

Comparison between B-scan Ultrasonography and Optical Coherence Tomography in Evaluation of Macular Oedema

Thesis

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Tist of Abbreviations

Abb.	Full term
μm	micrometer
	Three Dimensional
	. Anterior to the Equator
	Age Related Macular Degeneration
	. Best Corrected Visual Acuity
	. Branch Retinal Vein Occlusion
CF	
	. Cystoid Macular Edema
	. Choroidal Neovascularization
	. Cone Outer Segment Tip
	. Central Retinal Artery
	. Central Retinal Thickness
CRV	. Central Retinal Vein
CRVO	. Central Retinal Vein Occlusion
dB	. Decibel
DM	. Diabetes mellitus
DME	. Diabetic Macular Edema
DMT	. Diffuse macular thickening
DR	. Diabetic Retinopathy
DRT	. Diffuse Retinal Thickness
E	. Equator
ELM	. External Limiting Membrane
ERM	. Epiretinal Membrane
ETDRS	. Early Treatment Diabetic Retinopathy
	Study
	. Fluorescein angiography
	Fundus Autofluorescence
	. Ganglion Cell Layer
	. Hypoxia-inducible factor 1-alpha
HM	
	. Hemiretinal vein occlusion
HTN	
ICP	. Intra Cranial Pressure

Tist of Abbreviations cont...

Abb.	Full term
IL	Interlouking
	Internal Limiting Membrane
	Inferial Limiting Membrane Inferior Nasal Artery
	Inner Nuclear Layer
	Intra Ocular Pressure
	Inner Plexiform Layer
	Inner segment/outer segment
15/05	photoreceptor junction
ITA	Inferior Temporal Artery
	Juxtafoveal Retinal Telangiectasis
	Longitudinal Scan
ME	9
MHz	
	Neodymium:Yttrium Aluminum Garnet
NFL	
	Optical Coherence Tomography
	Outer Nuclear Layer
	Outer Plexiform Layer
PE	Posterior to the Equator
PHT	Posterior hyaloid Traction
PL	Perception of Light
PRP	Panretinal Photocoagulation
PVD	Posterior Vitreous Detachment
	Relative afferent pupillary defect
RMG	Retinal Muller Glial cells
ROP	Retinopathy of prematurity
	Retinitis Pigmentosa
	Retinal pigment epithelium
RVO	Retinal Vein Occlusion
SD	-
	Super Luminescent Diode
	Scanning laser ophthalmoscopy
SNA	Superior Nasal Artery

Tist of Abbreviations cont...

Abb.	Full term	
SRD	Serous Retinal Detachment	
SRF	Subretinal Fluid	
STA	Superior Temporal Artery	
T	Transverse Scan	
TD	Time Domain	
TNF-α	Tumor necrosis factor-α	
TRD	Tractional Retinal detachment	
U/S	Ultrasound	
UVR	Ultra Violet Rays	
VA	Visual Acuity	
VEGF	Vascular endothelial growth factor	
	Vitreomacular Traction Syndrome	

Introduction

Macular edema is a common phenomenon in various diseases where fluid accumulates in between the retinal cells. The fluid originates from the intravascular compartment. The focal, diffuse, and cystic forms are all characterized by extracellular accumulation of fluid, specifically in Henle's layer and the inner nuclear layer of the retina. The compartmentalization of the accumulated fluid is likely to be due in part to the relative barrier properties of the inner and outer plexiform layers (Tranos et al., 2004).

The classic pattern of cystoid macular edema (CME) with the petaloid appearance originating from the fluorescein leakage from perifoveal capillaries may be seen in cases of advanced edema of various origins. This includes postsurgical CME as well as CME associated with one of the following conditions: diabetes, vascular occlusion, hypertensive retinopathy, epiretinal membranes, intraocular tumors (e.g., melanoma, choroidal hemangioma), intraocular inflammation (e.g., pars planitis), macroaneurysm, retinitis pigmentosa, choroidal neovascularization, and radiation retinopathy (*Tranos et al.*, 2004).

Cystoid macular edema may have severe implications for the function of the retina, including decreased visual acuity and contrast sensitivity. Acute or chronic edema causes anatomical disruption that may result in cellular dysfunction and death. Treatment of CME is important because chronic edema may result in degenerative changes in the macula and permanent loss of vision (*Brown et al., 2011*). In addition, large cystic changes in the retina may lead to thinning and loss of inner retinal tissue, or the formation of lamellar hole (*Tsukada et al., 2011*).

Early detection of CME is critical for diagnosis and management. Traditional methods of assessing macular edema include contact and noncontact slit lamp biomicroscopy, indirect ophthalmoscopy, fluorescein angiography (FA), and fundus stereo photography. However the interpretation of their results can be subjective, and subtle changes in retinal thickness in early CME may not be evident.

Optical coherence tomography (OCT) correlates well with retinal histology and can be used to quantitatively and qualitatively monitor retinal thickness over time. Compared to biomicroscopy and FA, OCT is more sensitive in detection of macular edema and subretinal fluid, and subclinical macular edema is often only detected by OCT (*Hee et al.*, 1995a).

Fluorescein angiography and OCT have limitations. Both tests require the ocular media to be of sufficient clarity to image the retina. Yet in certain patients, opacities in the ocular media limit biomicroscopy, FA, and OCT. Furthermore, a high degree of patient cooperation is required to ensure reliable and accurate testing. However, certain patients, such as children,

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often cannot tolerate a FA or follow the specific fixation instructions for OCT testing.

Ophthalmic ultrasonography is a well-accepted noninvasive diagnostic tool. Ultrasonography has the advantage of reliably imaging the posterior segment regardless of the ocular media status. Furthermore, ultrasonography is less dependent on patient cooperation for reliable testing than either FA or OCT (*Fisher et al.*, *1991*).