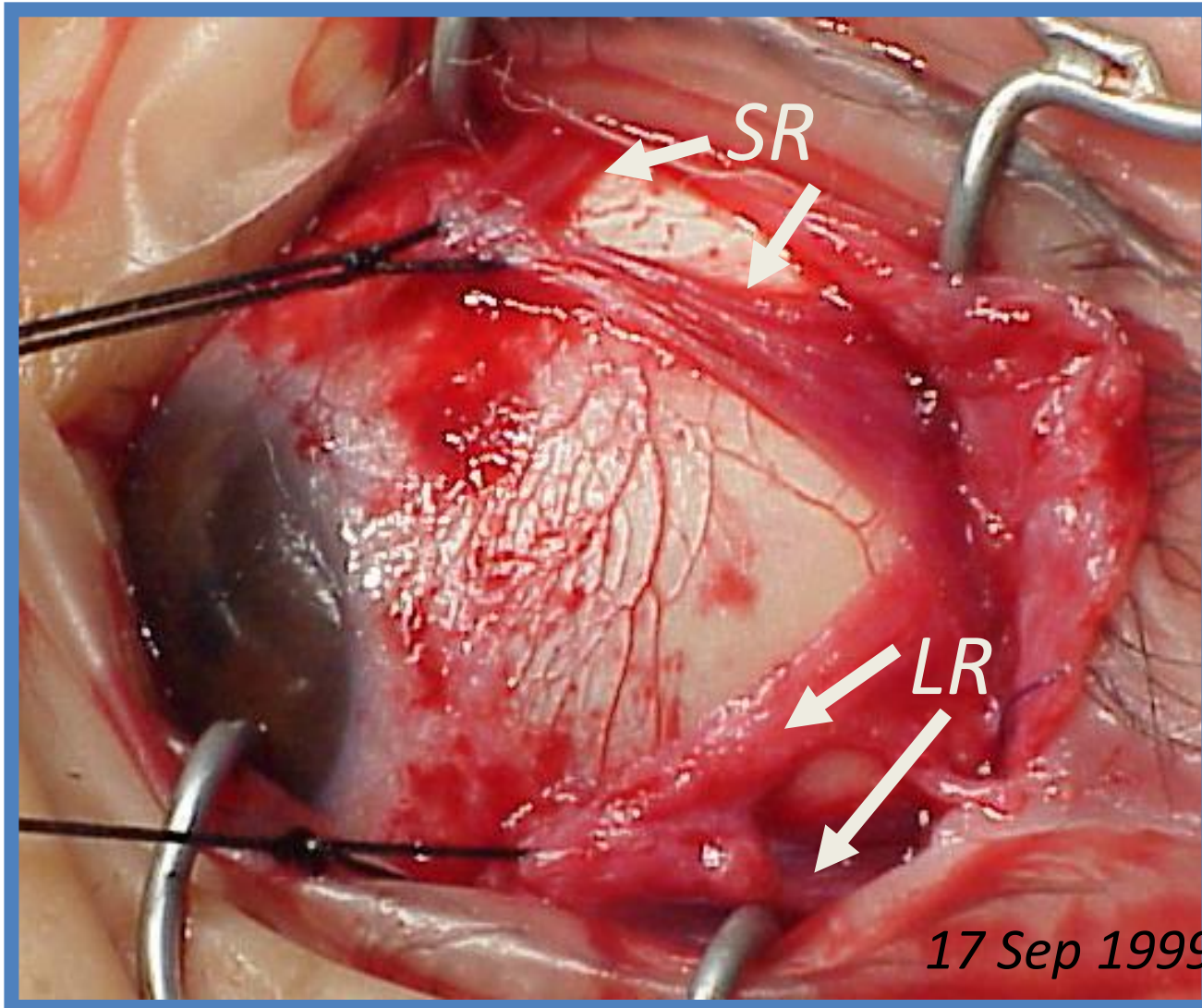


*Postoperative*

*The center positions were measured with Scion Image® software.*



*Joining the SR and LR  
After Splitting (into halves)*



17 Sep 1999

Pathological associations

Thank you