

Type of the Paper (Research Article)

Impact of Acute Unilateral Anterior Ischemic Optic Neuropathy on

Central Macular Thickness and Foveal Avascular Zone: An Optical

Coherence Tomography and Angiography Study

Esraa O. M. Eissa^{1*}, Ahmed T. S. Saif², Omar M. S. Said², Mahrous H. Shaheen²

¹Fayoum Ophthalmology Hospital, Fayoum, 63611 Egypt.

²Ophthalmology Department, Faculty of Medicine, Fayoum University, Fayoum, 63514 Egypt.

*Correspondence: Esraa O. M. Eissa, <u>ie1119@fayoum.edu.eg</u>, Tel: (002) 01018055526.

Received:	27 August, 2024	Reviewed:	11 September, 2024
Accepted:	18 February, 2025	Published online:	20 March, 2025

Abstract:

Introduction: Non-Arteritic Anterior Ischemic Optic Neuropathy (NAION) is a known ischemic change that causes changes in the optic disc. Fundus fluorescein angiography and field perimetry are useful in the diagnosis of NAION. Optical Coherence Tomography (OCT) and Optical Coherence Tomography Angiography (OCTA), are advanced non-invasive modalities that can be used to evaluate the structural and vascular changes at different retinal layers.

Aim of the study: To assess the impact of acute NAION on the central macular thickness and foveal avascular zone via OCT and OCTA during the acute phase (1-3 weeks from onset) before the resolution of disc edema.

Subjects and Methods: 30 patients with unilateral acute NAION presenting at the outpatient department at Fayoum University Hospitals were recruited. All cases (diseased and normal eyes) were examined using OCT and OCTA.

Results: showed a significantly lower CMT in diseased eyes after the resolution of edema. Reduction in FAZ was statistically non-significant.

Conclusions: We can take into consideration the quantitative differences in CMT during follow-up of the progression in NAION cases.

Keywords: Optical Coherence Tomography Angiography; Non-Arteritic Anterior Ischemic Optic Neuropathy; Optical Coherence Tomography; Edema.

1. Introduction

Anterior type of ischemic optic neuropathy (AION) is a known optic neuropathy characterized by severe, acute, painless visual loss. It is believed to be due to ischemic changes in the anterior segment of the optic disc. AION is subdivided into two types: an arteritic (AAION) and a nonarteritic (NAION) [1]. NAION is considered the most common optic neuropathy (other than glaucoma) above the age of 50 years. It represents up to approximately 85% of all AION cases. Pathogenesis is believed to be a result of hypoperfusion in short posterior ciliary arteries (PCAs) that can be due to systemic or local hypoperfusion [2].

NAION usually related is to hypertension, arteriosclerosis, diabetes hyperlipidemia, mellitus, hyperhomocysteinemia, nocturnal hypotension, apnea and also the sleep structural predisposition in a crowded disc and a small or absent cup. Vessel occlusion is distal to PCAs, affecting para-optic tributaries within the optic disc [1]. Patients are commonly discovered on awakening. On examination, visual acuity is moderately to severely impaired in around 70% of patients. In 30% of cases, visual acuity can be non-affected or only slightly diminished. The majority of cases show inferior altitudinal visual field defects but other pictures like central, paracentral, quadrantic or arcuate defects can either be found [2].

OCTA is a trendy, non-invasive investigation which can demonstrate vessels of the retina and optic disc depending on blood flow. It easily can show microvasculature through the detection of blood flow motion contrast with no dye administered [3]. While FFA cannot provide enough demonstration of deep vasculature, OCTA is more efficient in showing macular and radial peri-papillary capillary network [4, 5], which originate from the central retinal artery (superficial plexus in addition to the deep ONH micro-vessels coming out from PCAs (deep plexus) [6].

2. Subjects and Methods

2.1 Subjects

This series included thirty persons visiting the Ophthalmology outpatient department in Fayoum University Hospitals, complaining of acute N-AION between April 2021 and April 2022.

Inclusion criteria

Patients who are ≥ 40 years old, complaining of unilateral NAION with acute/ abrupt painless diminution of vision and a non-affected other eye.

Exclusion criteria

Patients diagnosed with AAION or NAION in the non-acute stage or acute N-AION beside another neuropathy affecting the other side were excluded from this study.

3. Results

On comparing the demographic characteristics of the patients, the mean age among the study group was (53.1 ± 7.5) years old (ranging between 42 and 67 years). 60% were females. Regarding co-

Subjects that had cataracts or corneal opacities, high intraocular pressure (more than 22 mmHg), and errors >6 diopters were also excluded. Non-cooperative patients were not included.

2.2 Study design

A prospective, observational, comparative study. Normal eyes on the other side were considered the control sample.

2.3 Statistical Methods

This was done via SPSS (version 22; SPSS, Chicago, IL, USA). Data was reported as mean ±standard deviation values. Probability values <0.05 were assumed significant.

morbidities, 23.3% of cases were hypertensive, 16.7% were diabetic, and 13.3% of cases had both HTN and DM (**Table 1**).

Variable		Frequency (n=30)
Age (years) Mean ±SD		53.1 ±7.5
	Range	42-67
Sex	Sex Male	
	Female	18 (60%)
Co-morbidities	None	7 (23.3%)
	Hypertension (HTN)	7 (23.3%)
	Diabetes mellitus (DM)	5 (16.7%)
	Both HTN & DM	4 (13.3%)
	HTN, anemia, & thyroid	2 (6.7%)
	Behcet's disease / HCV	2 (6.7%)
	HTN – Cardiac disease	2 (6.7%)
	Thyroid disease	1 (3.3%)

Table 1: Analysis of demographic characteristics of cases.

On analyzing the visual acuity, there was a highly statistically significant difference between diseased and normal eyes with a lower mean VA in the diseased group during all visits. Inside each group, changes in visual acuity over the three follow-up visits were insignificant. On the other side, there was no statistically significant difference as regards intraocular pressure (IOP) (**Table 2**).

Table 2: Comparison of IOP and VA between groups and inside each group.

Variable		Diseased (n =30)	Diseased (n =30) Fellow (n =30)	
IOP		14.63 ±2.3 14.6 ±0.7		0.9
	1 st visit	0.178 ± 0.24	0.465 ± 0.29	< 0.001*
VA	2 nd visit	0.18 ± 0.24	0.465 ± 0.291	< 0.001*
	3 rd visit	0.173 ±0.229	0.465 ±0.291	< 0.001*

* Significant at p < 0.05.

CMT in OCTs of diseased eyes was significantly lower in the second and third visits. While in the first visit, CMT was insignificantly lower in diseased eyes. Inside the diseased group, we found a statistically significant drop in CMT during the second visit. During the third visit, no significant change was seen. The control group (normal eyes) had no significant change in all visits (**Table 3, Figure 1**).

Variable		Diseased (n =30) Fellow (n =		P -value
	1 st visit	261.43 ± 44.49	253.73 ±42.02	0.5
CMT	2 nd visit	237.37 ± 23.25	251.13 ±27.53	0.04*
	3 rd visit	235.47 ±23.45	250.6 ±27.18	0.02*
P -value		0.008*a	0.6 ^a	
		0.1 ^b	0.08^{b}	

Table 3 : Comparison of OCT	(CMT) changes in	different study	groups.
------------------------------------	------	--------------	-----------------	---------

* Significant at p < 0.05, a: change between 1st and 2nd visit, b: change between 2nd and 3rd visit



Figure 1: Comparison of CMT between study groups.

On Examining FAZ during all visits, a non-significant difference between case and control groups in all visits was observed. Changes in FAZ between consequent visits in both diseased and normal eyes were also non-significant (Figure 2, Tables 4, 5).



Figure 2: Comparison of mean FAZ in different study groups.

Table 4. Comparison of TAZ during follow-up in different study groups	Table 4:	: Com	parison	of FAZ	during	follow-u	p in	different	study	group
--	----------	-------	---------	--------	--------	----------	------	-----------	-------	-------

Variable		Diseased (n =30)	Fellow (n = 30)	P -value
	1 st visit	0.455 ± 0.465	0.355 ±0.15	0.3
FAZ	2 nd visit	0.451 ± 0.411	0.35 ±0.131	0.2
	3 rd visit	0.458 ± 0.406	0.326 ±0.122	0.09
	0.0 -			

* Significant at p <0.05.

Variable		Diseased (n =30)	Fellow (n =30)	
	1 st visit	0.455 ± 0.465	0.355 ± 0.15	
FAZ	2 nd visit	0.451 ± 0.411	0.35 ±0.131	
	3 rd visit	0.458 ± 0.406	0.326 ± 0.122	
P -value		0.7 ^a	0.8 ^a	
		0.6^{b}	0.1 ^b	

Table 5: Comparison of FAZ follow-up inside each study group.

* Significant at p <0.05, a: change between 1st and 2nd visit, b: change between 2nd and 3rd visit

4. Discussion

NAION is known to be the most common acute optic neuropathy in patients above fifty years old [1].

the On analyzing demographic characteristics of our patients, the mean and range of age were not much different from most studies in the literature, but the female predominance found in this series was uncommon in other series. A systematic review and meta-analysis published in 2022 by Khalili et al. reported that only five out of the 26 studies included in the review had female predominance [6]. Although most publications have demonstrated no sex predisposition, one large case-control series found males to be more vulnerable to developing NAION [7].

Regarding co-morbidities, 23.3% of our patients had hypertension, 16.7% had diabetes mellitus, and both hypertension and diabetes mellitus were found in 13.3% of patients.

We found a statistically considerable drop of visual acuity (VA) in diseased eyes in all patients during all visits. The drop in visual acuity over time in diseased eyes was also significant. In contrast, we found nonconsiderable differences in intraocular pressure (I.O.P). This was found consistent with most studies in the literature [6, 7].

Apart from the first visit, analysis of CMT showed a significantly lower mean in diseased eyes. There was also a significant decrease in most OCTs of diseased eyes during follow-up. Fernández-Buenaga et al. (2009) did a cross-section study on 24 patients with unilateral NAION (eleven females, thirteen males; mean age was $64 \pm$ 9 years) [7]. All subjects had a full examination. Normal and diseased eyes were scanned with Stratus OCT. Diseased eyes had considerably lower average CMT (195 ±26 µm) than non-affected eyes (209 ±23 µm) with a probability value of 0.023.

Our results found that differences in FAZ between case and control groups during all visits were non-significant with p-values >0.05. Changes inside each group between consequent visits were also non-significant.

Depending on our results, we can take into consideration the quantitative differences in CMT during follow-up of the progression in NAION cases. OCT and OCTA are fast and noninvasive investigations that can easily detect structural defects and quantitative changes in vascularity, respectively. It can give data about vessels of the retina, choroid and optic disc via one scan.

5. Conclusion

We have used OCT for quantitative analysis and comparison of central macular thickness in affected and fellow eyes. Changes were recorded from the onset till

Ethical committee approval: This research has been granted the prerequisite acceptance from the Research Ethical Committee in the Faculty of Medicine, Fayoum University, Egypt. Participants were educated about the objectives, examination, investigations, confidentiality and their privilege to decline participation. Every patient has given an informed consent.

References

- Hayreh SS. Ischemic optic neuropathies—where are we now? Graefes Arch Clin Exp Ophthalmol. 2013;251(8):1873-84. doi:10.1007/s00417-013-2399-z.
- Arnold AC. The 14th Hoyt lecture: ischemic optic neuropathy: the evolving profile, 1966– 2015. J Neuroophthalmol. 2016;36(2):208-215. doi:10.1097/WNO.00000000000395.
- Koustenis A Jr, Harris A, Gross J, Januleviciene
 I, Shah A, Siesky B. Optical coherence

12 weeks in the case and control groups. The results showed significant thinning of CMT in diseased eyes after the resolution of edema. FAZ analysis showed no significant differences between diseased and normal eyes during all visits. Changes inside each group were also non-significant. Changes found in fellow normal eyes were not significant all over the follow-up period.

Competing interests: There are no conflicts of interest for the authors.

Funding: No particular grants from public, commercial, or nonprofit funding organizations were given to this research.

AI declaration statement: Not applicable.

tomography angiography: an overview of the technology and an assessment of applications for clinical research. Br J Ophthalmol. 2017;101(1):16-20. doi:10.1136/bjophthalmol-2016-309389.

 Spaide RF, Klancnik JM Jr, Cooney MJ. Retinal vascular layers imaged by fluorescein angiography and optical coherence tomography angiography. JAMA Ophthalmol. 2015;133(1):45-50.

doi:10.1001/jamaophthalmol.2014.3616.

- Gao SS, Jia Y, Zhang M, Su JP, Liu G, Hwang TS, Bailey ST, Huang D. Optical coherence tomography angiography. Invest Ophthalmol Vis Sci. 2016;57(9):OCT27-36. doi:10.1167/iovs.15-19043.
- 6. Khalili MR, Abedini A, Fadakar K, Moghadas Sharif N, Yaseri M, Karkhaneh R. Optical

coherence tomography angiography in anterior ischemic optic neuropathy: a systematic review and meta-analysis. Eur J Ophthalmol. 2023;33(1):530-545.

doi:10.1177/11206721211070720.

 Fernández-Buenaga R, Rebolleda G, Muñoz-Negrete FJ, Contreras I, Casas-Llera P. Macular thickness. Ophthalmology. 2009;116(8):1587. doi:10.1016/j.ophtha.2009.04.017.