Prevalence of Hepatitis C Virus Antibodies in Hemodialysis Patients in Kafr El- Sheikh Governorate (Sector B)

Chesis

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By

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List of Contents

Title		
♦ List of Abbreviations	II	
♦ List of figures	V	
♦ List of Tables	VI	
♦ Introduction	1	
♦ Aim of the Work	5	
• Review of the Literature		
• Chapter (1): Hepatitis C Virus Overview	6	
 Chapter (2): HCV and Chronic Kidney Diseases 	51	
 Chapter (3): Methods of Infection Control in 		
Hemodialysis Units	79	
♦ Patients and Methods		
♦ Results		
♦ Discussion		
Summary and Conclusion		
♦ Recommendations		
♦ References		
♦ Arabic Summary		

List of Abbreviations

ADH Antidiuretic hormone

ALT Alanine Aminotransferases

ANA Antinucler antibody

ANP Atrial natriuretic peptide ASK Atrophic single kidney

AST Aspartate Aminotransferases

AVF ArterioVenous Fistula
AVG ArterioVenous Graft
BCM Bio compatible membrane
BICM Bio incompatible membrane
BRA Bilateral renal atrophy

C Complement

C.ANCA Cytoplasmic pattern antinutrophil

cytoplasmic Abs

CAPD Continous Ambulatory Peritoneal Dialysis

CD Cluster of Differentiation

CDC Center for Diseases Control and Prevention

CHC Chronic hepatitis C

CKD Chronic kidney disease

CRRT continuous renal replacement therapy

CRYO cryoglobulinemia

DM Diabetes mellitus

DMSADi mercapto succinic acidDNADeoxyribo neucleic acidEEnvelop glycoproteins

EHMs Extrahepatic manifestations

EIA Enzyme immunoassay

ELISA Enzyme Linked Immunosorbent Assay

EPA Environmental Protection Agency

ESRD End stage renal disease

Et Endothelin

ETR End-of treatment response

EVR Early viral response

FDA Food and Drug Administration
GFR glomerular filtration rate

GN Glomerulonephritis

GSH Glutathione

HBV Hepatitis B Virus

HCC Hepatocellular carcinoma

HCV Hepatitis C Virus

HCWs Health care workers

HD Hemodialysis

HGF Hepatocyte growth factor

HIV Human Immunodeficiency Virus

HRS Hepato renal syndrome

HTN Hypertension

IB Immuno blot

ICU intensive care unit

IFNs Interferons

Ig Immunoglobulin

KDIGO Kidney Disease Improving Global Outcome

LDL Low Density Lipoprotien

MALT Mucosa associated lymphoma tumor

MGMyasthenia graviesmmMultible mylomaMOFMultiorgan faliur

MPGN Membranoproliferative glomerulonephritis

MRI Magnetic resonant image

NIH National Institute of Health

NIPD Nocturnal intermittent peritoneal dialysis

NO Nitric oxid

OBS Obstructive uropathy

PCR Polymerase Chain Reaction
PCT Porphyria cutenea tarda
PEG-IFN Pegylated-Interferon

r.HUEPO Recombinant human erythropoietin

RAS Renal artery stenosis

RBV Ribavirin

Rt-PCR Reverse transcriptase Poly chain reaction

RNA Ribo neucleic acid RT Reverse Transcriptase

RVR Rapid Virological Response
SLE Systemic Lupus Erythematosis
SLED Sustained law efficiency dialysis

SR Scavenger Receptor
SS Sjogren Syndrome

STD Sexually Transmitted Diseases

SVR Sustained viral response

TMA Transcription mediated amplification

TPN Total parentral nutrition

UOP Urin out put

UTI Urinary tract infectionWHO World Health Organization

List of Figures

Fig. No.	Title	Page No.	
1	HCV viral component.	8	
2	Simplified diagram of HCV replication cycle.	10	
3	Prevalence of HCV and HBV in studied group.	101	
4	HCV seroconvertion.	103	
5	Causes of chronic renal failure.	104	

List of Tables

Table No.	Title	Page No.
1	Annual reports of Egyptian renal registry from 1996 to 2008.	56
2	Brief details of studies showing benefit of isolation of HCV patients in HD units.	65
3	Recommended Treatment of HCV Infection in Patients with CKD and Their Associated Adverse Events.	67
4	Questionnaire form in patients &methods by history taking from all patients.	98
5	Age, sex, HBV and HCV status of the studied group and overall prevalence of HCV Ab.	100
6	Risk factors for HCV Ab seroconvertion.	102
7	Causes of chronic renal failure in the studied group.	105
8	Comparison between HCV sero-converters and free cases regarding age and Duration of dialysis free from HCV.	`106
9	Categorical risk factors for seroconversion – Univariate analysis.	109
10	Risk factors for seroconversion – Multivariable analysis using logistic regression.	110



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Introduction

Hepatitis C virus (HCV) is a small (40 to 60 nanometers in diameter), enveloped, single-stranded RNA virus of the family Flaviviridae and genus hepacivirus. Because the virus mutates rapidly, changes in the envelope proteins may help it to evade the immune system. There are at least six major genotypes and more than 50 subtypes of HCV. The different genotypes have different geographic distributions (*Meyers et al.*, 2003).

HCV infection is the most common chronic bloodborne infection in the United States; approximately 3.2 million persons are chronically infected. Although HCV is not efficiently transmitted sexually, persons at risk for infection through injection drug use might seek care in sexual transmitted diseases (STD) treatment facilities, human immune-deficiency virus (HIV) counseling and testing facilities, correctional facilities, drug treatment facilities, and other public health settings where STD and HIV prevention and control services are available (*Fissell et al.*, 2004).

Patients on hemodialysis (HD) have a high risk of acquiring HCV infection. Transfusion of unscreened blood, duration of dialysis and nosocomial transmission within HD

units is implicated as the main transmission routes of HCV infection in HD patients (Meyers et al., 2003).

HCV infection in patients on HD has been associated with greater morbidity and mortality (*Froio et al.*, 2003).

Virological diagnosis and monitoring of HCV infection are based on two categories of laboratory tests, namely serologic assay detecting specific antibody to HCV (indirect test) and assay that can detect, quantify or characterize the component of HCV viral particles (direct tests), such as HCV RNA (*Lok et al.*, 1998).

Antibody detection tests are considered important tools to assess the magnitude of HCV infection in patients on HD. The window phase in HD patients can be longer as these patients are immunocompromised and the anti HCV Enzyme Linked ImmunoSorbent Assay (ELISA) test alone may fail to detect the infected patients in the acute phase of the disease (*Lok et al.*, 1998).

Chronic hepatitis C varies greatly in its course and outcome. At one end of the spectrum are patients who have no signs or symptoms of liver disease and completely normal levels of serum liver enzymes. Liver biopsy usually shows some degree of chronic hepatitis, but the degree of injury is usually mild, and the overall prognosis may be good (*Fabrizi et al.*, 2002).

At the other end of the spectrum are patients with severe hepatitis C who have symptoms, HCV RNA in serum, and elevated serum liver enzymes, and who ultimately develop cirrhosis and end stage liver disease. In the middle of the spectrum are many patients who have few or no symptoms, mild to moderate elevations in liver enzymes, and an uncertain prognosis. So, the treatment may be in the form of symptomatic liver support, interferon, ribavirin and even liver transplantation (*Fabrizi et al.*, 2002).

At present, the main tools of preventing new cases of hepatitis C are to screen the blood supply, encourage health professionals to take precautions when handling blood and body fluids, and inform people about high-risk behaviours. Programs to promote needle exchange offer some hope of decreasing the spread of hepatitis C among injection drug users (*Moreira et al.*, 2003).

Effective screening of blood and blood products virtually eliminated HCV transmission by blood transfusions a decade ago, and a subsequent decline in HCV incidence and prevalence within HD units in developed countries occurred (*Fabrizi et al.*, 2010).

Vaccines and immunoglobulin (Ig) products do not exist for hepatitis C, and development seems unlikely in the near future because these products would require antibodies to all

Datroduction

the genotypes and variants of hepatitis C. Nevertheless, advances in immunology and innovative approaches to immunization make it likely that some form of vaccine for hepatitis C will eventually be developed (*Moreira et al.*, 2003).

Aim of the Work

To study the prevalence of hepatitis C antibodies among the patients undergoing regular HD in Kafr El- Sheikh Governorate Sector (B) by a questionnaire form.

They include patients in the following hospitals.

Kafr El- Sheikh city, Biala city, El Hamol city, El Riad city and Baltem city in the period of 1/1/2012 to 30/7/2012.

Chapter (1)

Hepatitis C Virus Overview

Hepatitis C virus (HCV) is the leading cause of chronic liver disease worldwide. It is estimated that about 170 million people are chronically infected with HCV, which is a major cause of cirrhosis and hepatocellular carcinoma (*Boyer & Marcellin*, 2000).

Hepatitis C virus is the single most important cause of liver disease in Egypt (*El-Zayadi* · 1997; *Habib et al.*, 2001), where high HCV rates were reported among several population groups reaching up to 20% (*Mohamed et al.*, 2006).

HCV is a small, enveloped virus with a single stranded RNA genome of positive polarity. Based on its features, it has been classified as the sole member of the genus hepacivirus in the Flaviviridae family, which include 2 other genera with important human and animal pathogens: Flaviviruses (e.g. West Nile virus, dengue virus and yellow fever virus) and pestiviruses (e.g. bovine viraldiarrhea virus and classical swine fever virus) (*Ivanyi-Nagy et al.*, 2006).

HEPATITIS C VIRUS (HCV):

Hepatitis C virus (HCV) is a single stranded ribonucleic acid (RNA) virus that replicates at a rapid rate. As a