Abnormal Ocular Findings in Chronic Renal Failure Patients on Hemodialysis

Thesis

Submitted for partial fulfillment of Master Degree in Ophthalmology

Presented by

Aml Mahmoud Ahmed El-Azab

M.B.B.Ch

Supervised by

Prof. Dr. Dina Ezzat Mansour

Professor of Ophthalmology

Faculty of Medicine - Ain-shams University

Dr. Ahmed Mohamed El-Bayomi

Lecturer of Ophthalmology
Faculty of Medicine - Ain-shams University

Faculty of Medicine
Ain Shams University
2018



Acknowledgement

First of all, all gratitude is due to Allah almighty for blessing this work, until it has reached its end.

Words do fail to express my deepest gratitude and appreciation to Prof. Dr. Dina Ezzat Mansour, Professor of ophthalmology, Faculty of medicine, Ain Shams University for her guidance, advices and marvelous support to conduct this work.

Great thanks to Dr. Ahmed Mohamed El-Bayomi, Lecturer of Ophthalmology, Faculty of Medicine, Ain-shams University, for his precious time, he gave me for the completion of this work.

Great thanks to my colleagues in Dialysis unit at Agouza hospital for their co-operation and help.

Thanks to my dearest friend **Dr**. **Eman Nagy Ahmed** who was always there giving me all kinds of support throughout this work.

I cannot forget to express my thanks to my lovely family: my father, my brothers and sister.

Lastly, but not the least, I dedicate this work to my long-sleeping angel; Mom who taught me the language of life.

Aml El-Azab (2018)

List of Abbreviations

AAO American academy of ophthalmology

ACR Albumin creatinine ratio

ADPKD Autosomal dominant polycystic

kidney disease

AMD Age-related macular degeneration

BCVA Best corrected visual acuity

BUT Break up time

CFT Central foveal thickness

CKD Chronic kidney disease

CRF Chronic renal failure

CSME clinically significant macular edema

DM Diabetes mellitus

DME Diabetic macular edema

eGFR Estimated glomerular filtration rate

EOM Extra ocular muscles

ESRD End stage renal disease

ETDRS Early Treatment Diabetic Retinopathy

Study

GN Glomerulonephritis

GPA Granulomatosis with Polyangiitis

HD Haemodialysis

HTN Hypertension

IOP Intra-ocular pressure

IRMA Intraretinal microvascular

abnormalities

MPA Microscopic Polyangitis

NPDR Non proliferative diabetic retinopathy
NSAIDs Non steroidal anti-inflammatory drugs

NSRD Neurosensory retinal detachment

NVD New vessels on optic disc

NVE New vessels elsewhere

OCT Optical coherence tomography

OPP Ocular perfusion pressure

PDR Proliferative diabetic retinopathy

PHO Primary hyperoxaluria

PTDs Post trachomatous degenerations

RNFL retinal nerve fiber layer

RPE retinal pigment epithelium

SD Standard deviation

SD-OCT Spectral domain optical coherence

tomography

SLE Systemic lupus erythematosus

SPSS Statistical Package for Social Science

SRD Serous retinal detachment

TINU Tubulo-interstitial Nephritis and

Uveitis Syndrome

USRDS United States Renal Data System

VA Visual acuity

VMT Vitreo-macular traction

List of Tables

Page	Subject	
Review of the literature		
10	Table (I): International Clinical Diabetic	
	Retinopathy Disease Severity Scale	
11	Table (II): International Diabetic Macular	
	Clinical Edema Severity Scale	
18	Table (III): Other autoimmune diseases linking	
	kidney and eye	
Results		
30	Table (1): Demographic data of studied patients	
31	Table (2): Medical history and past ophthalmic	
	history of studied patients	
32	Table (3): BCVA in the eyes of studied groups	
33	Table (4): Causes of visual impairment in the	
	eyes with BCVA <6/18	
33	Table (5): IOP measurements after hemodialysis	
	session	
34	Table (6): Abnormal anterior segment findings as	
	regard 1ry diseases causing chronic kidney disease	

37	Table (7): Abnormal posterior segment findings in studied group
40	Table (8): Abnormal OCT finding in studied group.
42	Table (9): Comparison between CFT as regard 1ry cause of CKD

List of Figures

Page	Subject	
Review of the literature		
6	Figure (I): Band keratopathy in a patient on chronic HD for 9 years	
7	Figure (II): A picture of pinguecla in renal patient on dialysis.	
23	Figure (III) Retinal thickness map: Top left image shows the color-coded macular thickness map showing thickening at the macular region and macular thickness values are seen in 1 mm, 3 mm, and 6 mm circles in the top right image. Bottom image shows the corresponding spectral domain optical coherence tomography image	
	Results	
31	Figure (1): Causes of CKD in studied patients	
35	Figure (2): Photo of one of the studied patients showing lid edema.	
35	Figure (3): Photo of one of the studied patients showing pterygium in the right eye.	
36	Figure (4): Photo of one of the studied patients showing yellow sclera.	
36	Figure (5): Photo of one of the studied patients showing senile cortical cataract.	

38	Figure (6): A fundus photo of one eye in our study showing NPDR without CSME (microaneurysms) in the left eye.
38	Figure (7): A fundus photo of one eye in our study showing NPDR with CSME (microaneurysms, retinal hemorrhage, exudates) in the right eye.
39	Figure (8): A fundus photo of one eye in our study showing PDR with CSME (microaneurysms, flame shaped hges, exudates) and hypertensive retinopathy (attenuated arterioles and dilated tortous veins, A-V crossing) in the right eye.
39	Figure (9): A fundus photo of one eye in our study showing dry macular degeneration (drusen)in the right eye.
40	Figure (10): A fundus photo of one eye in our study showing macular hge in hypertensive patient in the right eye.
41	Figure (11): An OCT photo of one eye in our study showing diabetic macular edema (multiple retinal cysts, subretinal fluid, exudates).
41	Figure (12): An OCT photo of one eye in our study showing age related macular degeneration (drusen)
42	Figure (13): An OCT photo of one eye in our study showing macular hemorrhage

43	Figure (14): A photo of color macular thickness
	map of one eye of diabetic patient in our study
	showing increased macular thickness shown by
	color code.
44	Figure (15): A photo of OCT section showing
	DME with impending macular hole and
	subretinal fluid with color macular thickness map
	of one eye of diabetic patient in our study
	showing increased macular thickness.
44	Figure (16) A photo of OCT macula section
	showing well defined retinal layers with color
	macular thickness map of one eye of
	hypertensive patient in our study showing
	decrease in macular thickness.
45	Figure (17) A photo of OCT macula section
	showing retinal layers with color macular
	thickness map of one eye of non-diabetic non-
	hypertensive patient with unknown cause of
	CKD in our study showing normal retinal layers
	and normal macular thickness.

List of Contents

Page	Subject
1	Introduction& Aim of the work
4	Review of literature:
	Ocular manifestations of CRF and effect of hemodialysis on the eye
	Oculorenal diseases
	• OCT
26	Subjects and methods
30	Results
46	Discussion
55	Summary& Conclusion
58	Recommendations
59	References
	Arabic Summary

ABSTRACT

Background: chronic kidney disease (CKD) is a global public health problem and diabetes is a leading cause for it. Those patients with renal disease (ESRD) are generally treated hemodialysis (HD). HD causes numerous metabolic changes which in turn induce osmotic changes in blood and extracellular fluids including the agueous and vitreous humor, so there are many ocular **ESRD** either due abnormalities in patients to common pathophysiological mechanisms and risk factors between eye and kidney diseases or due to hemodialysis itself. Aim of the Work: the aim of this study is to evaluate ocular findings in CKD patients on hemodialysis, including the best-corrected visual acuity (BCVA), measurement of intraocular pressure (IOP), corneal, conjunctival, tear film changes, lens and fundus changes. Patients and Methods: this cross sectional observational study was carried out on twenty three patients (6 males and 17 females) with 44 eyes, with chronic kidney disease on chronic hemodialysis for 2-10 years in the Dialysis Unit at Agouza Specialized Hospital in the period from November 2017 to April 2018. The hemodialysis duration was approximately four hours three times per week. Results: forty three percent of total 44 eyes had BCVA > 6/18. As regard anterior segment, we found yellow sclera was the commonest finding (75%), cataract (61.4%), lid edema (56.8%), dry eye (43.2%), conjunctival congestion (47.7%), and pterygium (13.6%). As regard posterior segment; Diabetic retinopathy was the commonest finding (38.6%), hypertensive retinopathy (4.5%), AMD (2.3%), macular hemorrhage (2.3%). Regarding OCT findings, there was statistically significant difference between eyes of diabetic, hypertensive and other patients as regard central foveal thickness. **Conclusion:** our results demonstrate high prevalence of abnormal ocular findings in chronic renal failure patients on hemodialysis. So those patients should be evaluated regularly by the ophthalmologist.

Keywords: Chronic Renal Failure, Hemodialysis, Ocular Findings,

Central Foveal Thickness, OCT

Introduction

Chronic kidney disease (CKD) is a slow progressive loss of kidney function over a period of several years, which also is the presence of either reduced renal function as evaluated by estimated glomerular filtration rate (eGFR $<60 \,\mathrm{ml/minute/1.73\,m^2}$) or albuminuria evaluated by urinary albumin to creatinine ratio (ACR $\ge 30 \,\mathrm{mg/g}$) (1).

Chronic kidney disease has become a major public health problem worldwide and has been associated with premature morbidity and mortality ⁽²⁾.

Chronic kidney disease has numerous causes. The most common causes of CKD are diabetes mellitus and long-term uncontrolled hypertension ⁽³⁾. Overuse of common drugs such as ibuprofen, acetaminophen can also cause chronic kidney disease ⁽⁴⁾.

Chronic renal failure patients have had a wide range of findings, including refractive changes, dry eye, increased tear osmolarity, conjunctival calcium deposits, band keratopathy, corneal endothelium changes and lenticular opacity ⁽⁵⁾.

Patients with chronic kidney disease are generally treated using a blood filtration mechanism such as hemodialysis. The primary objective of hemodialysis is to correct the composition and volume of body fluids. Correction of body fluid aims to resolve the excessive accumulation and abnormal distribution of body fluids and

is concerned with the change in plasma colloid osmotic pressure during haemodialysis ⁽⁶⁾.

Although hemodialysis is relatively effective, it has certain impairments that persist over time and many patients experience multi-organ changes. Hemodialysis usually reduces body weight and blood pressure and increases the plasma/interstitial osmotic pressure in patients with ESRD ⁽⁷⁾.

During hemodialysis, numerous metabolic parameters including blood urea, sodium, potassium and sugar will be changed ⁽⁸⁾, which also induce osmotic changes in blood and extracellular fluids. This includes the aqueous and vitreous humor, any change in osmotic pressure of these fluids could affect the refractive status ⁽⁹⁾.

The relationship between chronic hemodialysis and intra-ocular pressure (IOP) has been widely investigated in the past. There are several studies, which demonstrate a significantly increased IOP and decreased OPP occur during HD, bringing both to levels that increase the risk of glaucoma development and progression ⁽¹⁰⁾. In contrast, other studies have demonstrated the opposite effect ⁽¹¹⁾, while some studies reported no IOP change during HD ⁽¹²⁾.

Ocular posterior segment changes such as, retinopathy secondary to diabetes mellitus, hypertension, anemia, and uremia, are also observed frequently in CRF patients ⁽¹³⁾.

Pahor et al. found that CFT was significantly thinner in hemodialysis patients than normal healthy subjects ⁽¹⁴⁾.

Aim of the work

The aim of this study is to evaluate abnormal ocular findings in CKD patients on hemodialysis, including the best-corrected visual acuity (BCVA), measurement of intraocular pressure (IOP), corneal, conjunctival, tear film changes, lens and fundus changes.

Ocular Manifestations of CRF and Effect of Hemodialysis on The Eye

The eye shares striking structural, developmental and genetic pathways with the kidney, suggesting that kidney disease and ocular disease may be closely linked. Richard Bright, an early pioneer in morbid anatomy and clinical signs and symptoms of kidney disease first reported the association between renal disease and blindness in 1836 (15).

Renal microvascular pathology is thought to play an important role in the development of renal insufficiency. The assessment of renal vascular pathology requires invasive procedures. The retinal vasculature, conversely, can be observed non-invasively in humans and therefore offers a unique opportunity to explore the association between systemic microvascular disease and renal function. These organs share pathophysiological mechanisms, such as endothelial dysfunction and the inflammatory process, leading to circulatory abnormalities and decreased vascular reactivity ⁽¹⁶⁾.

We are going to discuss abnormal ocular findings associated with hemodalysis:

1- Refractive state:

Hemodialysis patients, especially the elderly, have visual acuity (VA) levels much lower than their agematched counterparts (17).