

Clinical Assessment of Pharmacotherapeutic Plan for Virologic Response of Hepatitis C Virus

Faiza Naeem*

Institute of Pharmacy, Lahore College for Women University, Pakistan

***Corresponding author:** Faiza Naeem, Institute of Pharmacy, Lahore College for Women University, Lahore, Pakistan, Email: faizanaeem12@hotmail.com

Case report

Volume 3 Issue 4 Received Date: August 05, 2019 Published Date: September 06, 2019 DOI: 10.23880/vij-16000220

Abstract

Hepatitis C is triggered by the presence of hepatitis C virus (HCV) where acute and chronic hepatitis causes mild to severe illness lasting for few weeks to lifelong illness. The HCV is a bloodborne virus having most common modes of infection through exposure to even small quantities of contaminated blood and this may trigger through contaminated injection drug use, unsafe injection practices, hazardous health care, transfusion of unscreened blood or blood products and sexual practices. In 2016, WHO estimated roughly 399, 000 people died from HCV frequently from liver cirrhosis and hepatocellular carcinoma (HCC) which is primary liver cancer. Presently, there is no effective vaccine in contradiction of hepatitis C. Fever, fatigue, decreased appetite, abdominal pain and joint pain are common clinical presentations. Investigation for anti-HCV antibodies alongwith serological testing identifies individuals who have been infected with HCV. Ribavirin and sofosbuvir are the main anti-viral drugs which can treat the hepatitis C. In this case scenario, referred patient of 40 years old female is suffering from hepatitis C having severe abdominal pain along with generalized body weight. Confirmation of hepatitis C for this patient was done by anti-HCV antibodies, LFT report and ultrasound which also reveal fatty liver. Symptomatically, she was treated by antiviral drug therapy which finally stabilizes her severe abdominal pain and other clinical presentations along with the condition of hepatitis C.

Keywords: Hepatitis C; Pharmacotherapeutic Plan; Clinical Assessment; Ribavirin; Sofosbuvir

Abbreviations: HCV: Hepatitis C Virus; HIV: Human Immunodeficiency Virus; INF: Interferon; HCC: Hepatocellular Carcinoma.

Introduction

Hepatitis C virus (HCV) is a foremost leading reason of liver diseases globally and an impending source of

significant morbidity as well as mortality in future [1]. HCV is considered in Hepacivirus genus and Flaviviridae family [2]. Up to 7 million individuals internationally are infected with both of the viruses including human immunodeficiency virus (HIV) & heaptitis C virus (HCV) [3]. HCV infection is connected with higher rates of liver cirrhosis, fibrosis, hepatocellular carcinoma and overall higher mortality [4]. Re-use of contaminated and inadequately sterilized syringes as well as needles used in medical, paramedical & dental measures [5]. The annual rate for HCC development between patients of liver cirrhosis is presently predicted as 1.6% [6]. Combination of sofosbuvir with ribavirin for 24 weeks has also revealed proper treatment for patients with genotype 1 of HCV [4]. Ribavirin when used in combination with pegylated interferon alfa (INF), leads to improved restoration of patient's immune response from HCV [7].

Case Presentations

A 40 years old female comes to physician for the clinical presentations of severe abdominal pain, generalized body weakness, Sore muscles and anorexia (decreased appetite). She is also suffering from fever. Her past medical history includes peptic ulcer. She frequently

Virology & Immunology Journal

uses Panadol, in case of mild pain and fever. She belongs to a Poor Socio-economic status.

On Examination (O/E)

Sign	Normal Range	Results	
Blood Pressure	120/80 mmHg	110/80 mmHg	
Heart Rate	72 BPM	78 beats/min	
Temperature	98°F	102°F	

Table 1: Vital signs of patient.

Laboratory Findings

Ultra sound of patient shown that the size of liver is 15.8cm and also fatty liver is present. Anti-HCV test is positive for this patient. Other diagnostic tests are described in Tables 2 and 3.

Tests	Normal Range	Unit	Values	Comment
Wbcs	4-11	x 10 ³ /μL	12.7	Above Normal
Total Rbc Count	4.0-5.5	x 10 ⁶ /μL	4.5	Normal
Hemoglobin	13-17 (MALE),	a /dI	13	Normal
	12-15 (FEMALE)	g/uL		
Platelets	150-400	x 10 ³ /μL	230	Normal
HCT (PCV)	40-75	%	40	Normal
MCV	20-45	fl	50	Above Normal
МСН	1-20	Pg	22	Above Normal
МСНС	65-110	%	33	Normal

Table 2: Complete blood count of (CBC) patient.

Liver Function Test							
Test	Normal Range	Unit	Value	Comments			
Bilirubin Total	0.2-1.2	(mg/dl)	1.4	Above Normal			
Bilirubin conjugated	< 0.5	(mg/dl)	0.7 Above Nor				
Bilirubin unconjugated	0.1-1	(mg/dl)	0.5	Normal			
ALT (SGPT)	5-55	(U/L)	73	Above Normal			
AST (SGOT)	5-35	(U/L)	77	Above Normal			
Alkaline Phosphatase	40-150	(U/L)	120	Normal			
GAMMA G.T.	10-64	(U/L)	72	Above Normal			
Total protein	6-8.5	(g/dl)	8.1	Normal			
Serum Albumin	3.5-5.0	(g/dl)	3.6	Normal			
Serum Globulin	1.8-3.4	(g/dl)	3.9	Above Normal			
A/G Ratio	1.2-2.2	(g/dl)	0.9	Below Normal			

Table 3: Liver function test (LFT) of patient.

Current Prescribed Medication						
Brands	Generics	Dosage form	Frequency	Dose	Indications	
Sovaldi,	Sofosbuvir	Tablet	OD	400mg	HCV infection	
Daclavia	Daclatasvir	Tablet	OD	60mg	HCV infection	
Risek	Omeprazole	Capsule	OD	40mg	Treat Gastric disturbance	

Pharmacotherapeutic Plan

Table 4: Prescribed Medication for hepatitis C treatment.

Clinical Pharmacist Interventions

Tab. Sovaldi & Daclavia are more effective in patients with compromised liver function, when given with Ribavarin, 800 mg OD. Hence, Ribavirin 800 mg OD (400mg, 2 caps. BD) was added to prescribed pharmacotherapeutic plan of this patient.

Care Plan

- Follow up for routine tests and examination is advised.
- LFTs and other lab tests to be repeated after every 3 months.
- Adherence to therapy is advised and assessed at each follow up.

Follow Up Requirement

To assess the following parameters

- HCV progression into HCC (hepatocellular carcinoma)
- Liver Function test
- Adherence to therapy

Discussion

Hepatitis C virus (HCV) contaminates an expected 170 million individuals globally and therefore epitomizes a viral disease as pandemic case [8]. According to WHO Report in 2001, chronic liver diseases were accountable for 1 to 4 million deaths, comprising 796,000 due to liver cirrhosis and 616,000 deaths due to liver cancer [9]. In this case study, 40 years old patient is complaining about the severe abdominal pain, generalized body weakness, Sore muscles and anorexia. Her previous history is about peptic ulcer which has been cured after triple therapy.

According to clinical assessment Under prescription of HCV medications for patient was done. So, the first priority should be given to prescribing all required drugs to the patient during problem prioritization. As the goal of therapy is reduction of HCV infection in patient with ultimate cure. Hence Ribavarin, 800 mg OD (400mg, 2 caps. BD) was recommended by clinical pharmacist for this patient. Oral regimen of sofosbuvir along with ribavirin for 12 or 24 weeks predominantly treats hepatitis C [10]. Daclatasvir with sofosbuvir and ribavirin for almost 12 weeks resulted in a treatment of virologic response in patients either coinfected with HIV-1 and HCV genotypes 1 or only HCV [11]. Follow up for routine tests and examination was advised to patient for evaluation of treatment. Adherence to therapy was advised for prevention of hepatocellular carcinoma.

Conclusion

In this case scenario, the patient was suffering from hepatitis C. Recently, hepatitis is leading liver disorder so our aim was to treat severe abdominal pain of patient due to HCV and further progression of disease into hepatocellular carcinoma (HCC). She was treated with Antiviral therapy. Abdominal pain, generalized body weakness of patient was relieved after using sofosbuvir, daclatasvir and Ribavarin. Patient was stable after treatment with these antiviral agents.

References

- 1. Shepard CW, Finelli L, Alter MJ (2005) Global epidemiology of hepatitis C virus infection. The Lancet infectious diseases 5(9): 558-567.
- Moradpour D, Penin F, Rice CM (2007) Replication of hepatitis C virus. Nature reviews microbiology 5(6): 453-463.
- 3. Soriano V, Vispo E, Labarga P, Medrano J, Barreiro P (2010) Viral hepatitis and HIV co-infection. Antiviral Res 85(1): 303-315.
- 4. Weber R, Sabin CA, Friis Møller N, Reiss P, El Sadr WM, et al. (2006) Liver-related deaths in persons infected with the human immunodeficiency virus: the D:A:D study. Arch Intern Med 166(15): 1632-1641.

Virology & Immunology Journal

- 5. Hutin YJ, Chen RT (1999) Injection safety: a global challenge. Bull World Health Organ 77(10): 787-788.
- 6. The Global Burden of Hepatitis C Working Group (2004) Global burden of disease (GBD) for hepatitis C J Clin Pharmacol 44(1): 20-29.
- 7. Herrmann E, Lee JH, Marinos G, Modi M, Zeuzem S (2003) Effect of ribavirin on hepatitis C viral kinetics in patients treated with pegylated interferon. Hepatology 37(6): 1351-1358.
- 8. Lauer GM, Walker BD (2001) Hepatitis C virus infection. New England journal of medicine 345(1): 41-52.

- 9. Lai CL, Ratziu V, Yuen MF, Poynard T (2003) Viral hepatitis B. The Lancet 362(9401): 2089-2094.
- 10. Sulkowski MS, Naggie S, Lalezari J, Fessel WJ, Mounzer K, et al. (2014) Sofosbuvir and ribavirin for hepatitis C in patients with HIV coinfection. JAMA 312(4): 353-361.
- 11. Wyles DL, Ruane PJ, Sulkowski MS, Dieterich D, Luetkemeyer A, et al. (2015) Daclatasvir plus sofosbuvir for HCV in patients coinfected with HIV-1. The New England Journal of Medicine 373(8): 714-725.

