UC Davis

Ophthalmology and Vision Science

Title

Ocular manifestations and visual outcomes of patients with giant cell arteritis

Permalink

https://escholarship.org/uc/item/8tv5b6m1

Authors

Lau, Janelle Celiker, Pelin Moghadam, Mohammad et al.

Publication Date

2024-04-01

Supplemental Material

https://escholarship.org/uc/item/8tv5b6m1#supplemental

Data Availability

The data associated with this publication are not available for this reason: NA



Visual Outcomes of Giant Cell Arteritis

Janelle Lau, BS¹; Pelin Celiker, MD²; Mohammad Johari Moghadam, MD²; Fateme Montazeri, MD, MPH²; Yin Allison Liu, MD, PhD¹; ¹University of California, Davis School of Medicine; ²Department of Ophthalmology, UC Davis Health, Sacramento, CA

Introduction

Giant cell arteritis (GCA) is a large vessel vasculitis classically affecting branches of the common carotid artery and is the leading cause of blindness in neuro-ophthalmology if left untreated.

- Temporal artery biopsy (TAB) is the gold standard for diagnosing GCA but has variable sensitivity due to urgent need for treatment with high-dose corticosteroids.¹
- Previous studies on ocular manifestations have only included small cohorts.²
- Additionally, longitudinal evaluation of the visual outcomes have not yet been assessed.

We hypothesized that diagnosis of GCA, ocular manifestations, and treatment response can be better evaluated using neuro-ophthalmic evaluation and imaging techniques.

• We also hypothesized that changes in the retina & optic nerve correlate with disease status and can predict visual outcomes.

Materials & Methods

We retroactively reviewed neuro-ophthalmology visits of GCA patients performed at a tertiary center.

- Patients split into TAB+ or clinically diagnosed (CD) groups
- Eyes sorted into visual involvement (VI) or no VI (NVI)
- Measurements separated into intervals of 0-6 months, 6-12 months, and beyond 12 from date of TAB or CD

Assessments of ocular involvement:

- Best-corrected visual acuity ("BCVA" in logMAR and Snellen)
- Intraocular pressure ("IOP" in mmHg)
- Humphrey visual fields ("HVF" in median deviation)
- 24-2, Ishihara color plates ("% Color" in % correct)
- Erythrocyte sedimentation rate ("ESR" in mm/hr) and Creactive protein ("CRP" in mg/dL) were also recorded

Optical Coherence Tomography (OCT) measurements:

- Retinal nerve fiber layer ("RNFL" in µm)
- Ganglion cell complex ("GC-IPL" in μm)

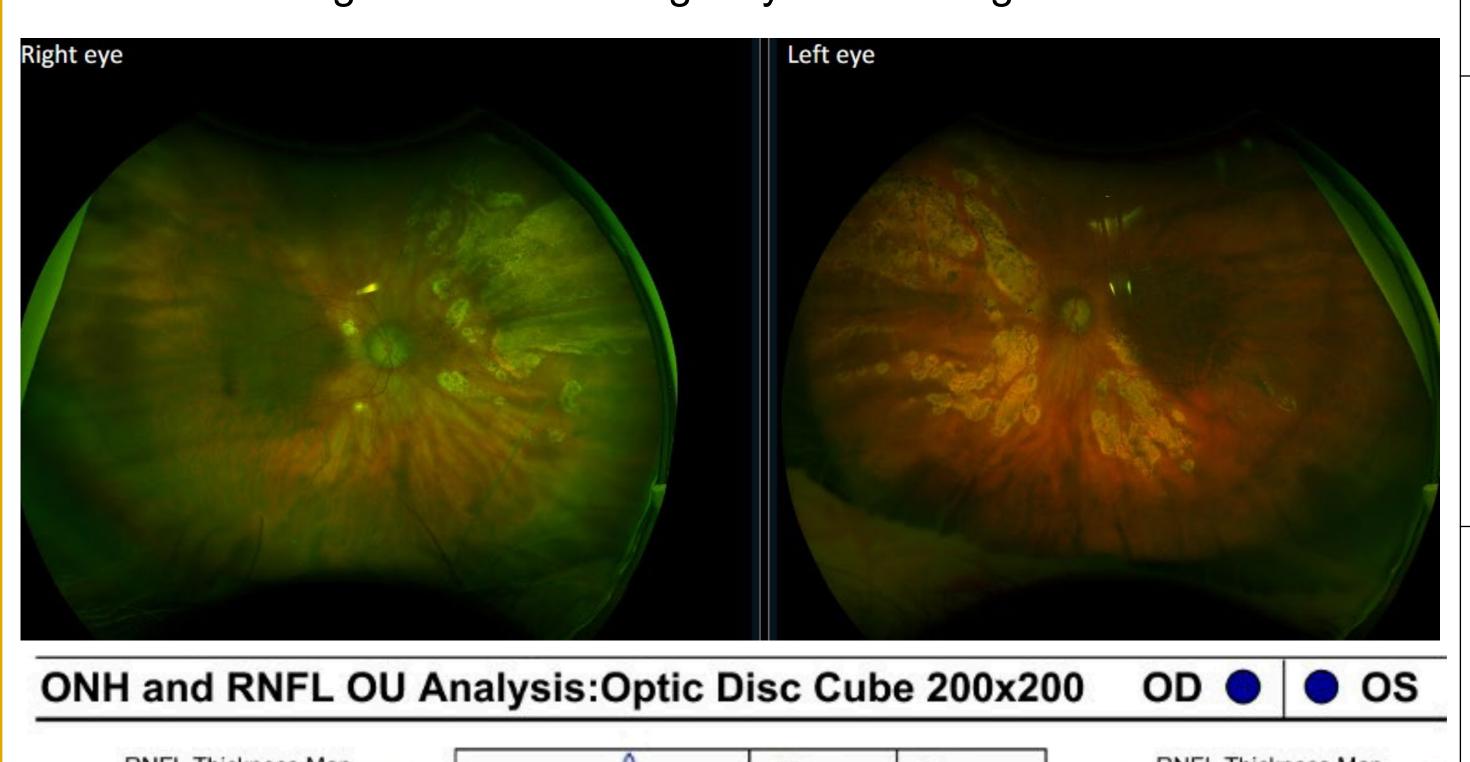
OCT-Angiography (OCT-A) measurements:

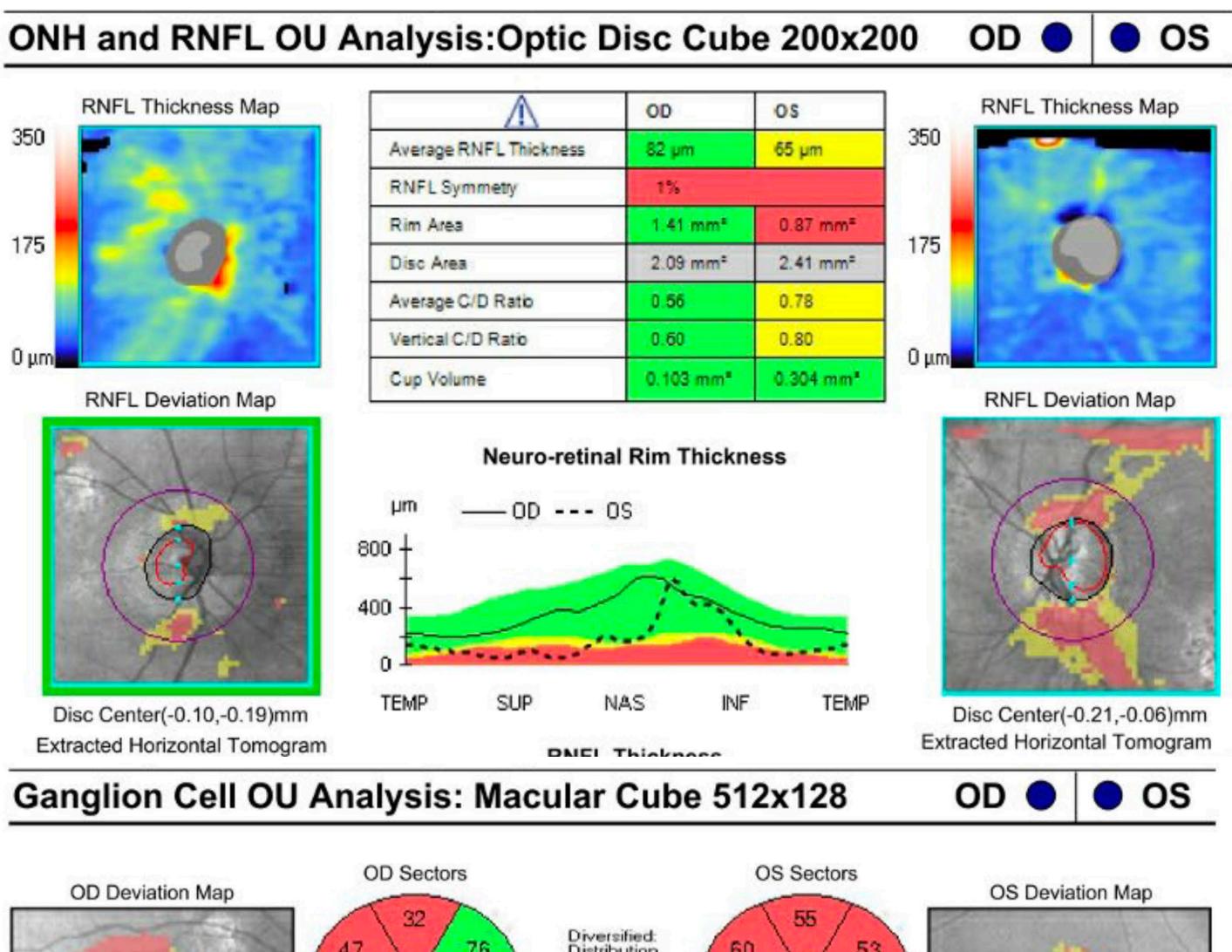
- Microvascular density of the inner retina ("ILM-IPL")
- Outer retina ("IPL-OPL")
- Radial peripapillary capillary ("RPC") vessels
- Area of foveal avascular zone ("FAZ" in mm²)

Results

A total of 219 visits from 25 patients were included (146 TAB+ eyes, 73 CD eyes).

 All patients received high-dose intravenous steroids at the time of diagnosis in an emergency care setting.





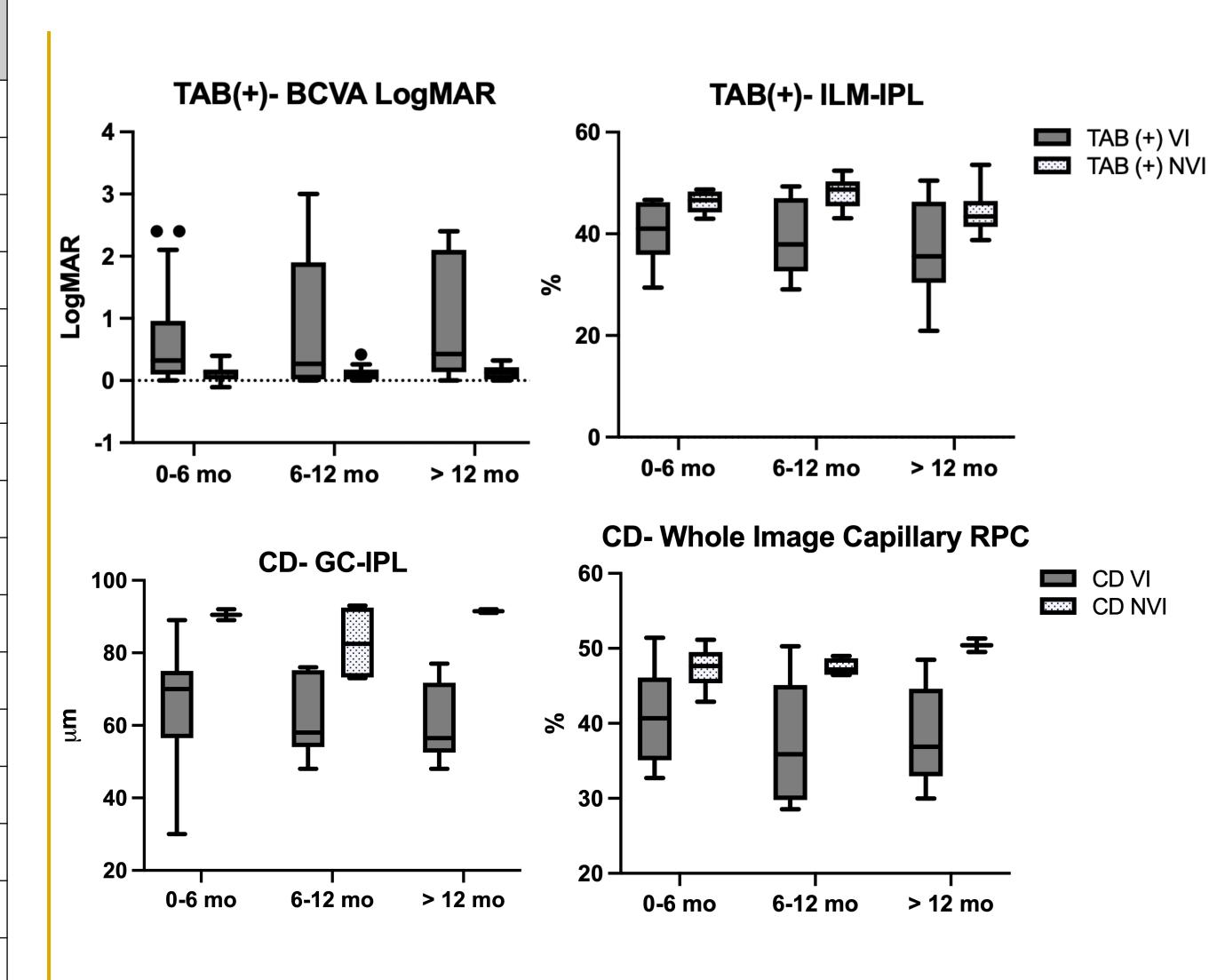
Average GCL+IPL Thickness

Minimum GCL + IPL Thickness

• os	
ickness Map	> 12 Montl
eviation Map	Clinica
	0-6 Mo
-0.21,-0.06)mm ontal Tomogram	6.44
ation Map	6-12 Mont
	> 12 Mont

TAB (+)	Measurement	Visual Impairment	No Visual Impairment	P-value
0-6 Months	BCVA	20/90	20/26	<0.001
	% Color	56.72%	85.30%	0.021
	HVF	-16.73	-1.90	<0.001
	GC-IPL	59.27	78.35	<0.001
	ILM-IPL	40.19	46.28	0.044
6-12 Months	BCVA	20/139	20/26	0.006
	% Color	25.00%	88.39%	0.001
	HVF	-17.23	-1.58	<0.001
	RNFL	71.31	93.17	<0.001
	GC-IPL	54.12	71.70	<0.001
	RPC	37.33	48.52	0.003
	ILM-IPL	39.10	47.85	0.005
> 12 Months	BCVA	20/166	20/26	<0.001
	% Color	37.25%	91.67%	<0.001
	HVF	-11.13	-2.43	0.01
	RNFL	68.07	87.75	<0.001
	GC-IPL	57.07	75.82	<0.001
	RPC	32.86	45.52	<0.001
	ILM-IPL	37.34	44.44	0.01
Clinically	Measurement	VI	NVI	P-Value

Clinically Diagnosed	Measurement	VI	NVI	P-Value
0-6 Months	IOP	16.26	11.25	0.008
	CRP	2.60	1.12	0.040
	RNFL	76.32	103	<0.001
	GC-IPL	65.84	90.50	<0.001
	RPC	40.81	47.41	0.024
6-12 Months	IOP	15.64	10	0.008
	CRP	0.15	1.66	<0.001
	RNFL	75.41	105.66	<0.001
	GC-IPL	62.07	85.66	0.003
	RPC	37.40	47.75	0.008
> 12 Months	RNFL	72	106.50	0.003
	GC-IPL	60	91.50	<0.001
	RPC	38.17	50.43	0.014
	ILM-IPL	37.38	49.40	0.008
	IPL-OPL	37.20	54.43	0.008



Conclusions

- Thinner RNFL and GC-IPL in eyes with visual impairment: ischemic injury causing retinal thinning.
- Decreased retinal density in the capRPC and ILM-IPL layers at certain time points indicating GCArelated hypoperfusion in those respective layers.

Visual relapse of GCA is rare after steroid initiation.

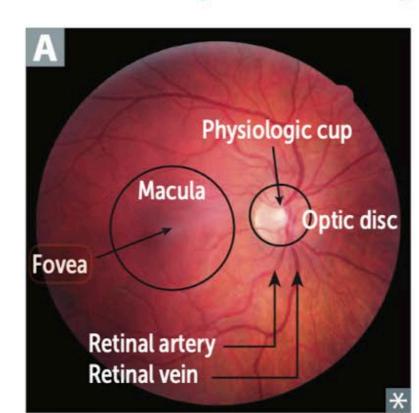
 Efficacy of steroids in managing symptoms several months post-diagnosis and the efficacy of tocilizumab in long-term prevention of GCA relapse.

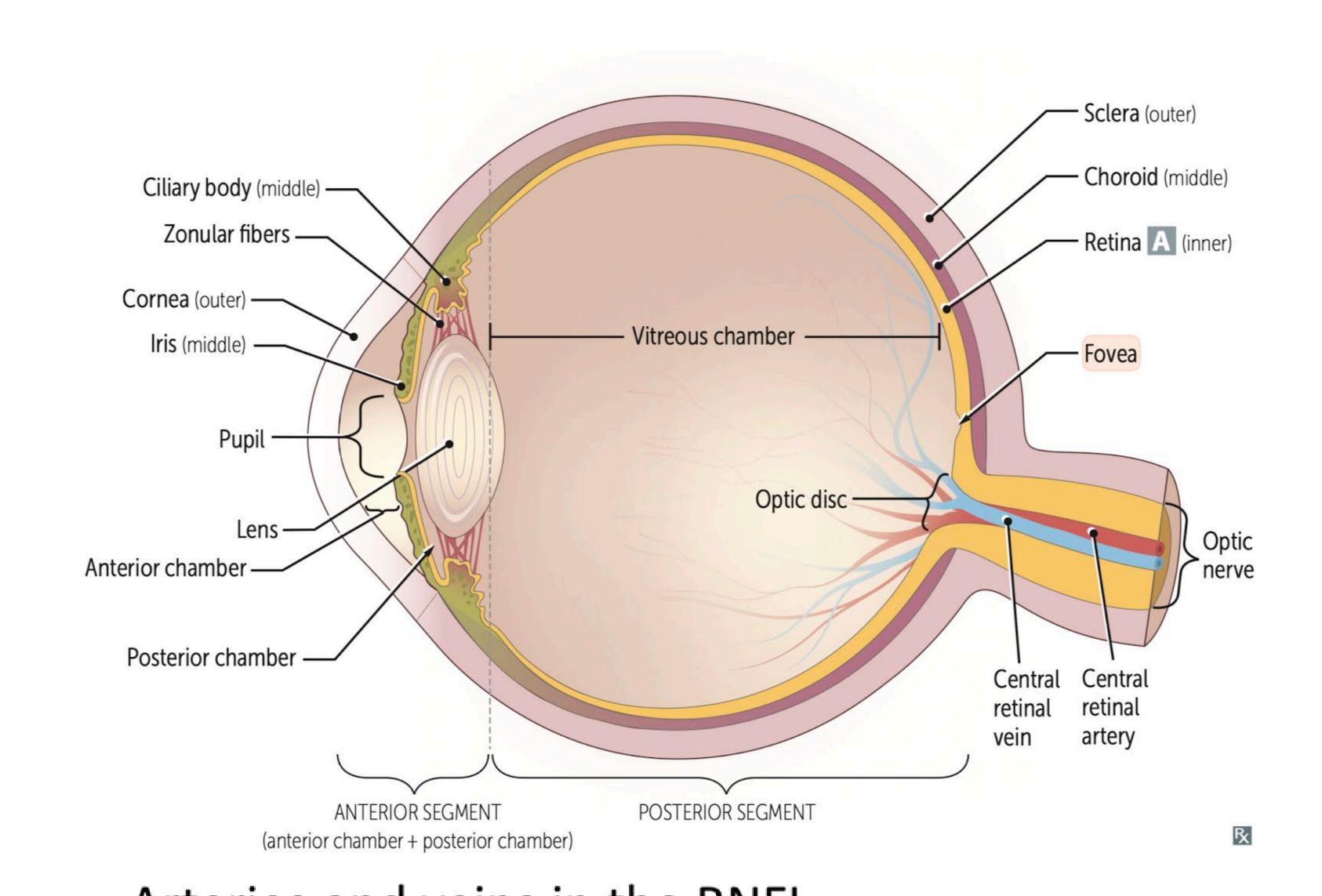
References

¹ Davies CG, May DJ. The role of temporal artery biopsies in giant cell arteritis. Ann R Coll Surg Engl. 2011 Jan;93(1):4-5. doi: 10.1308/003588411X12851639107476. PMID: 21418754; PMCID: PMC3293260.

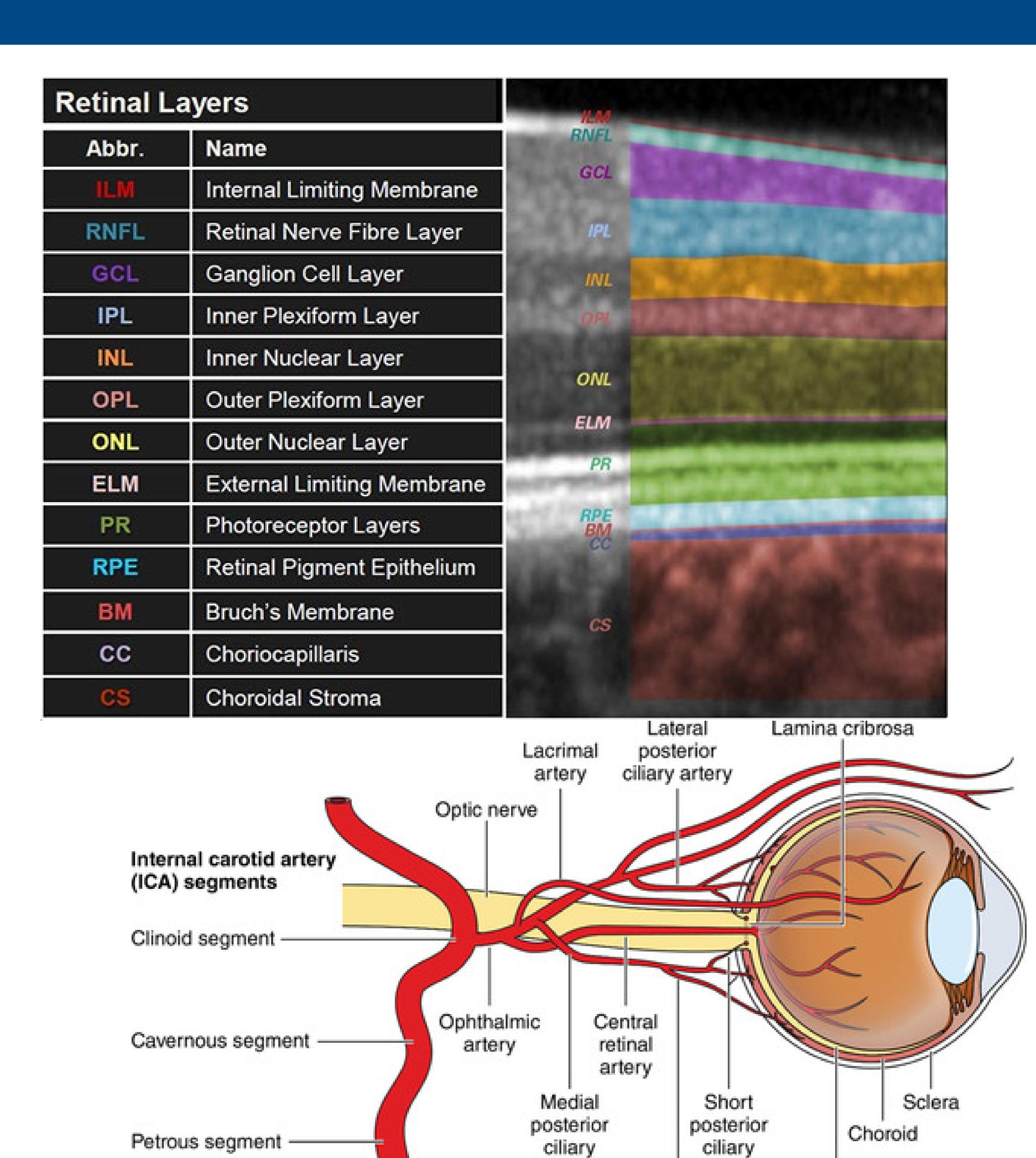
² Gaier ED, Gilbert AL, Cestari DM, Miller JB. Optical coherence tomographic angiography identifies peripapillary microvascular dilation and focal non-perfusion in giant cell arteritis. Br J Ophthalmol. 2018 Aug;102(8):1141-1146.

Normal eye anatomy





RNFL — Capillary networks (perfusion) IPL — Choroidal Vessels (diffusion)



arteries

Long posterior

ciliary artery

Retina

artery