



ARGOMENTI DI NEUROFTALMOLOGIA LE NEURITI OTTICHE

Oftalmologo vs Neurologo

Cona, 29 settembre 2018

Dott.ssa Giuseppe Lamberti

U.O. Oculistica



U.O. OCULISTICA
DIR. : PROF. PAOLO PERRI



UNIVERSITÀ
DEGLI STUDI
DI FERRARA
- EX LABORE FRUCTUS -

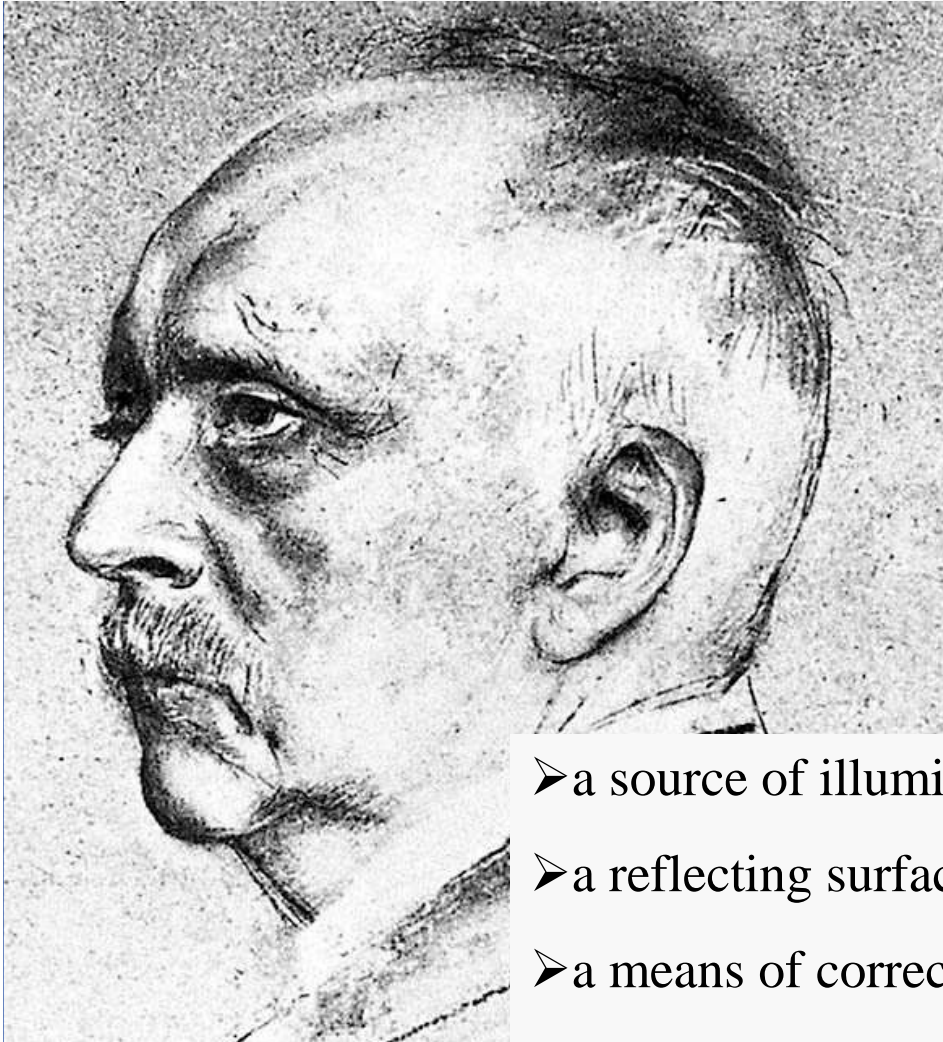
PICCOLI PROBLEMI ALL'OCCHIO CAUSANO GRANDI PROBLEMI AL PAZIENTE

I DISTURBI OCULARI POSSONO ESSERE SEGNI DI PATOLOGIE SISTEMICHE

L'oculista dovrebbe inviare al
neurologo tutti i casi di sospetta N.O.

- *Corretta diagnosi differenziale*
- *Trattamento*
- *Monitoraggio a lungo termine*

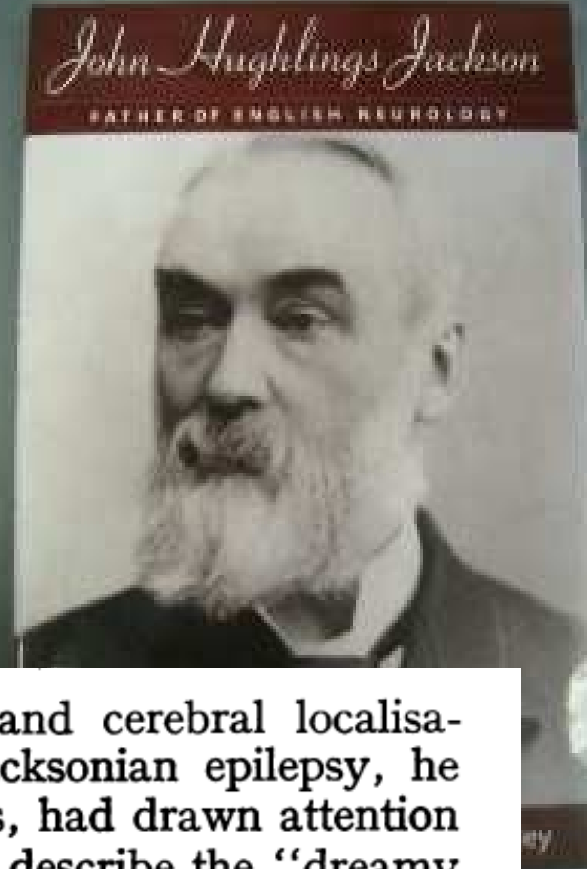
**1851: viene risolto il mistero di come
osservare l'interno dell'occhio
passando attraverso il foro nero della pupilla**



- a source of illumination,
- a reflecting surface to direct light toward the eye, and
- a means of correcting an out-of-focus image on the fundus.

Hermann von Helmholtz

Non è un nervo
nell'accezione abituale del
termine, *morfologicamente*
e funzionalmente è una
proiezione diencefalica

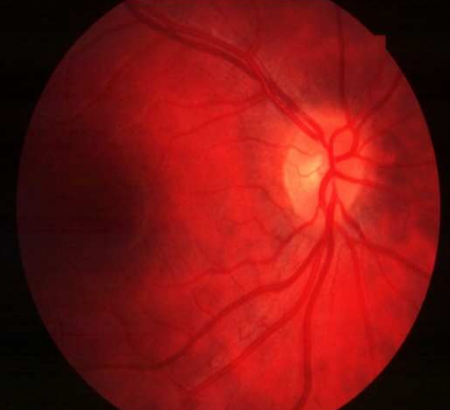


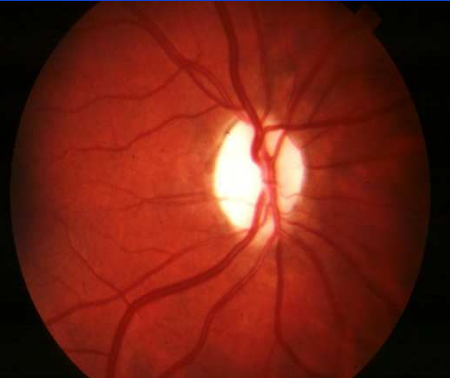


Though his publications on epileptiform convulsions and cerebral localisation soon led to this form of epilepsy being known as Jacksonian epilepsy, he never ceased to point out that the French physician, Bravais, had drawn attention to the type forty years previously. Jackson was the first to describe the “dreamy state” and olfactory aura associated with epileptic attacks in tumours of the temporal lobe, named by him “uncinate epilepsy.”

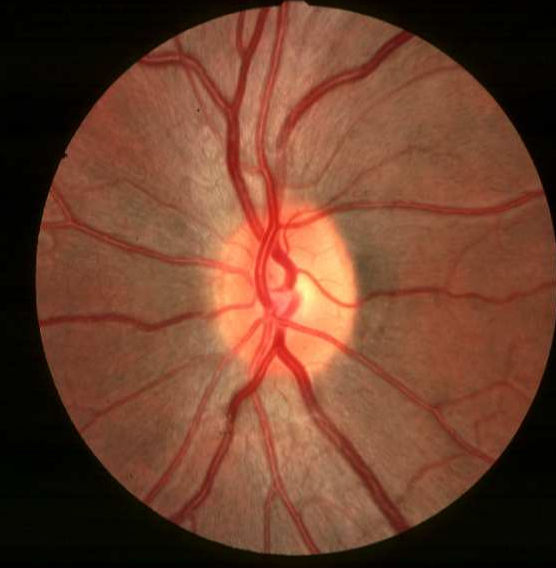
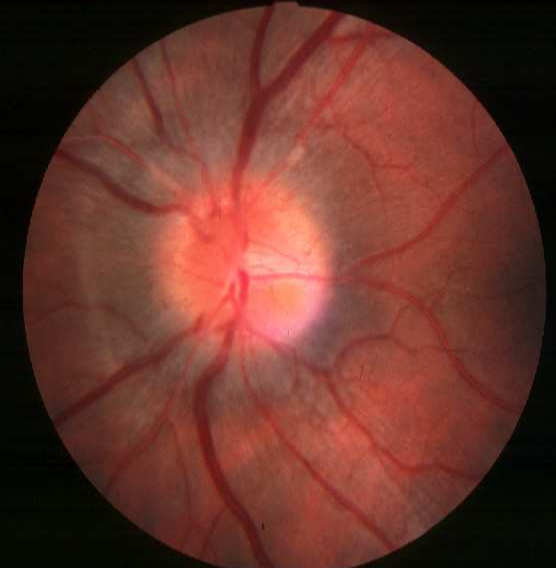

“The ophthalmoscope, as an aid to the study of diseases of the brain” .

Med Times Gaz, 1862, ii: 598–601 12 Dec 62

Optic disc changes

Normal	Swelling
	
<ul style="list-style-type: none"> • Retrobulbar neuritis • Early compression 	<ul style="list-style-type: none"> • Papilloedema • Papillitis and neuroretinitis • AION
Optico-ciliary shunts	Atrophy
	
<ul style="list-style-type: none"> • Optic nerve sheath meningioma • Occasionally optic nerve glioma 	<ul style="list-style-type: none"> • Postneuritic • Compression • Hereditary optic atrophies

Classification of optic neuritis

Retrobulbar neuritis (normal disc)	Papillitis (hyperaemia and oedema)	Neuroretinitis (papillitis and macular star)
		
<ul style="list-style-type: none">• Demyelination - most common• Sinus-related (ethmoiditis)• Lyme disease	<ul style="list-style-type: none">• Viral infections and immunization in children (bilateral)• Demyelination (uncommon)• Syphilis	<ul style="list-style-type: none">• Cat-scratch fever• Lyme disease• Syphilis

Neurite ottica

SEMEIOTICA LOW-TECH

SOSPETTO

AMBULATORIALE

Monoculare

Riduzione anche sfumata

AV

Dolore (movimenti oculari)

Discromatopsia

SEMEIOTICA EPONIMICA

Uhthoff's *symptom*

Flashes (Lhermitte)

Pulfrich *phenomenon*

Alterazioni del CV

(Seidel; Bjerrum)

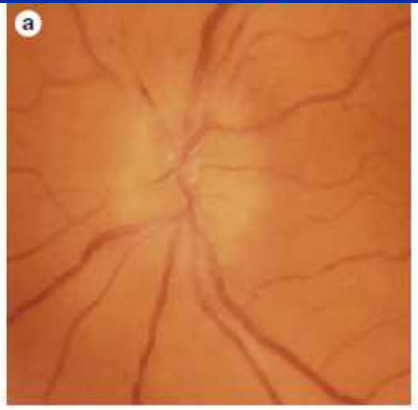
Pupilla di Marcus Gunn

WILHELM UHTHOFF



09. Uthoff

- n Wilhelm Uthoff (1853-1927)
- n *Eye Symptoms in Diseases of the Nervous System (Published 1915)*
- n Described by Bielschowsky as the “true originator” of clinical neuro-ophthalmology



Il sintomo di Uthoff

Uthoff descriveva tre pazienti in cui la fatica fisica causava una discromatopsia

Il paziente XVIII aveva un calo del visus dopo aver passeggiato intorno all'ambulatorio

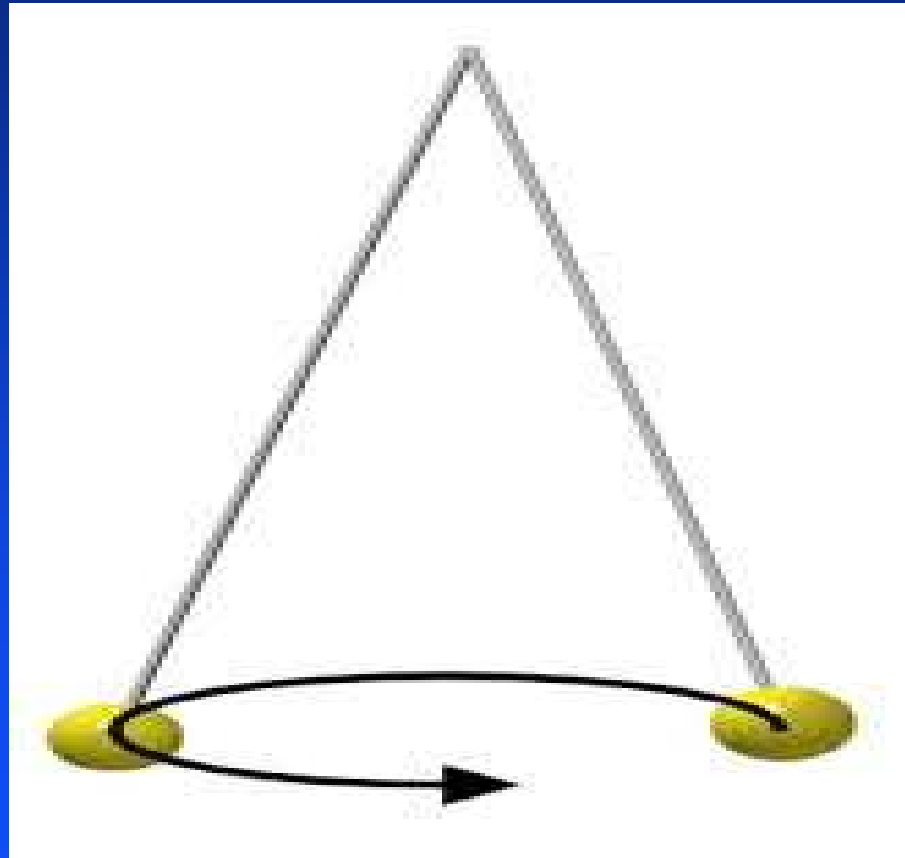


UTHOFF

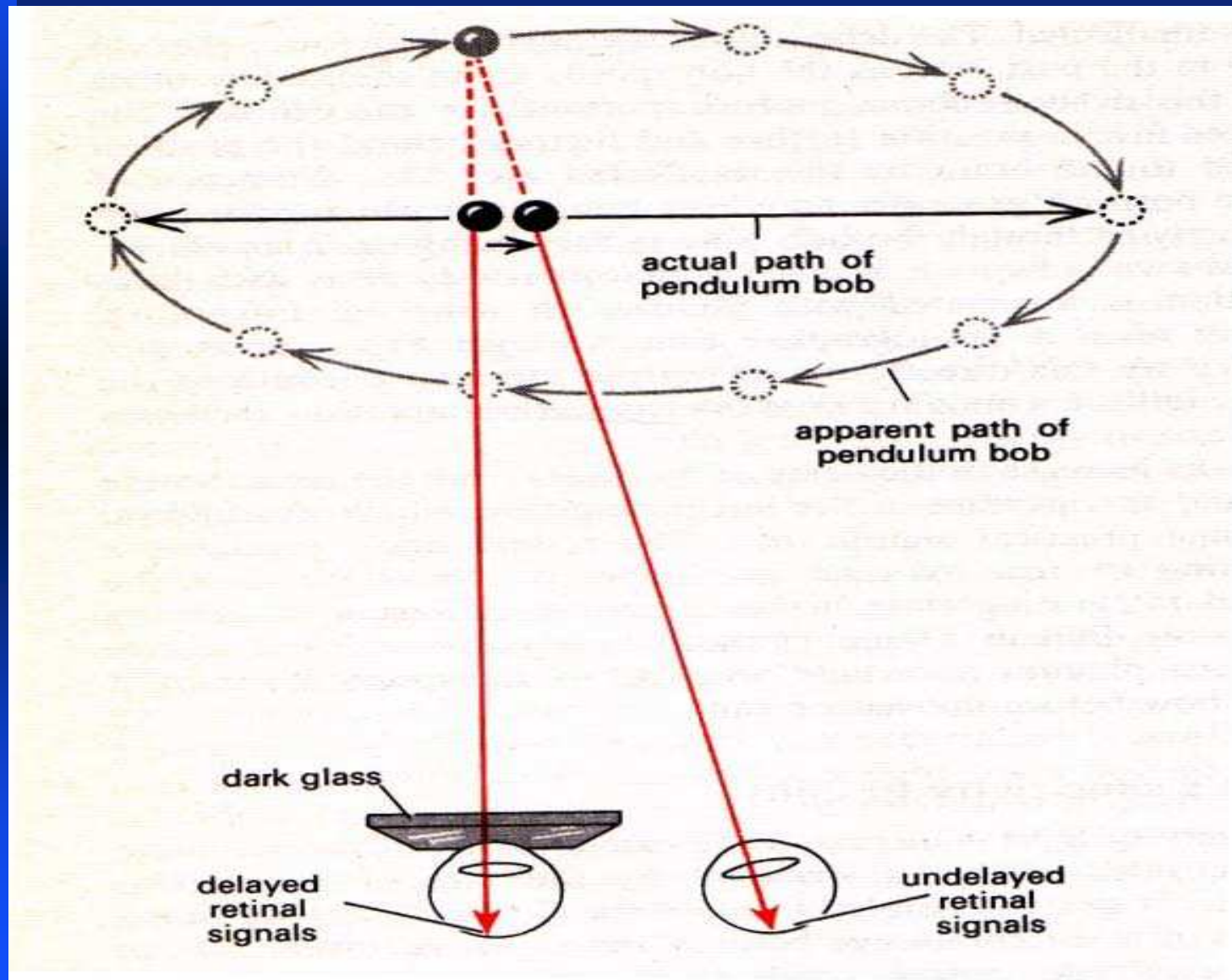
Uhthoff's Symptom

- n *Uhthoff's symptom in optic neuritis: relationship to MRI and development of MS.* (Scholl GB, Song HS, Wray SH) *Ann Neurol* 1991; 30(2):180-4
- n *Uhthoff and his Symptom* (Selhorst JB, Saul RF) *Journal of Neuro-ophthalmology* 1995; 15(2):63-9

PULFRICH



Pulfrich Phenomenon



Flashes (scotomi scintillanti)

Movement phosphenes in optic neuritis: A new clinical sign (Davis F, Bergen D, Schauf C, McDonald I, Deutsch W)
Neurology 1976; 26: 1100-1104.

bagliori luminosi al buio accentuati dai movimenti oculari

dx differenziale con fosfeni trazionali

rappresentano l'equivalente oftalmologico dei sintomi di Lhermitte

RIFLESSO FOTOMOTORE

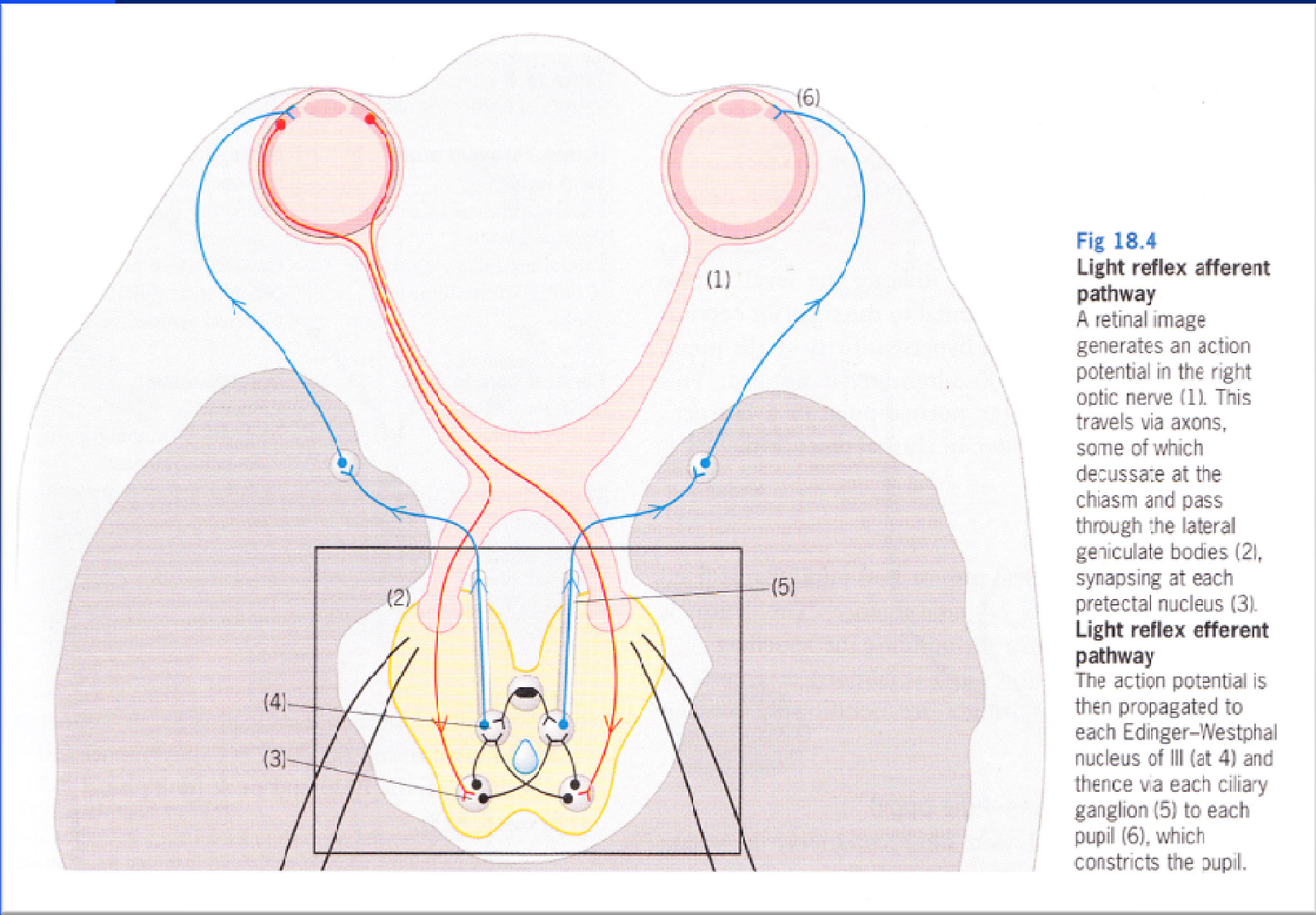


Fig 18.4

Light reflex afferent pathway

A retinal image generates an action potential in the right optic nerve (1). This travels via axons, some of which decussate at the chiasm and pass through the lateral geniculate bodies (2), synapsing at each pretectal nucleus (3).

Light reflex efferent pathway

The action potential is then propagated to each Edinger-Westphal nucleus of III (at 4) and thence via each ciliary ganglion (5) to each pupil (6), which constricts the pupil.

RELATIVE AFFERENT PUPILLARY DEFECT

- n Incomplete damage to AFFERENT pathway
 - u Partial retina or optic nerve damage
- n For LEFT RAPD
 - u Light into left eye: left and right pupil constrict
 - u Light into right eye: both pupils constrict further
 - u Back to left eye: **both pupils dilate, but not completely**
 - u Light away: both pupils dilate completely

The abnormally Reacting Pupil

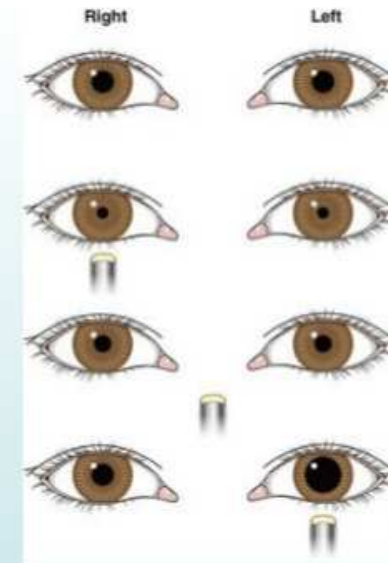
2- Swinging flashlight test

when the pupil exhibits an RAPD, it is described as a Marcus Gunn pupil. It suggests:

- Optic nerve disease, central retinal artery or vein, A mild RAPD may also occur in amblyopia .

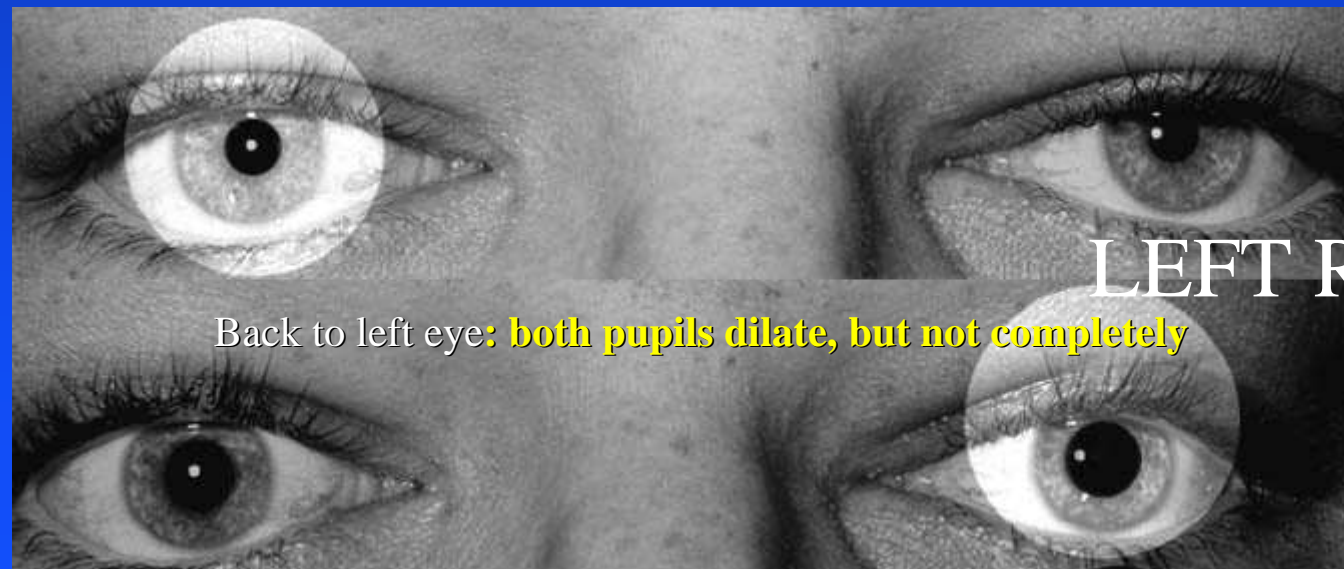


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Marcus Gunn pupil

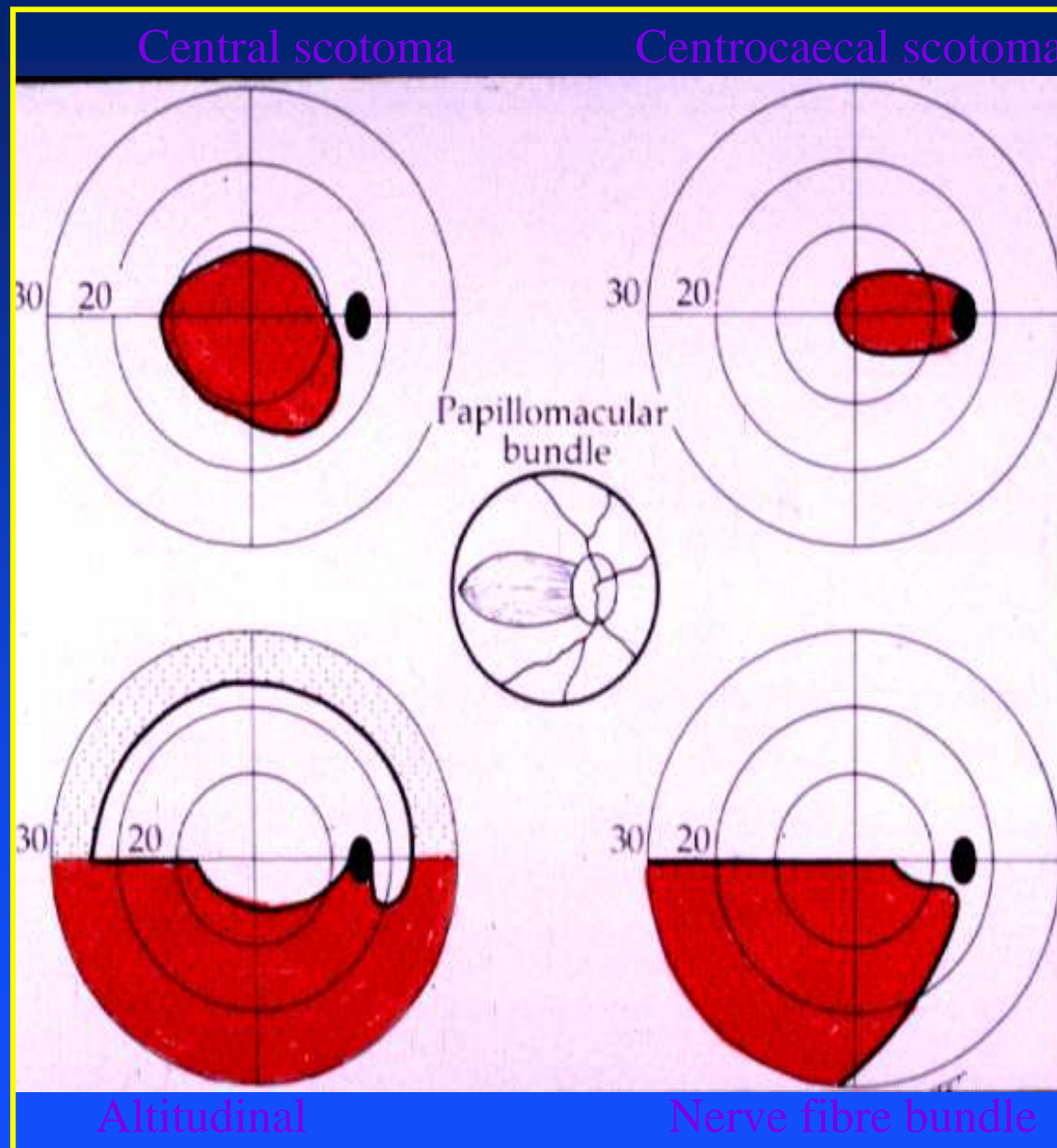
relative afferent pupillary defect caused by optic nerve lesion or extensive retinal disease



LEFT RAPD

Back to left eye: both pupils dilate, but not completely

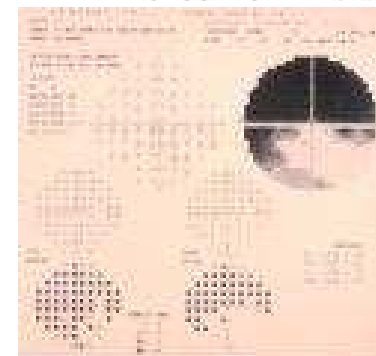
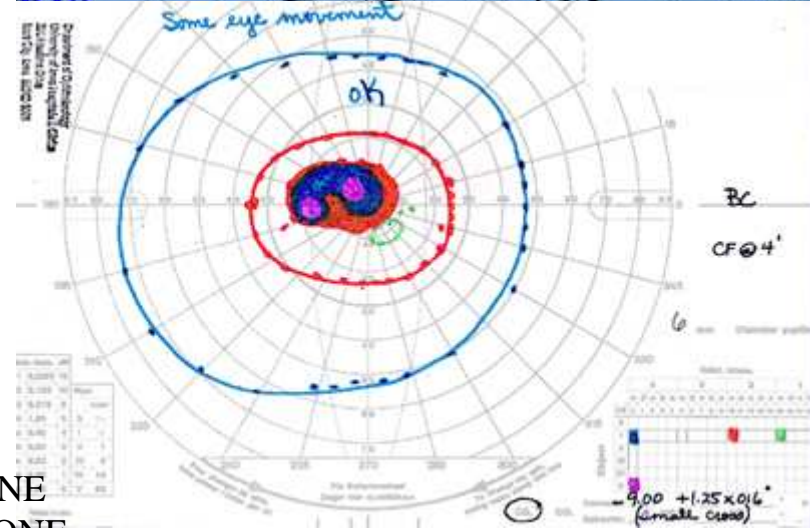
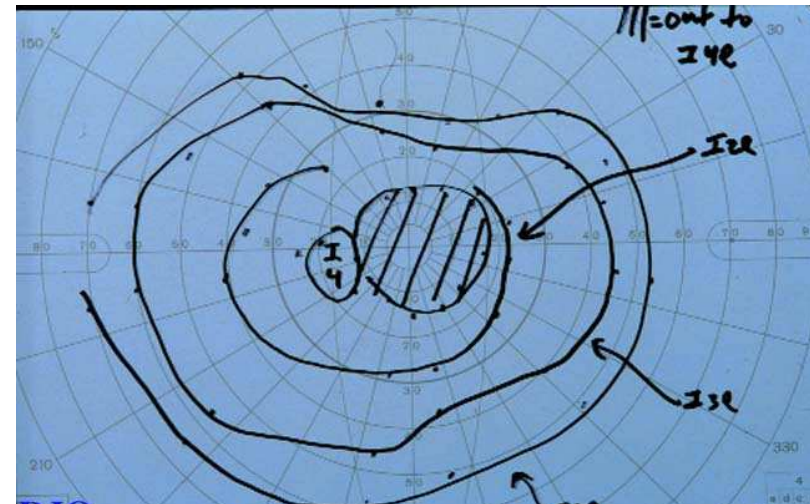
Visual field defects



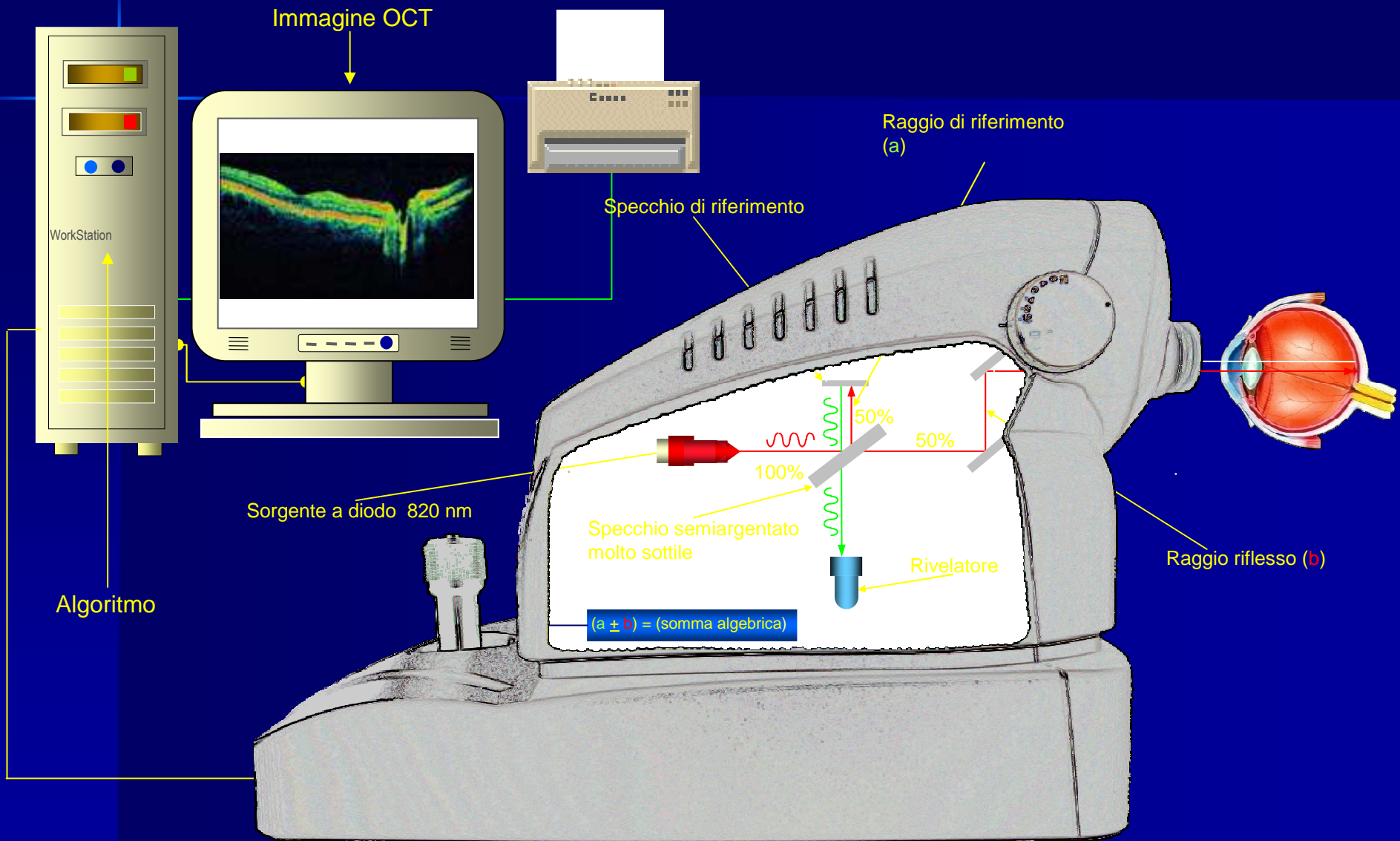
Visual Fields

- Central scotomas
- Paracentral scotomas
- Altitudinal defects

L'OCT FORNISCE IMMAGINI AD ALTA RISOLUZIONE NELL'ORDINE DEI MICRON SOTTO FORMA DI SEZIONE TRASVERSALE DETTA TOMOGRAFICA DELLE MICROSTRUTTURE DEI TESSUTI BIOLOGICI TRAMITE LA MISURA DEL RITARDO O DELLA VARIAZIONE SPETTRALE DELLA LUCE RIFLESSA IN RETRODIFFUSIONE



OCT (Optical Coherence Tomography)



Prognosis

- Severity of initial visual loss is related to final visual outcome
- Most recover well
 - 74% \geq 20/20
 - 92% \geq 20/40



Optic neuritis presenting with amaurosis fugax

Amer M. Awad · Bachir Estephan ·
Worthy Warnack · Olaf Stüve

J Neurol (2009) 256:2100–2103

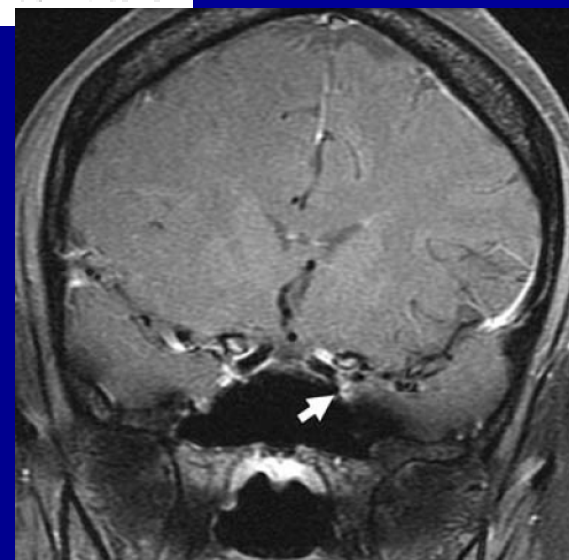
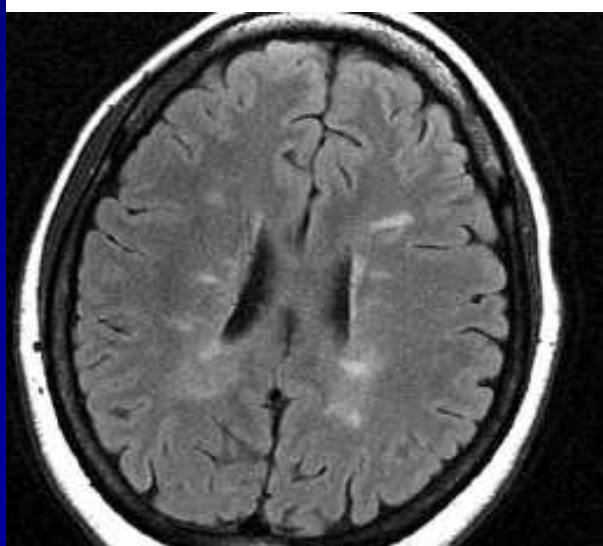
The onset of visual loss is relatively acute and progresses over hours or days [7]. It is associated with intra- or retro-orbital pain on the affected side in 92.2% of the cases [3]. Pain may precede visual loss and lasts for days to weeks [3]. Visual function usually begins to improve rapidly after a median of 2 weeks, and resolution continues over several months [4]. Complete recovery of visual acuity is common, even after near blindness, and functional levels of sight following remission are the rule [7]. ON is most commonly associated with multiple sclerosis (MS).

Optic neuritis presenting with amaurosis fugax

Amer M. Awad · Bachir Estephan ·
Worthy Warnack · Olaf Stüve

J Neurol (2009) 256:2100–2103

The patient is a 39-year old right-handed previously healthy African American woman who presented to a tertiary referral center after experiencing ten episodes of painless vision loss. Over the course of 72 h, she exhibited sudden and complete loss of vision in the left eye lasting between 10 s and 45 min each. During the episodes of visual loss there was no light perception in the left eye. In the intervals



SOGLIA TONI DI GRIGIO

SOGLIA (DB)

DEVIAZIONE TOTALE

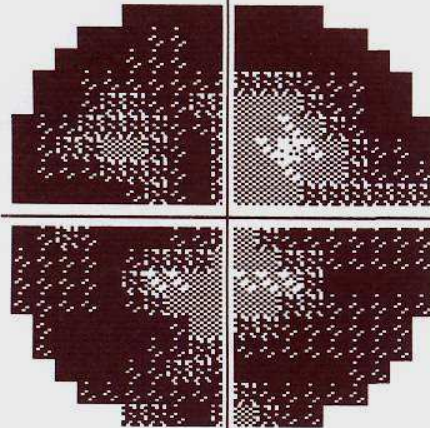
DEVIAZIONE DAL MODELLO

04-05-2011

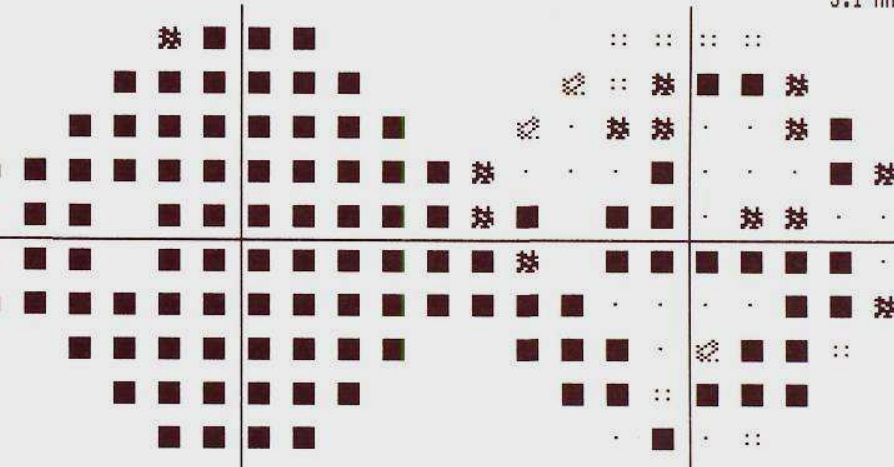
SITA-STANDARD

GHT: FUORI DAI LIMITI NORMALI

5.1 MM



<0	<0	<0	<0						
<0	3	0	<0	<0	<0				
0	7	4	6	13	13	4	<0		
0	10	11	10	<0	12	24	14	1	<0
0	<0	0	8	<0	13	11	9	9	6
<0	6	<0	<0	4	11	3	3	4	5
2	1	<0	19	14	16	16	<0	0	<0
<0	<0	<0	11	10	3	3	5		
<0	3	9	<0	0	<0				
8	<0	13	3						



FOVEA: 19 DB ■

FL: 0/18

FN: N/A

FP: 0%

MD: -24.82 DB P < 0.5%

PSD: 7.25 DB P < 0.5%

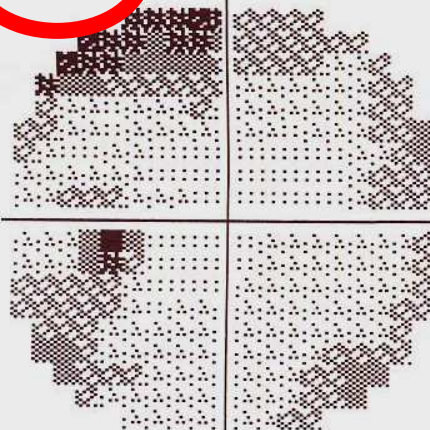
BOLO CORTISONICO

19-07-2011

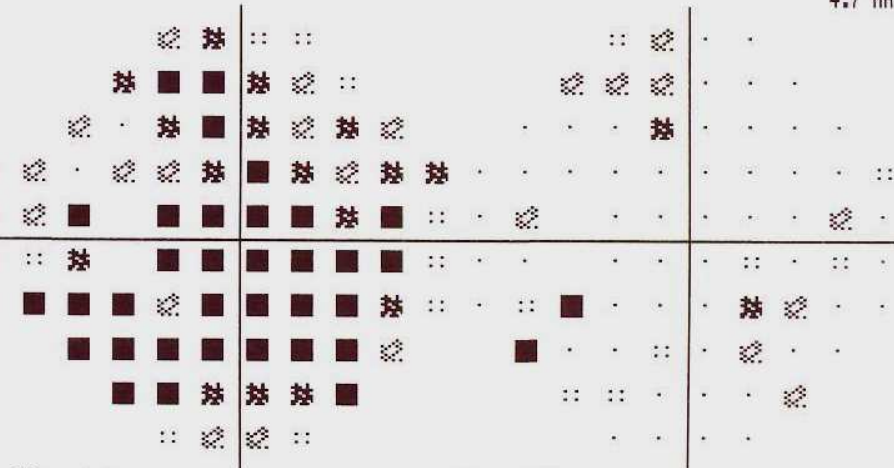
SITA-STANDARD

GHT: SENSIBILITÀ RIDOTTA

4.7 MM



9	4	17	18						
9	14	14	19	20	21				
21	26	23	19	24	25	21	20		
20	27	25	26	26	26	27	26	21	14
23	19	19	25	26	27	26	27	19	20
25	23	<0	26	27	26	24	24	21	20
20	20	17	27	26	25	23	22	23	19
16	23	24	22	24	21	23	21		
20	21	23	23	21	16				
24	21	21	20						



FOVEA: 31 DB ■

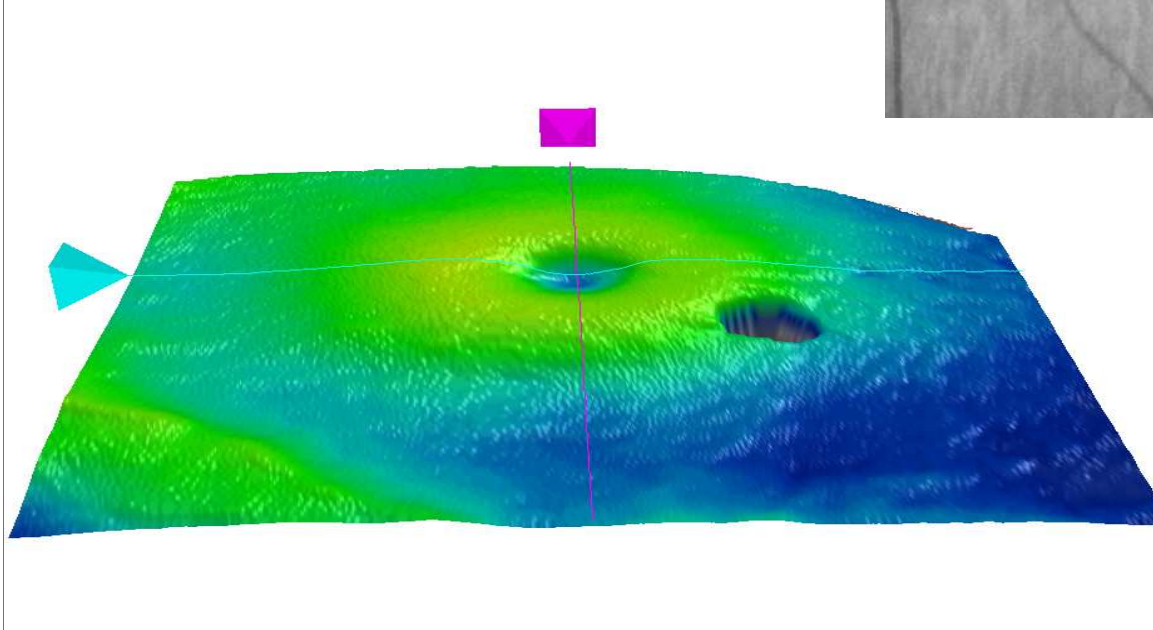
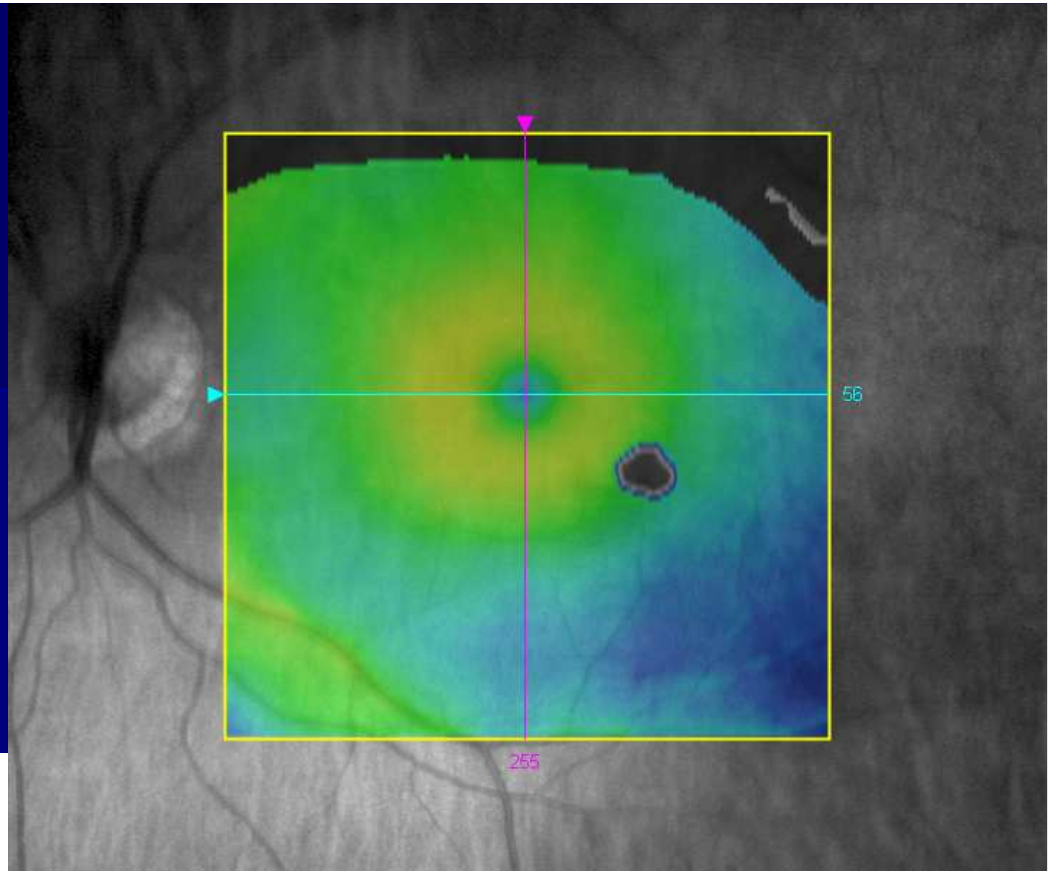
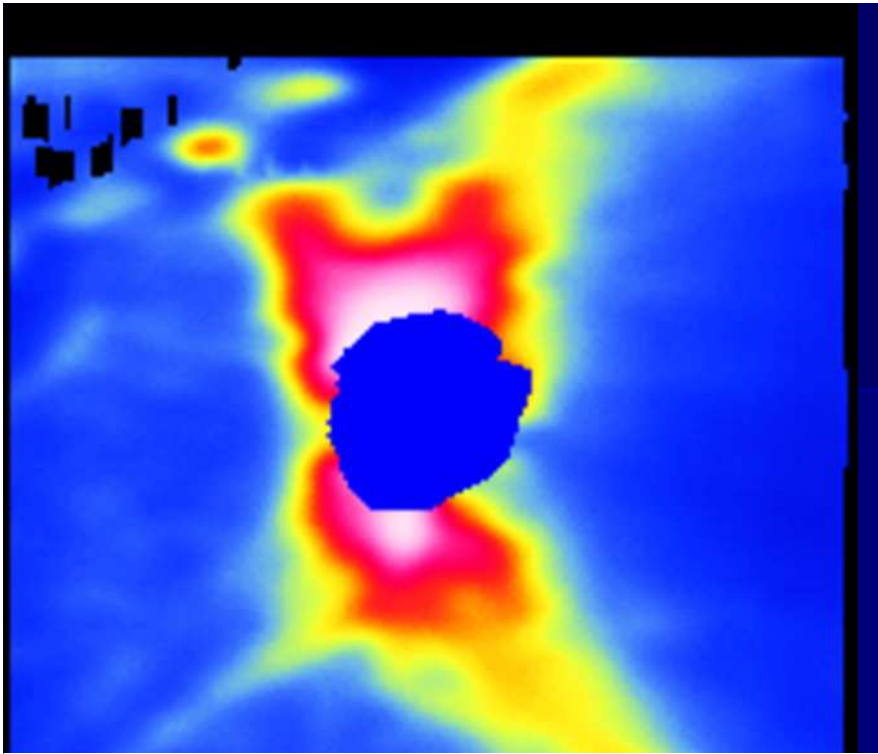
FL: 0/19

FN: 1%

FP: 0%

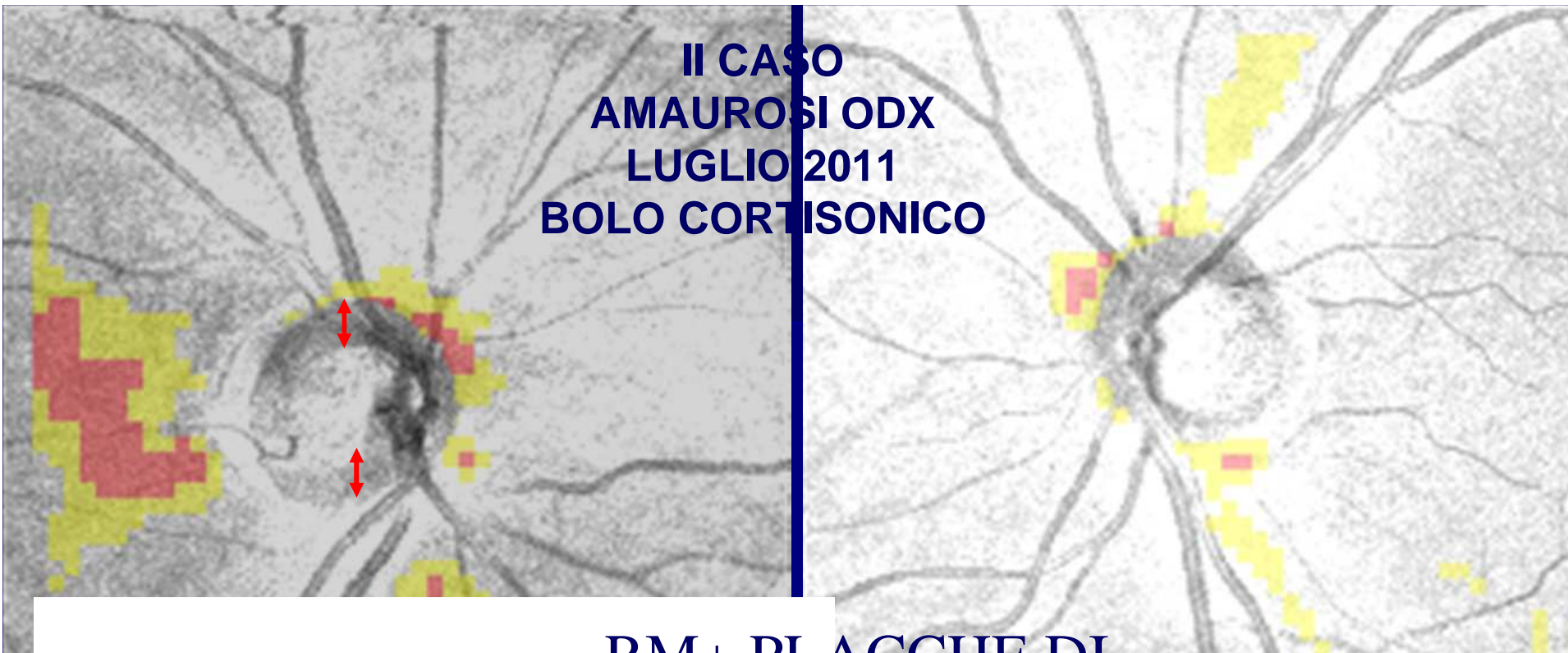
MD: -7.52 DB P < 0.5%

PSD: 9.04 DB P < 0.5%

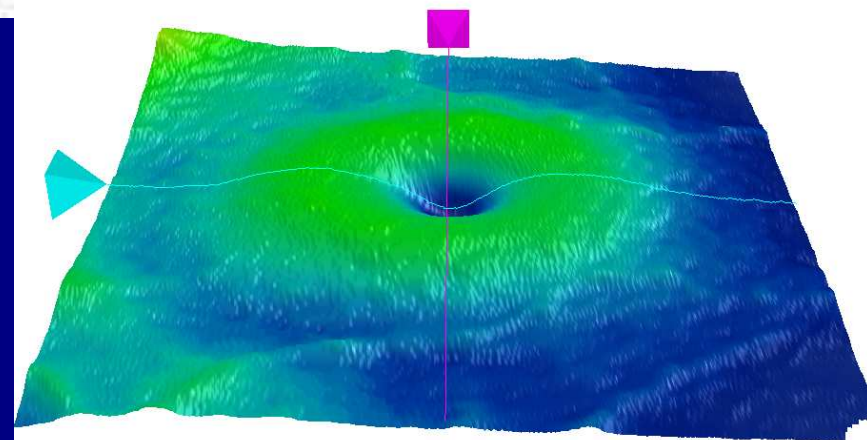
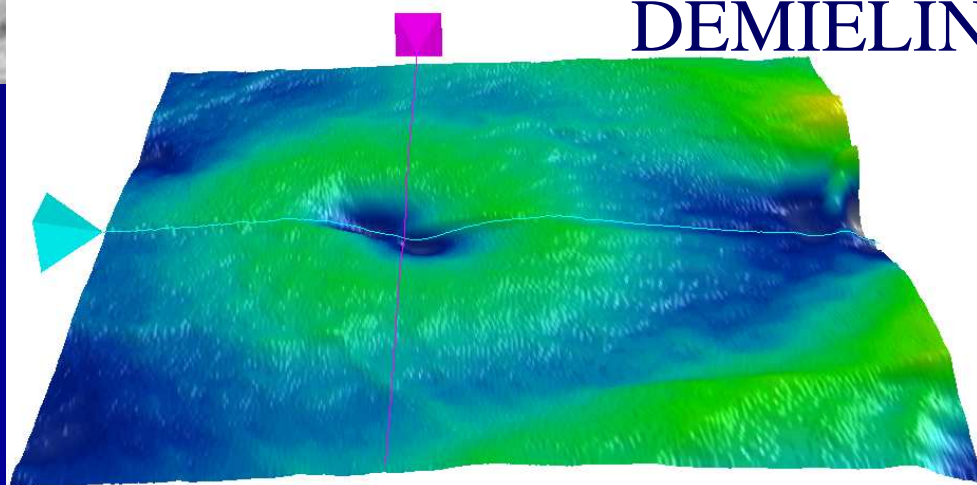


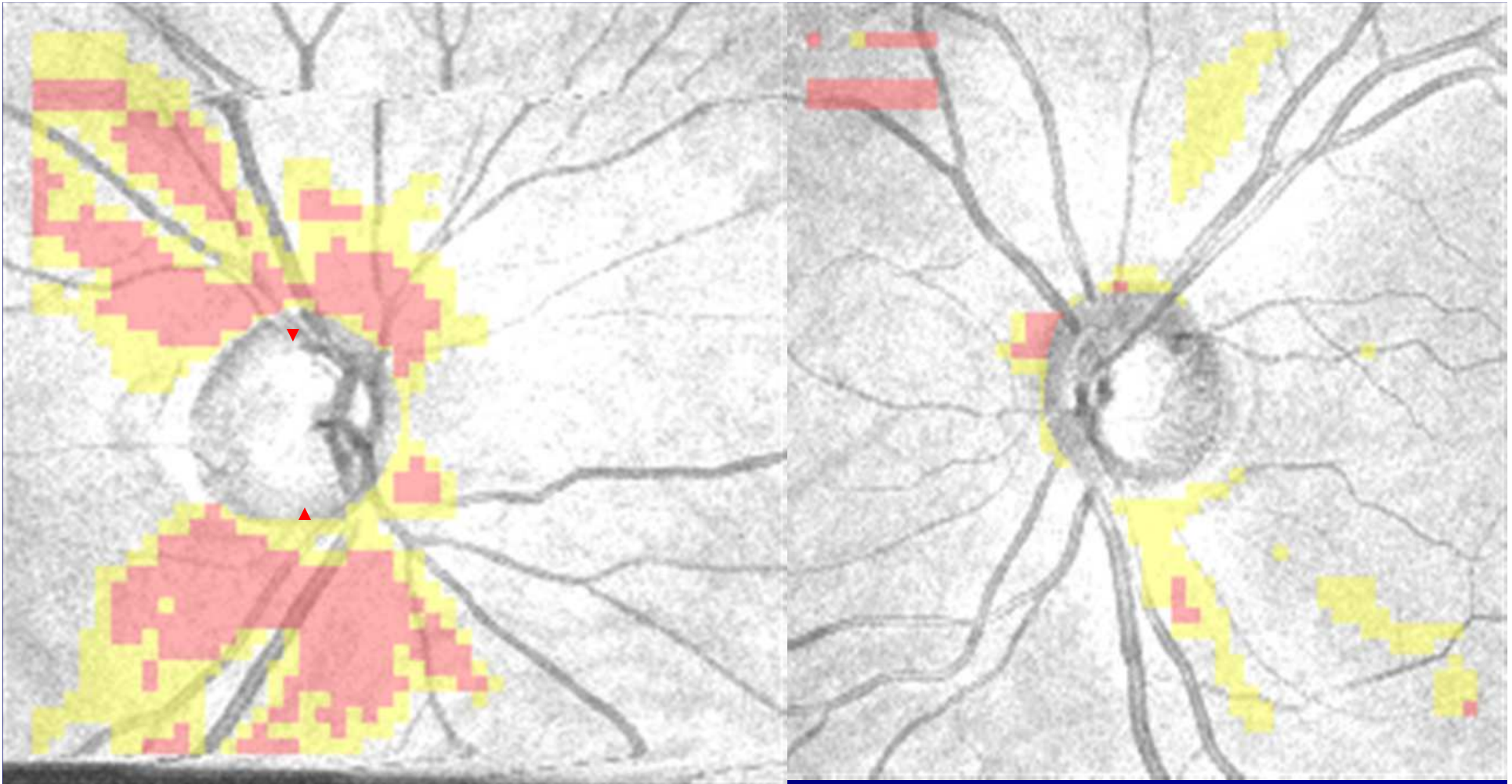
RM+ PLACCHE DI
DEMIELINIZZAZIONE

**II CASO
AMAUROSÌ ODX
LUGLIO 2011
BOLO CORTISONICO**



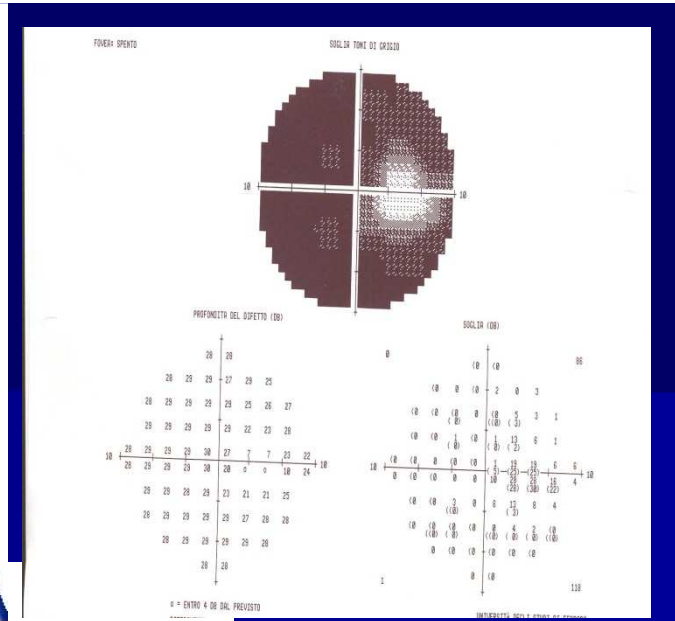
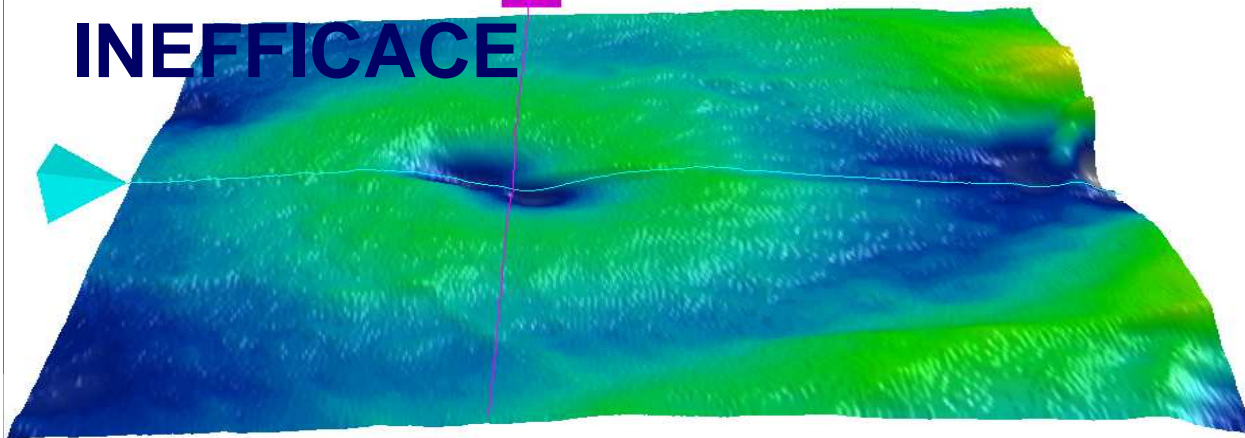
**RM+ PLACCHE DI
DEMIELINIZZAZIONE**



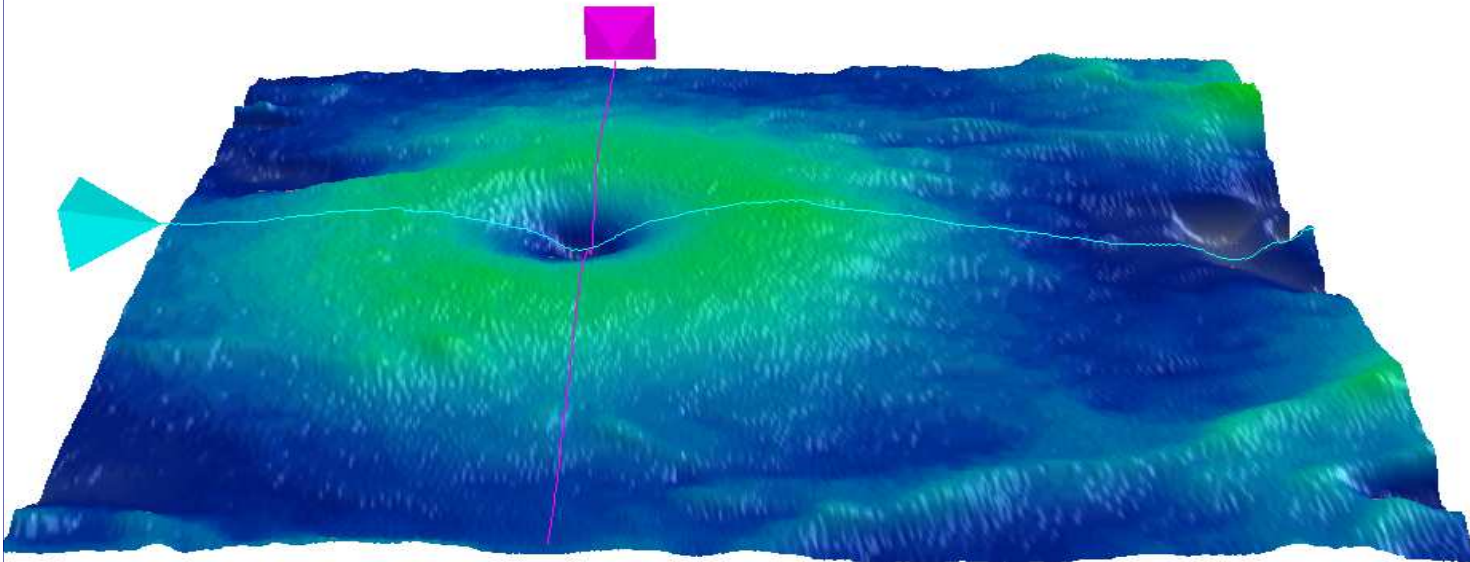


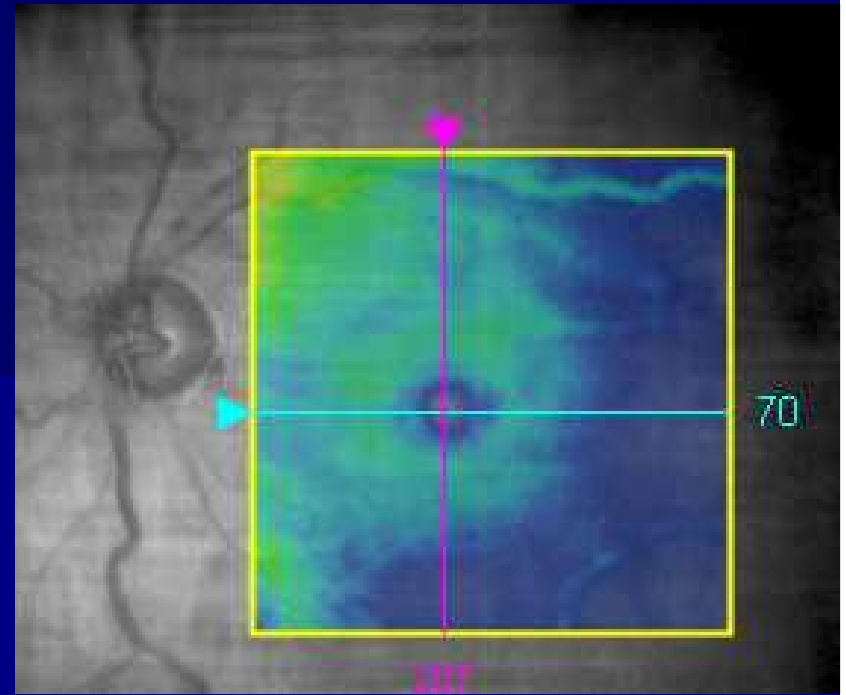
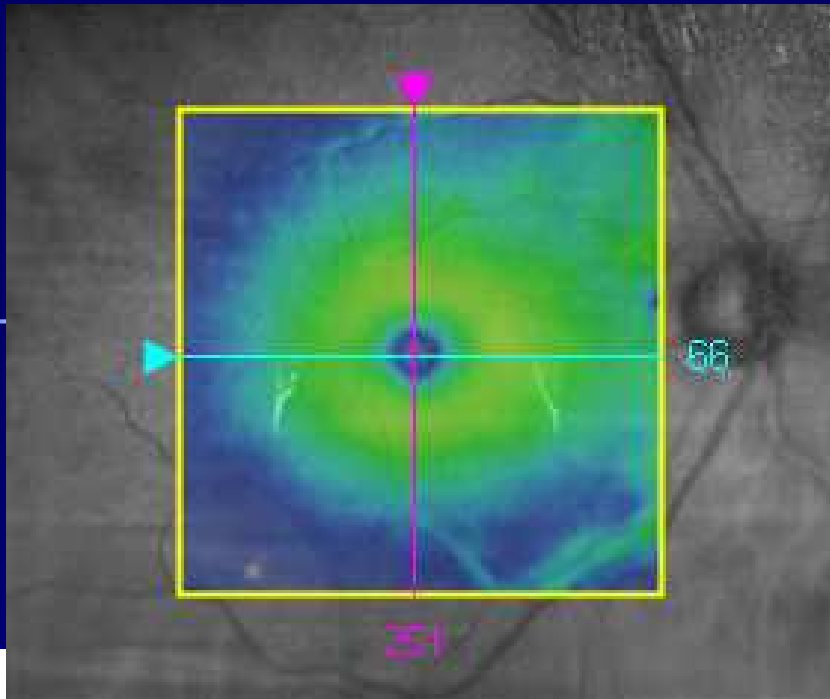
DOPO 1 MESE

BOLO CORTISONICO INEFFICACE

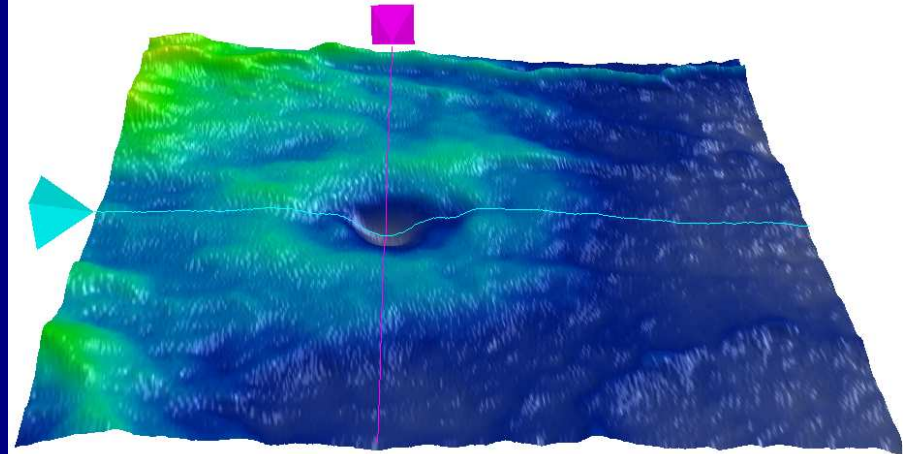
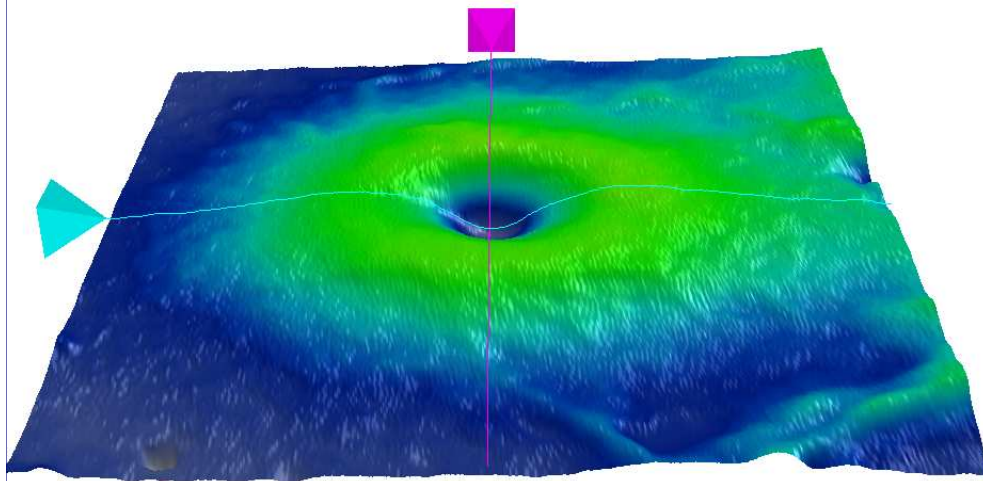


AGOSTO 2011





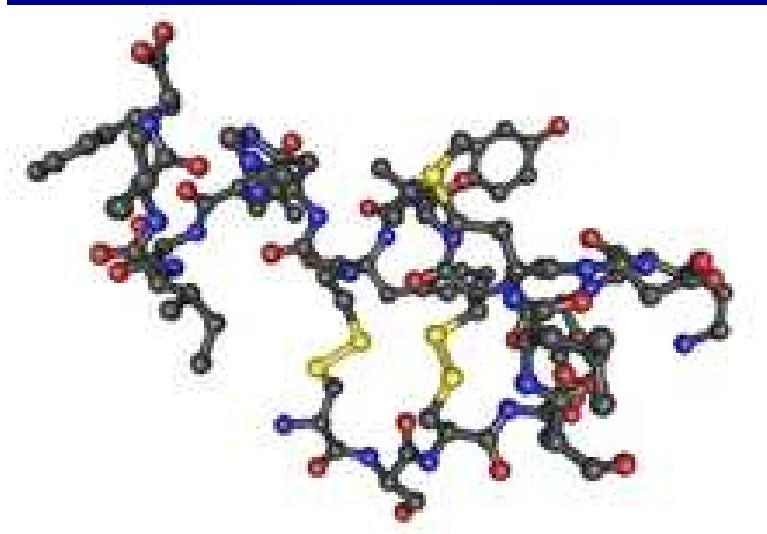
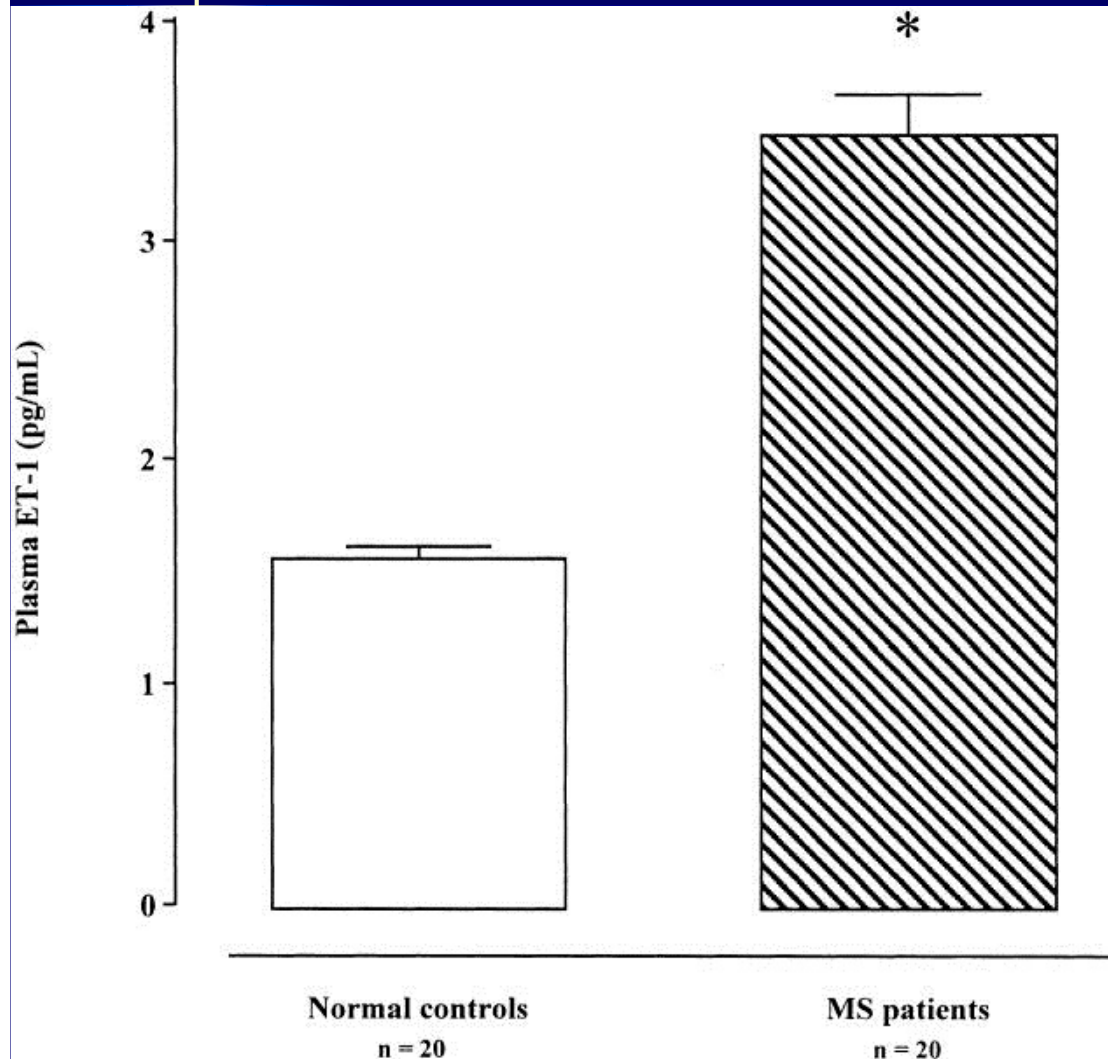
OACR



Increased Endothelin-1 Plasma Levels in Patients With Multiple Sclerosis

Haufschild, Timo MD; Shaw, Sidney G. PhD; Kesselring, Jürg MD; Flammer, Josef MD

Journal of Neuro-Ophthalmology:
March 2001 - Volume 21 - Issue 1 - pp 37-38



Vasospasm, its Role in the Pathogenesis of Diseases with Particular Reference to the Eye

Josef Flammer*, Mona Pache and Thérèse Resink

University Eye Clinic Basel, Mittlere Strasse 91, CH-4012 Basel, Switzerland

The exaggerated increase of endothelin-1 during an acute retrobulbar neuritis may account, in part, for the reversible visual loss in such patients.

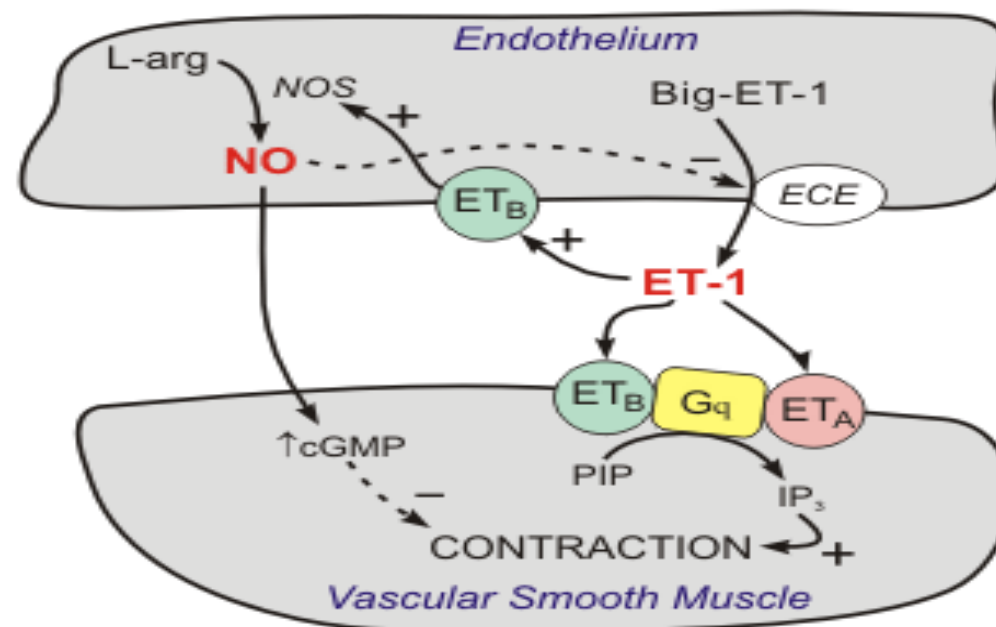


Figure 2. Endothelin receptors and interactions with nitric oxide.

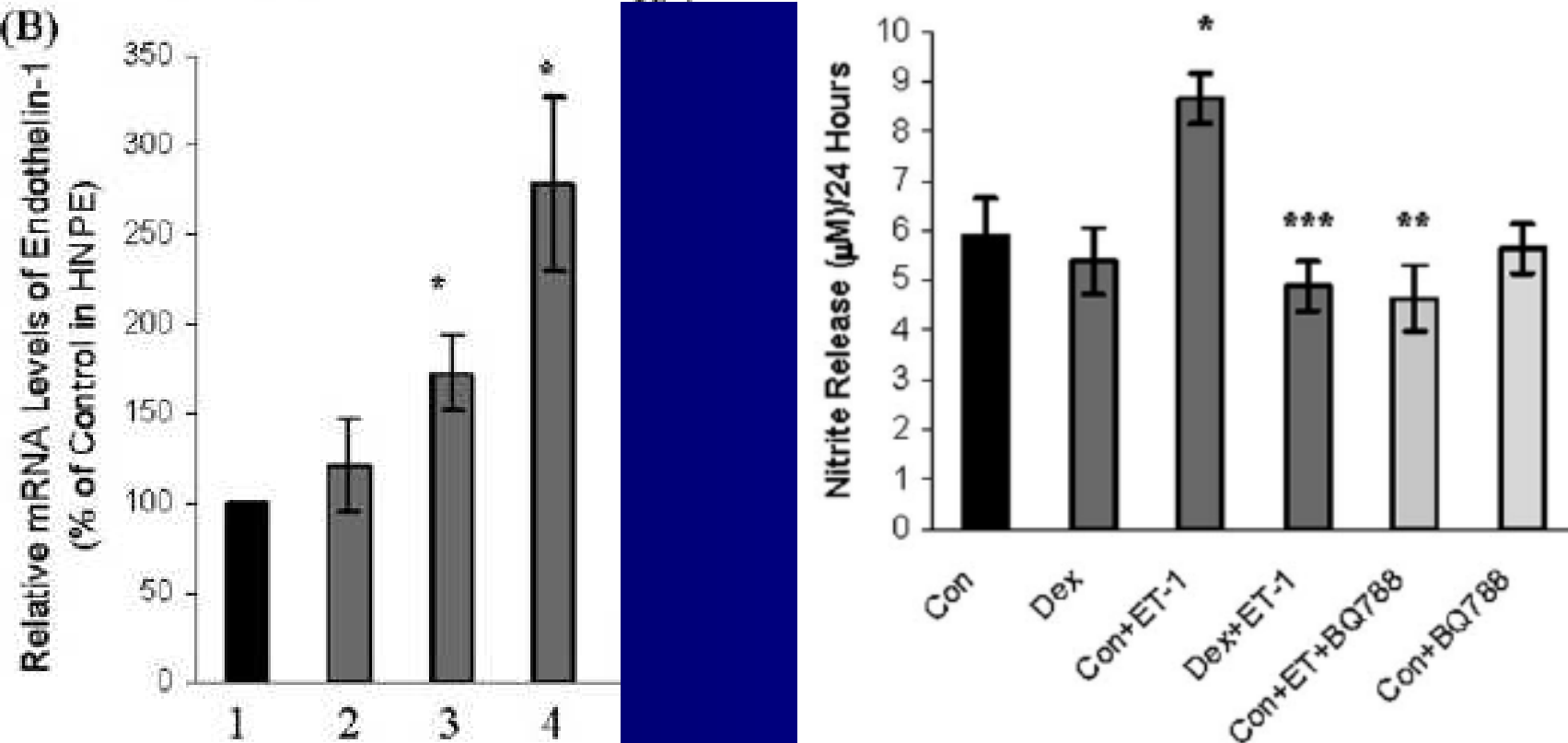
Glucocorticoids Regulate the Expression of the Human Osteoblastic Endothelin A Receptor Gene

abolic action on bone cell metabolism. Increased ET1 concentrations may also impair bone perfusion by vasoconstriction in a metabolically activated skeletal region.

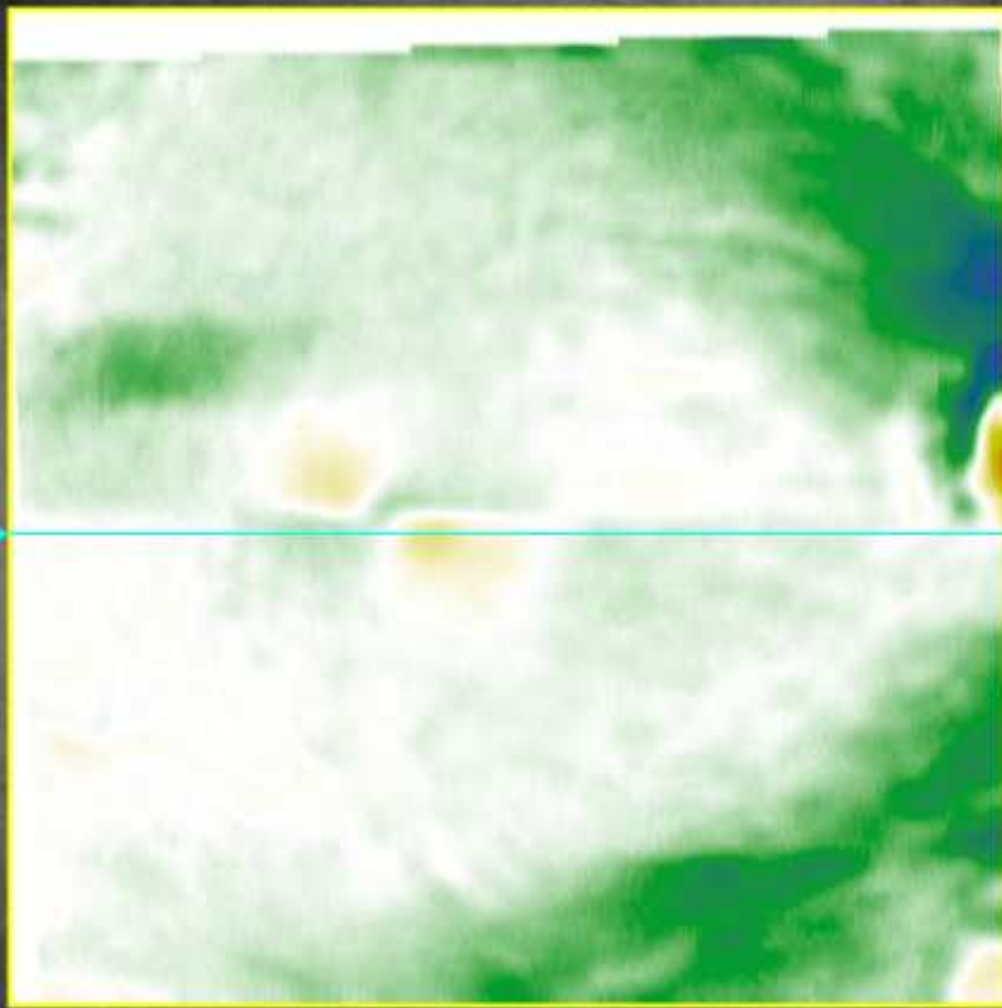
Interactions of Endothelin-1 with Dexamethasone in Primary Cultured Human Trabecular Meshwork Cells

Investigative Ophthalmology & Visual Science, December 2003, Vol. 44, No. 12

(B)



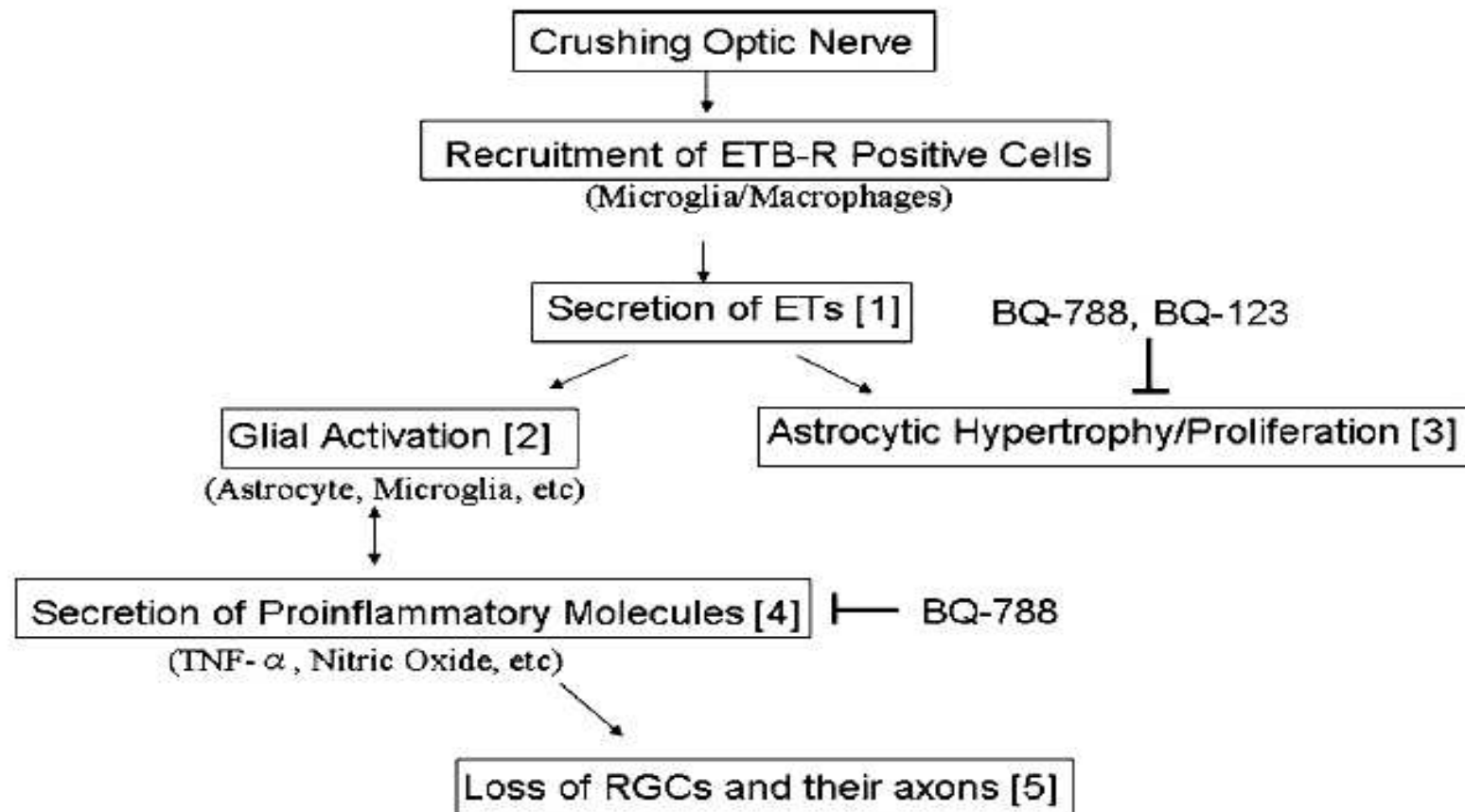
-70 MICRON



**BOLO CORTISONICO INEFFICACE
(CONTROINDICATO ?)**

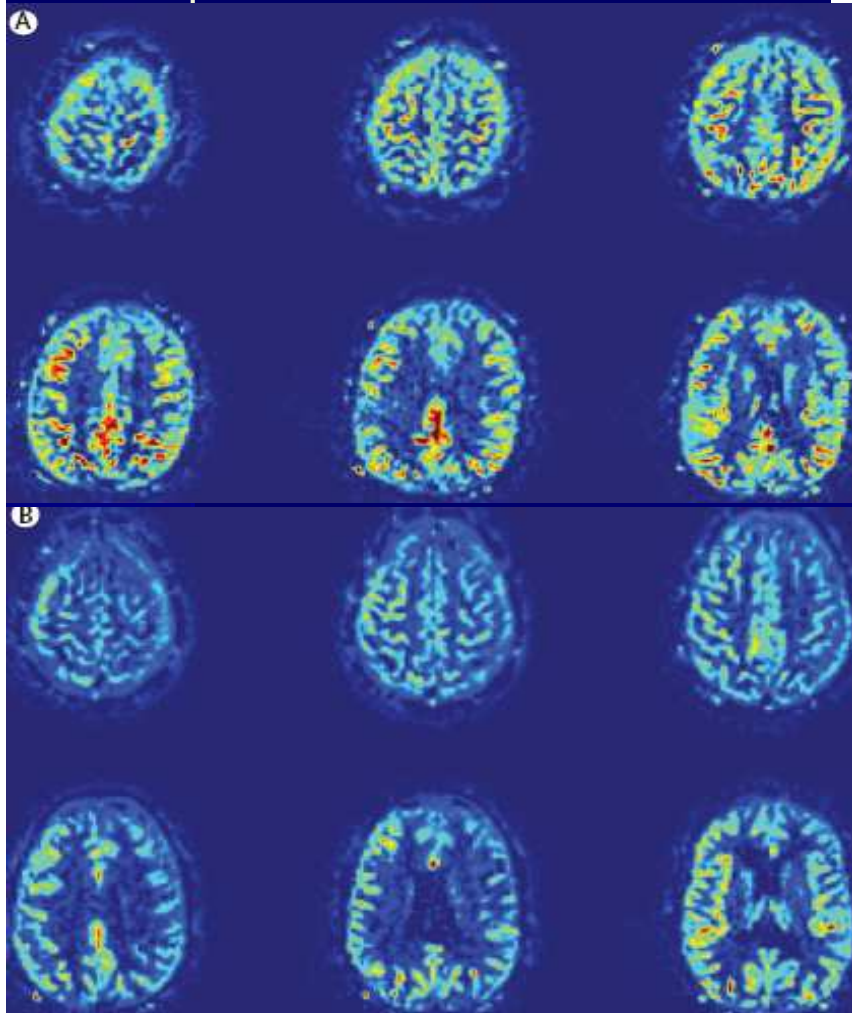
Blocking Endothelin-B Receptors Rescues Retinal Ganglion Cells from Optic Nerve Injury through Suppression of Neuroinflammation

Masabiro Tonari, Takuji Kurimoto, Taeko Horie, Tetsuya Sugiyama, Tsunehiko Ikeda, and Hidehiro Oku



Vascular aspects of multiple sclerosis

Miguel D'haeseleer, Melissa Cambron, Ludo Vanopdenbosch, Jacques De Keyser



Lancet Neurol 2011; 10: 657-66

Data suggest that a subtype of focal MS lesions might have an ischaemic origin

Infiammazione

Ischemia

2008

Fumatore 4-5 sigarette/die

Circa 2 anni fa traumatismo cranio-facciale dx. Da allora disturbi della vista parossistici a tipo fosfeni o “alone” intorno agli oggetti. Progressivo deficit del visus, in assenza di dolore.

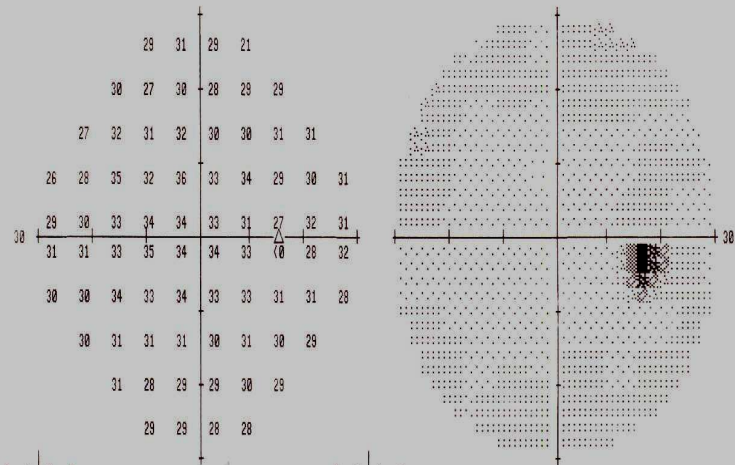
Eseguiva pertanto visita oculistica (con riscontro di deficit visivo 4/10 in occhio dx), PEV (a sin anomalie della conduzione della via ottica con aspetto sia assonali sia demielinizzanti), esami ematochimici per connettivopatie (nella norma), RMN encefalo (nella norma), non deficit all'obiettività neurologica.

Conclusioni: neurite ottica sin di n.d.d. a completamento diagnostico programmerei rachicentesi da effettuarsi c/o DH Neurologico di ferrara. Il paziente verrà contattato per un appuntamento.

Cordiali saluti

ORA: 08:13

FOVER: 37 DB



2	4	3	-6
1	-2	1	0
-2	2	0	0
-2	-2	3	0
0	-1	2	0
2	0	1	2
2	0	1	0
1	0	-1	-2
1	-2	-2	-1
1	0	-1	-2

0	2	2	-7
0	-3	0	-2
-3	1	-1	-1
-4	-3	2	-2
-1	-2	-1	-1
0	-2	-1	0
0	-2	0	-1
-1	-1	-3	-3
0	-3	-4	-3
-1	-2	-3	-3

GHT
ENTRO LIMITI NORMALI

MD -0.16 DB
PSD 1.72 DB

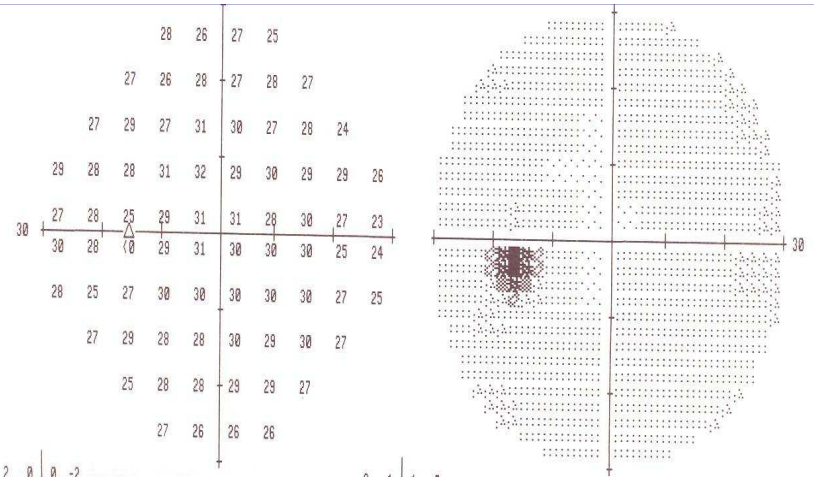
TOTAL
DEVIATIONE

MODELLO
DEVIATIONE

- ⋯ < 5%
- ⊗ < 2%
- ⊠ < 1%
- < 0.5%

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 AZIENDA OSPEDALIERA UNIVERSITARIA S. ANNA
 CLINICA OCULISTICA
 DIRETTORE PROF. ADOLFO SEBASTIANI
 ORTODONTISTA-SS. DI OTTALMOCIA

FOVER: 35 DB



2	0	0	-2
-1	-3	-1	-2
-2	-1	-3	0
-1	-2	-4	-1
-4	-3	-4	-2
-1	-3	-4	-3
-3	-6	-5	-2
-4	-2	-3	-5
-6	-3	-3	-2
-3	-3	-2	-3

3	1	1	0
1	-1	0	-1
-1	0	-2	1
0	-1	-2	0
-2	-1	-2	-1
0	-1	-3	-1
-1	-5	-4	-1
-3	-1	-2	-3
-4	-1	-2	0
-1	-2	-1	-1

GHT
ENTRO LIMITI NORMALI

MD -2.95 DB P < 2%
PSD 1.56 DB

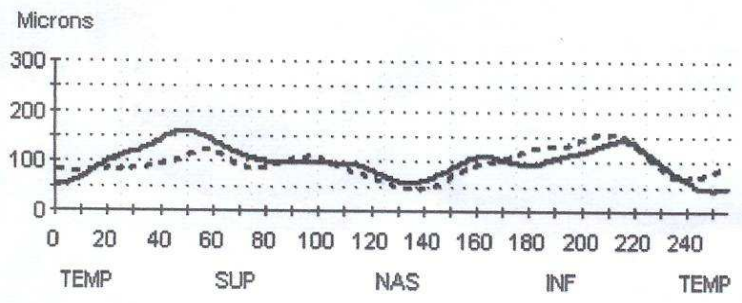
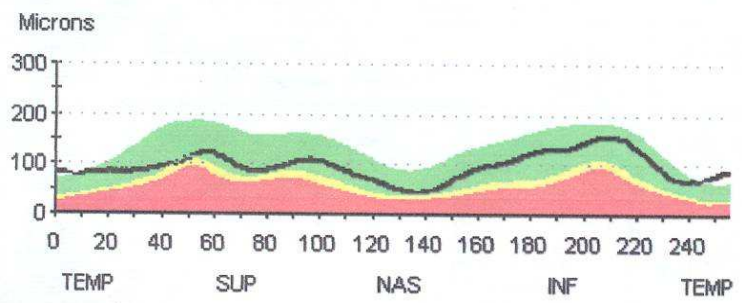
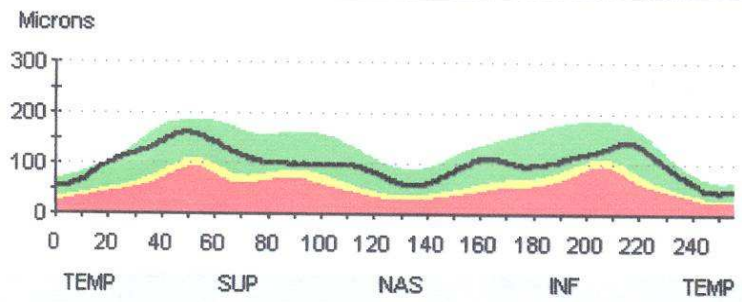
TOTAL
DEVIATIONE

MODELLO
DEVIATIONE

- ⋯ < 5%
- ⊗ < 2%
- ⊠ < 1%
- < 0.5%

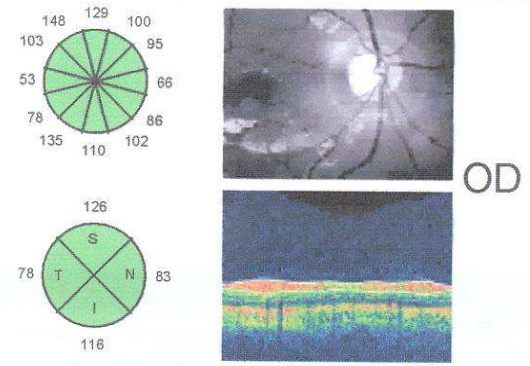
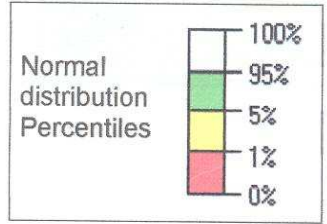
UNIVERSITÀ DEGLI STUDI DI FERRARA
 AZIENDA OSPEDALIERA UNIVERSITARIA S. ANNA
 CLINICA OCULISTICA
 DIRETTORE PROF. ADOLFO SEBASTIANI
 ORTODONTISTA-SS. DI OTTALMOCIA

OCT DEL
NERVO OTTICO
E
CALO VISUS DI
NDD

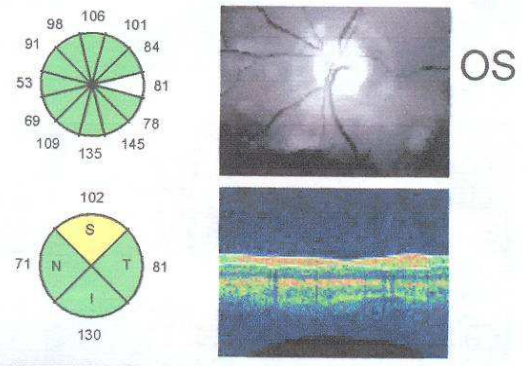


— OD - - - - OS

OD	Scans used	1, 2, 3
OS	Scans used	1, 2, 3

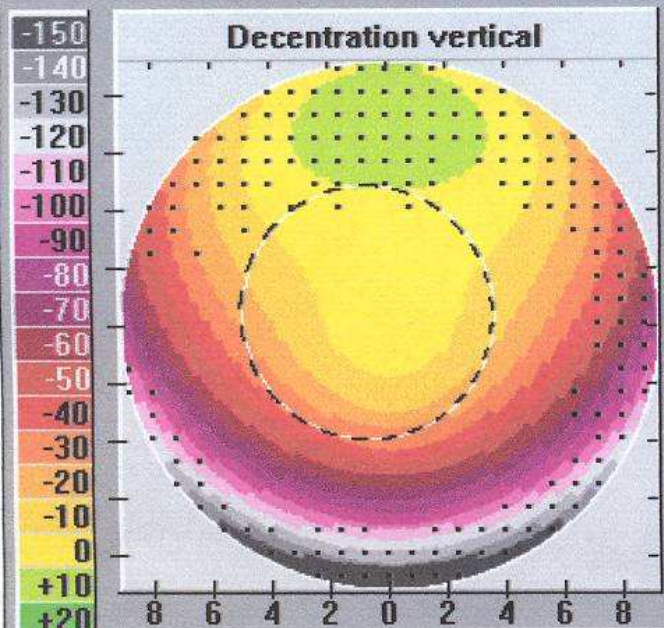


Signal Strength (Max 10)	9
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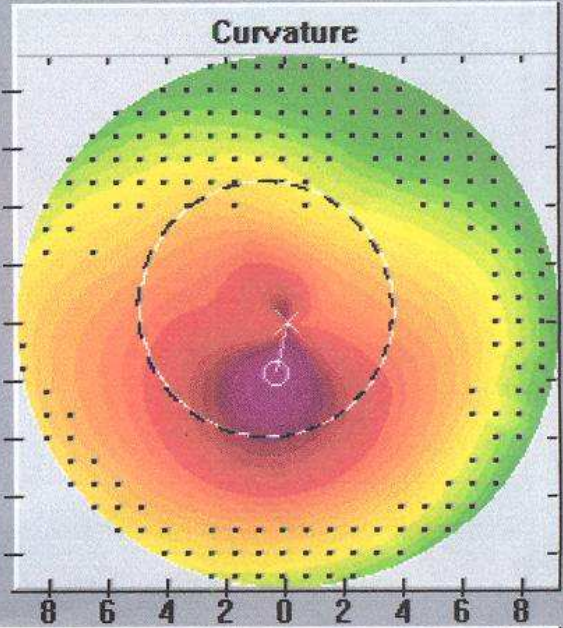
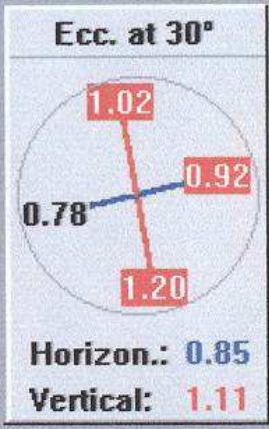
Signal Strength (Max 10)	8
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	OD (N=3)	OS (N=3)	OD-OS
lmax/smax	0.89	1.28	-0.39
smax/lmax	1.13	0.78	0.35
smax/tavg	2.05	1.50	0.55
lmax/tavg	1.82	1.92	-0.10
smax/navg	1.94	1.71	0.22
Max-Min	116.00	114.00	2.00
smax	160.00	122.00	38.00
lmax	142.00	156.00	-14.00
savg	126.00	102.00	24.00
lavg	116.00	130.00	-14.00
Avg.Thick	100.52	95.98	4.55



Indices

ISV:	66
IVA:	0.61
KI:	1.18
CKI:	1.05
Rmin:	6.65
IHA:	61.8
IHD:	0.045
ABR:	1.6
AA:	59%



Keratoconus Level Topogr.

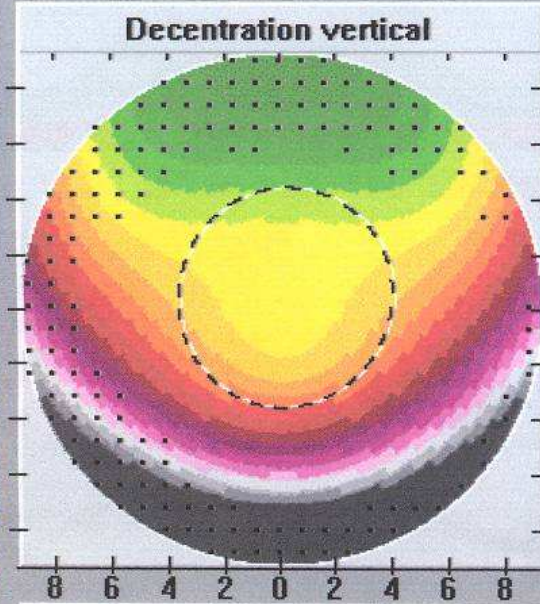
KC 2

- 150
 - 140
 - 130
 - 120
 - 110
 - 100
 - 90
 - 80
 - 70
 - 60
 - 50
 - 40
 - 30
 - 20
 - 10
 - 0
 - +10
 - +20
 - +30
 - +40
 - +50
 - +60
 - +70
 - +80
 - +90
 - +100
 - +110
 - +120
 - +130
 - +140
- µm

- 88.8
- 80.4
- 73.4
- 67.5
- 62.5
- 58.2
- 54.4
- 51.1
- 49.6
- 48.2
- 46.9
- 45.6
- 44.4
- 43.3
- 42.2
- 41.2
- 40.2
- 39.2
- 38.4
- 37.5
- 36.7
- 35.9
- 35.2
- 34.4
- 33.8
- 33.1
- 32.1
- 20.5
- 15.0
- 11.8
- D
- Sag
- Abs

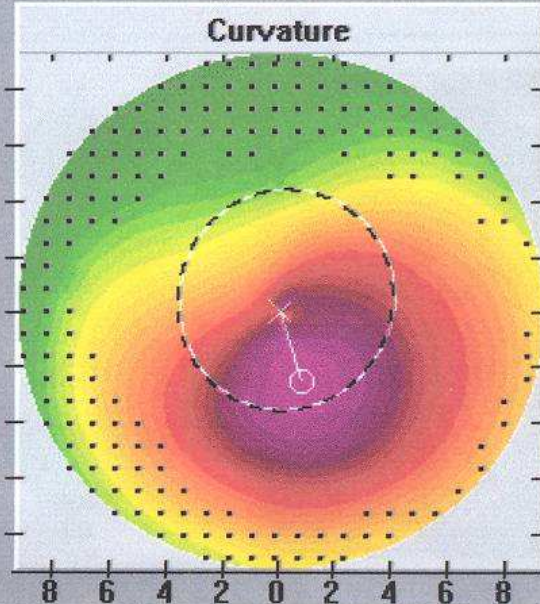
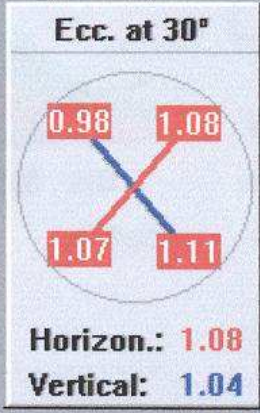
Sinistro	Progress of Indices	Destro (Current)
	15/04/08	Date: 15/04/08
	90	ISV: 66
	1.00	IVA: 0.61
	1.27	KI: 1.18
	1.06	CKI: 1.05
	6.37	RMin: 6.65
	109.2	IHA: 61.8
	0.079	IHD: 0.045
	2.8	ABR: 1.6
	2-3	KKS: 2
	63%	AA: 59%

-150
-140
-130
-120
-110
-100
-90
-80
-70
-60
-50
-40
-30
-20
-10
0
+10
+20
+30
+40
+50
+60
+70
+80
+90
+100
+110
+120
+130
+140
μm



Indices

ISV:	90
IVA:	1.00
KI:	1.27
CKI:	1.06
Rmin:	6.37
IHA:	109.2
IHD:	0.079
ABR:	2.8
AA:	63%



88.8
80.4
73.4
67.5
62.5
58.2
54.4
51.1
49.6
48.2
46.9
45.6
44.4
43.3
42.2
41.2
40.2
39.2
38.4
37.5
36.7
35.9
35.2
34.4
33.8
33.1
32.1
20.5
15.0
11.8
D
Sag
Abs

Keratoconus Level Topogr.

KC 2-3

Sinistro (Current)	Progress of Indices	Destro
	15/04/08	Date: 15/04/08
	90	ISV: 66
	1.00	IVA: 0.61
	1.27	KI: 1.18
	1.06	CKI: 1.05
	6.37	RMin: 6.65
	109.2	IHA: 61.8
	0.079	IHD: 0.045
	2.8	ABR: 1.6
	2-3	KKS: 2
	63%	AA: 59%

EXTENDED REPORT

Interpreting the multifocal visual evoked potential: the effects of refractive errors, cataracts, and fixation errors

B J Winn, E Shin, J G Odel, V C Greenstein, D C Hood

Br J Ophthalmol 2005;89:340–344. doi: 10.1136/bjo.2004.047910

Both refractive errors and cataracts blur the stimulus. Whether dioptric or diffusive, blur can be thought of as a spatial frequency filter selectively eliminating high spatial frequencies. The greater the blur, the greater the loss of high spatial frequency resolution. The amplitude of the response in

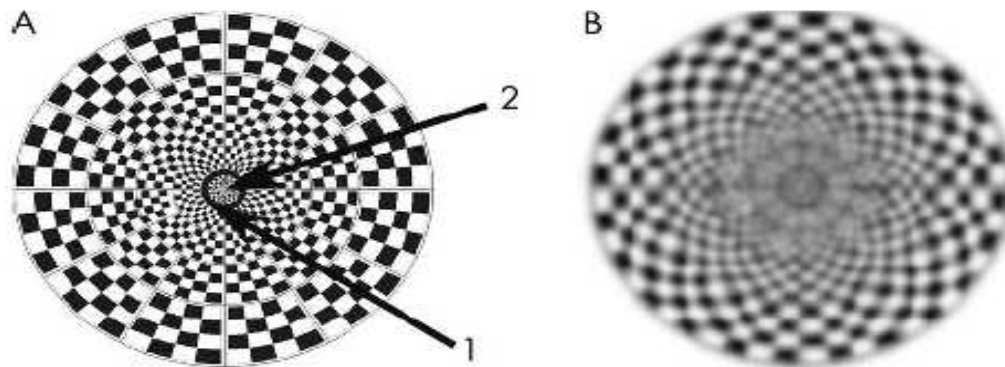


Figure 3 (A) The mfVEP display. Arrow 1 marks the position of fixation of the right eye for the steady, eccentric fixation condition. Arrow 2 marks the position of fixation of the left eye for all conditions. Note the relative difference in size of the stimulus sectors at fixation for the two eyes. (B) A blurred mfVEP display created by a computerised high spatial frequency filter. Note that pattern resolution is virtually lost in the central sectors but relatively preserved in the peripheral sectors.