

REVIEW

Hepatitis C and kidney disease: A narrative review



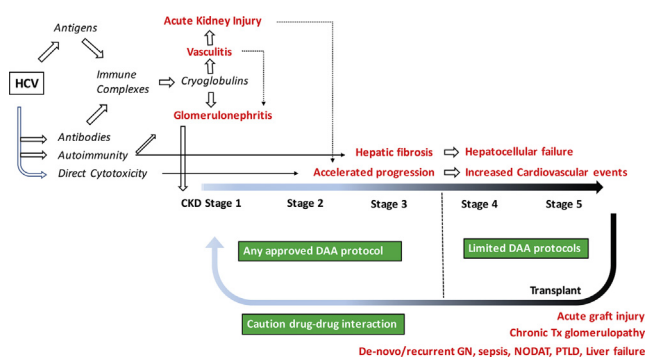
Rashad S. Barsoum^{a,b,*}, Emad A. William^{b,c}, Soha S. Khalil^b

^a Kasr-El-Aini Medical School, Cairo University, Cairo, Egypt

^b The Cairo Kidney Center, Cairo, Egypt

^c National Research Centre, Cairo, Egypt

GRAPHICAL ABSTRACT



ARTICLE INFO

Article history:

Received 19 April 2016

Received in revised form 7 July 2016

Accepted 17 July 2016

Available online 26 July 2016

Keywords:

Acute kidney injury

Chronic kidney disease

Glomerulonephritis

ABSTRACT

Hepatitis-C (HCV) infection can induce kidney injury, mostly due to formation of immune-complexes and cryoglobulins, and possibly to a direct cytopathic effect. It may cause acute kidney injury (AKI) as a part of systemic vasculitis, and augments the risk of AKI due to other etiologies. It is responsible for mesangiocapillary or membranous glomerulonephritis, and accelerates the progression of chronic kidney disease due to other causes. HCV infection increases cardiovascular and liver-related mortality in patients on regular dialysis. HCV-infected patients are at increased risk of acute post-transplant complications. Long-term graft survival is compromised by recurrent or de novo glomerulonephritis, or chronic transplant glomerulopathy. Patient survival is challenged by increased incidence of diabetes, sepsis, post-transplant lymphoproliferative disease, and liver failure. Effective and safe directly acting antiviral agents (DAAs)

* Corresponding author. Fax: +20 225790267.

E-mail address: Rashad.barsoum@gmail.com (R.S. Barsoum).

Peer review under responsibility of Cairo University.



Production and hosting by Elsevier

<http://dx.doi.org/10.1016/j.jare.2016.07.004>

2090-1232 © 2016 Production and hosting by Elsevier B.V. on behalf of Cairo University.

This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

List of abbreviations

AA-protein	Amyloid-A protein	HCV +ve	HCV infected
AASLD	American Association For The Study Of Liver Disease	HCV -ve	HCV non-infected
ADA	American Diabetes Association	HIV	Human Immunodeficiency virus
AKD	Acute kidney disease	IFN- α	Interferon-alpha
AKI	Acute kidney injury	IgA, IgG, IgM	Immunoglobulins A, G and M (respectively)
C1q, C3a, C4, C5a, C5-9	Respective complement components	κ -RF	Kappa-Rheumatoid factor
CKD	Chronic Kidney disease	MGN	Membranous glomerulonephritis
CLD	Chronic liver disease	MPGN	Membranoproliferative (Mesangiocapillary) glomerulonephritis
CYP-450	Cytochrome P-450	mTOR	Mammalian target of rapamycin
D +ve	HCV positive donor	NHL	Non-Hodgkin lymphoma
DAAs	Direct-acting antivirals	NODAT	New-onset diabetes after transplantation (Post-transplant diabetes mellitus)
DNA	Desoxyribonucleic acid	NSx	Non-structural viral protein-number (x)"
EASL	European Association For The Study Of The Liver	PCRC	Polymerase chain reaction for Hepatitis C virus
eGFR	Estimated glomerular filtration rate	PTLD	Post-transplant lympho-proliferative disorder
ELISA	Enzyme-linked immunosorbent assay	R +ve	HCV positive recipient
ESKD	End-stage kidney disease	RBV	Ribavirin
FCH	Fibrosing cholestatic hepatitis	RCT	Randomized controlled trial
FDA	Food and Drug Administration	RDT	Regular dialysis treatment
FSGS	Focal segmental glomerulosclerosis	RF	Rheumatoid Factor
GN	Glomerulonephritis	RNA	Ribonucleic Acid
GT	Genotype	RR	Relative Risk
HBV	Hepatitis B virus	SVRx	Sustained Viral Response In (x) weeks
HCV	Hepatitis C virus	USRDS	United States Renal Data System

Dialysis

Renal transplantation

Direct-acting antivirals

are currently available for treatment at different stages of kidney disease. However, the relative shortage of DAAs in countries where HCV is highly endemic imposes a need for treatment-prioritization, for which a scoring system is proposed in this review. It is concluded that the thoughtful use of DAAs, will result in a significant change in the epidemiology and clinical profiles of kidney disease, as well as improvement of dialysis and transplant outcomes, in endemic areas.

© 2016 Production and hosting by Elsevier B.V. on behalf of Cairo University. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).



Rashad Barsoum Emeritus Professor and former chairman of Internal Medicine and chief of Nephrology at Cairo University. He authored 44 chapters on kidney diseases in 25 international textbooks. He published \approx 200 papers with over 2500 citations. He served on the editorial boards of 18 and reviewer for 35 international journals. He holds the Egyptian First Class Order of Arts and Sciences and was the recipient of the Egyptian Nile Award and State Appreciation Prize, the ISN Roscoe

Robinson Award, the International Award of the USA National Kidney Foundation Tarek Suhaimat Award of the ASNRT, Pioneer Award of the ISN, and many others.



Emad A. William Graduated from Faculty of Medicine, Ain Shams University, Egypt, in 1997, obtained his Master's degree in Internal Medicine in 2004, and the Doctorate degree from the same University in 2011. He currently holds an academic position as a research lecturer at the National Research Center.

He is also a Clinical Consultant and Head of the Clinical Research Unit at the Cairo Kidney Center, Egypt. His main clinical expertise is in Clinical Nephrology, Dialysis and Transplantation, and his main research interest is renal transplantation.

Download English Version:

<https://daneshyari.com/en/article/5022830>

Download Persian Version:

<https://daneshyari.com/article/5022830>

[Daneshyari.com](https://daneshyari.com)