Management of Hepatitis C in the Indian Context: An Update

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Abstract

Hepatitis C, a leading cause of chronic liver disease globally, is caused by Hepatitis C Virus (HCV), a hepatotropic RNA virus. HCV infection starts with an acute infection, mostly subclinical, which ultimately leads to chronic hepatitis in about 80% of the infected cases.

HCV is classified into 6 major genotypes and numerous subtypes. The global prevalence of HCV infection is about 1.6% with a majority of these infections being in adults. There is widespread heterogeneity in the prevalence of different genotypes of HCV in different geographical regions of the world. While genotype 1 is the most common worldwide, different regions of the world report variations in the prevalence of the other genotypes. Genotype 3 is the commonest genotype in India, but there is a wide variation in the distribution of the other genotypes in different parts of the country; genotype 6, a comparatively rarer genotype, has been reported frequently from the northeast part of India.

With the introduction of new oral drugs, the Directly Acting Antivirals (DAA), the management protocols of hepatitis C has undergone dramatic transformation, with a paradigm shift towards an all-oral interferon-free regimen. However, developing countries like India still face a challenge with respect to accessibility and affordability of such newer regimens. Furthermore, the differences in genotypic distributions in India, with a higher prevalence of genotype 3, which is more difficult to treat, makes the situation even more challenging. As newer antivirals are being universally used to manage HCV infection, economically weaker countries like India should incorporate these changes in treatment guidelines soon. However, till substantial evidence on the efficacy of the newer regimens is enough to altogether discard the existing conventional modes of HCV therapy.

Keywords: Hepatitis C; HCV; Genotype; Treatment guidelines; Antivirals; Chronic hepatitis

Introduction

Hepatitis C is a blood-borne viral infection that is caused by Hepatitis C Virus (HCV), a hepatotropic RNA virus, with a propensity to affect the liver. It is recognised as a major public health problem worldwide responsible for chronic liver disease and a variety of extra-hepatic manifestations. It is primarily transmitted via the parenteral route which includes injection drug use, blood transfusion, unsafe injection practices, and other healthcare related procedures. HCV causes acute hepatitis which is mostly subclinical and gradually evolves into chronic hepatitis in about 80% of those infected [1]. For a long time hepatitis C remained obscure to researchers and one of the reasons is its clinically silent nature. Most patients with acute infection are symptom free and only a small proportion develops jaundice. Chronic HCV infection may be associated with vague, non-specific symptoms such as fatigue, joint pain, and discomfort in the right-upper quadrant of the abdomen. Patients usually become symptomatic when complications of chronic liver disease or extra-hepatic manifestations develop [2].

HCV is an enveloped positive stranded Ribonucleic Acid (RNA) virus belonging to genus Hepacivirus in the family Flaviviridae. It was first detected in 1989 using molecular biology techniques after extensive testing of serum from experimentally infected animals [3]. It is now widely recognized as one of the common aetiologic agents for cirrhosis of the liver and is the leading cause of liver transplantation and the most common chronic blood borne infection in developed countries like the USA [4]. HCV is classified into 6 broad genotypes and numerous subtypes, with wide genomic variations. This has led it to be an enigmatic challenge for both clinicians and researchers.

Global Epidemiology

The total global prevalence of HCV is estimated to be at an average of 1.6% (1.3-2.1%), corresponding to 115 (92-149) million viraemic infections. The majority of these infections, 104 (87-124) million, are among adults (defined as >15 years) with an anti-HCV infection rate of 2.0% (1.7-2.3%). The viraemic (RNA positive) prevalence is believed to be 1.1% (0.9-1.4%), which corresponds to 80 (64-103) million viraemic infections [5].

Globally, genotype 1 (G1) accounts for 46% of all HCV infections among adults making it the most common, followed by G3 (22%), G2 (13%), G4 (13%), G6 (2%), and G5 (1%) [5]. Undefined or combination of genotypes accounts for 3% of the total HCV infections. Among the subtypes, genotype 1b is the most common sub-type, accounting for 22% of all infections [5]. However there are significant variations in the global distribution of genotypes. Genotype 1 is the predominantly dominating genotype in Australasia, Europe, Latin America and North America (53-71% of all cases) [5]. Genotype 4 is the most common (71%) in North Africa and the Middle East [5]. In Asia the predominant genotype is G3 (39%) followed by G1 (36%) [5].

Epidemiology: The Indian Perspective

Currently India harbours an estimated 10–15 million chronic...
carriers of HCV, which is a major cause of liver related mortality and morbidity of the country [6]. The prevalence of Hepatitis C Virus (HCV) infection in the general population is estimated to be around 0.5%–1.5% [6]. However, the prevalence of HCV is variable in different high risk populations according to various studies and there is still a paucity of data from large multi-centric studies. Studies on voluntary or mixed donors have reported a prevalence of hepatitis C below 2% [4]. In a large community based systematic study from the Indian state of West Bengal the prevalence was documented to be around 0.71% [7]. On the contrary, a comparatively smaller study from the north eastern state of Arunachal Pradesh showed a much higher prevalence of hepatitis C of 7.89% [8]. A recent study done in Mizoram, another state in the north east of the country, showed a HCV prevalence of 71.2% among the active injection drug addicts [9]. There continues to be variations in reporting the HCV prevalence, depending upon the geographical region population sub-groups included in these studies.

Overall, genotype 3 is the predominant genotype (63.85%) followed by genotype 1 (25.72%) in India [6]. However there is a genotypic difference in different geographical regions of India, with genotype 3 being the commonest in northern, eastern and western India while genotype 1 is the commonest in the southern states of India (Figure 1). Besides the widespread prevalence of genotypes 3 and 1, there is also a trend of increased prevalence of genotypes 4 and 6 in certain regions of the country. Genotype 4 is found mostly in south Indian patients from the states of Andhra Pradesh and Tamil Nadu [6]. Genotype 6 is found to be prevalent exclusively in patients from north eastern parts of the India [6]. It is also to be noted that there is also increased prevalence of genotype 6 in various parts of the neighbouring country of Myanmar which shares its boundaries with the north-eastern states of India thus showing an ecological niche for HCV genotype 6 in north-eastern India and adjacent geographical areas [10]. Genotype 2 has rarely been reported from India whereas genotype 5 is yet to be reported5. Such epidemiological studies and trend analysis are important in the wake of licensure of the newer generation DAA which may revolutionize the management of HCV. However, it must be stressed that the reported distribution of the various genotypes can be expected to change with increasing migration of population and changes in high risk behaviour and lifestyle.

Virological Diagnosis of Hepatitis C

Virological diagnosis of HCV infection is based on two categories of laboratory tests: (i) indirect tests - serologic assays detecting specific antibody to HCV (anti-HCV) and (ii) direct tests - assays that can detect, quantify, or characterize the components of HCV viral particles, such as HCV RNA and core antigen. Anti-HCV is typically identified by using Enzyme-Linked Immunosorbent Assay (ELISA) or Chemiluminescence Assays (CIA) (Table 1). Newer generation assays have made the diagnosis by these tests highly specific. Confirmation of serological reactive tests may be done by a Nucleic Acid Test (NAT) for detection of HCV RNA. A third way for diagnosis is detection of the core antigen of HCV by ELISA. Evaluations in transfusion settings have shown that the HCV core Ag assay detects HCV infection as effective as NAT [11]. A simplified approach to the diagnosis of HCV is depicted in Figure 2.

Treatment Strategies of HCV

**Whom to treat**

According to the current guidelines antiviral treatment is recommended for all patients with chronic HCV infection, except those with limited life expectancy due to non-hepatic causes [12]. If resources limit the ability to treat all infected patients immediately as recommended, then it is most appropriate to treat those at greatest risk of disease complications before treating those with less advanced disease or persons with risk of HCV transmission or in whom treatment may reduce transmission (Tables 2 and 3).

**How to treat**

The selection of a treatment regimen will be determined by several factors. These include HCV genotype, treatment history, stage of fibrosis, medications for comorbid medical conditions, and adherence factors. The presence of associated co-morbid conditions like advancing age, chronic kidney disease, co-infection with HIV make the treatment protocols more complicated. Based on the joint recommendations issued by the American Association for the Study of Liver Diseases (AASLD) and the Infectious Diseases Society of America, in collaboration with the International Antiviral Society USA.

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**Figure 1:** Showing regional distribution of HCV genotypes in India.

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**Table 1:** Laboratory tests for virological diagnosis of HCV infection.

| Test Type       | Description                                           | Sensitivity (%) | Specificity (%) |
|-----------------|-------------------------------------------------------|----------------|----------------|
Antigenic targets for HCV diagnosis*.

Table 1: showing various ELISA techniques and their ability to detect different antigenic targets for HCV diagnosis*.

| Type of ELISA test (ELISA/CIA) | Antigenic targets for HCV diagnosis |
|-------------------------------|------------------------------------|
| ELISA 1st Generation          | Core - NS3 + NS4a - NS5b -         |
| ELISA 2nd Generation          | + + + -                           |
| ELISA 3rd Generation/ CIA     | + + + +                           |

*Modified from AASLD, IDSA, IAS-USA. Recommendations for testing, managing, and treating hepatitis C.

(IAS-USA2015), the brief summary of recent treatment guidelines for HCV is as follows [12].

**Genotype 1:** New interferon-free regimens comprising of Direct Acting Antiviral (DAA) are now the standard of care for the treatment of chronic hepatitis C genotype 1 infection. For initial therapy of treatment-naive patients with genotype 1 infection, four regimens with similar efficacy are recommended [12-15] (a) ledipasvir-sofosbuvir plus ribavirin, (b) ombitasvir-paritaprevir-ritonavir with or without ribavirin, (c) sofosbuvir plus simeprevir with or without ribavirin, or (d) daclatasvir plus sofosbuvir with or without ribavirin.

For retreatment of patients with genotype 1 who had no resolution despite prior therapy with peg-interferon and ribavirin, one of following regimens that have similar efficacy are recommended [12]: (a) ledipasvir-sofosbuvir with or without ribavirin, (b) ombitasvir-paritaprevir-ritonavir and dasabuvir with or without ribavirin, (c) sofosbuvir plus simeprevir with or without ribavirin, or (d) daclatasvir plus sofosbuvir with or without ribavirin. The use of ribavirin and the duration of therapy depend on the HCV genotype 1 subtype and on cirrhosis status. The duration of therapy with ledipasvir-sofosbuvir is 12 weeks without cirrhosis; those with cirrhosis can receive ledipasvir-sofosbuvir for 24 weeks or ledipasvir-sofosbuvir plus ribavirin for 12 weeks.

In patients with genotype 1 infection who previously failed sofosbuvir plus ribavirin, with or without peginterferon, the recommended regimen is ledipasvir-sofosbuvir plus ribavirin [12]. The duration of therapy is 12 weeks without cirrhosis and 24 weeks with cirrhosis. For treatment-naive and treatment-experienced patients with genotype 1 infection, the new benchmark for sustained virologic response rates is 90% or greater. The major barrier to treatment with all new therapies is the extremely high cost of a treatment course.

**Genotype 2:** The standard of care for initial treatment of HCV genotype 2 consists of sofosbuvir plus ribavirin, which typically achieves an SVR rate of 90% or better; the duration of therapy, is 12 weeks in patients without cirrhosis and 16 weeks in those with cirrhosis [12]. For initial treatment of patients with HCV genotype 2 who are intolerant to ribavirin, the 12-week regimen of daclatasvir plus sofosbuvir provides a new ribavirin-free option [16,17]. The recommended retreatment of patients with genotype 2 and prior treatment failure with peginterferon and ribavirin consists of sofosbuvir plus ribavirin for 16 or 24 weeks; the optimal duration remains unclear [12]. For retreatment of patients with genotype 2 who previously failed therapy with sofosbuvir plus ribavirin, the recommended regimen is either (a) daclatasvir plus sofosbuvir (with or without ribavirin) for 24 weeks or (b) sofosbuvir plus ribavirin plus peginterferon for 12 weeks [12]. The demand and interest for studies that examine future therapies for genotype 2 are somewhat limited given the very high SVR rates with the all-oral regimen of sofosbuvir plus ribavirin.

**Genotype 3:** With new direct-acting antiviral treatments, genotype 3 has emerged as the most difficult genotype to treat. The recent FDA approval of daclatasvir has significantly expanded and improved treatment options for patients with genotype 3 infection.

Three regimens are recommended for genotype 3 treatment-naive patients [12,18]: (1) daclatasvir plus sofosbuvir for 12 weeks (in patients without cirrhosis), (2) daclatasvir plus sofosbuvir, with or without ribavirin, for 24 weeks (in patients with cirrhosis), or (3) sofosbuvir plus ribavirin plus peginterferon for 12 weeks (with or without cirrhosis).

The recommended regimens for genotype 3 treatment-experienced patients who have failed prior treatment with peginterferon and ribavirin are [12]: (1) daclatasvir plus sofosbuvir for 12 weeks in patients without cirrhosis and 24 weeks along with ribavirin in those with cirrhosis, or (2) sofosbuvir plus ribavirin plus peginterferon for 12 weeks (with or without cirrhosis).

The recommended regimen for genotype 3 treatment-experienced patients who have failed prior treatment with sofosbuvir consists of either [12] (1) daclatasvir plus sofosbuvir plus ribavirin for 24 weeks, or
Furthermore, there are certain genetic variations, especially the one to treat genotype 5 or 6 infection [12].

A combination of grazoprevir-elbasvir is a promising investigational agent plus ribavirin plus peginterferon for 12 weeks. The fixed dose consists of [12] (a) ledipasvir-sofosbuvir for 12 weeks or (b) sofosbuvir genotype 5 or 6 who previously failed peginterferon plus ribavirin [20, 21].

Retreatment are based on in vitro data and limited experience in clinical cost of a treatment course.

The barrier to treatment of patients with genotype 4 infection is the high medications grazoprevir and elbasvir have promising activity against sofosbuvir plus ribavirin plus peginterferon [12]. The investigational therapy, except they also include the option to use a 12-week course of (b) ombitasvir-paritaprevir-ritonavir plus ribavirin for 12 weeks, or recommended [12,19] in the (a) ledipasvir-sofosbuvir for 12 weeks, (b) sofosbuvir plus ribavirin for 24 weeks. For retreatment of patients with genotype 4 who previously failed therapy with peginterferon (c) sofosbuvir plus ribavirin for 24 weeks. For retreatment of patients with genotype 4 infection, three regimens with similar efficacy are recommended [25]. It is with this background that one must consider the consensus statement of 2014 recommended by the HCV task force of the Indian National Association for Study of the Liver on HCV infection in India [25].

These guidelines recommend standard of care therapy for chronic hepatitis C patients with presently available drugs while awaiting the entry of newer DAA into the Indian market. At this point of time, when the entire Western scientific community is highlighting the positive effect of the DAA, with the AASLD-2015 guidelines considering Peg-IFNa/RBV therapy as “not recommended”, local compulsions in resource limited countries like India would probably still require the continued use of standard treatment regimes and would even consider use of conventional IFN. Where cost is a limiting factor in developing countries like India, relatively less efficacious but cheaper modes of therapies like INF may still be considered as an alternative option, as long as newer DAAs are unaffordable to the majority of the patients. As an example, in the neighbouring country of Pakistan, more than 20,000 persons have received therapy with conventional IFN, which has been funded by the state [28].

Another major point of discordance in the Indian guidelines, as compared to international guidelines, has been the recommendation to consider deferring treatment for certain groups of patients like those with (i) low levels of fibrosis, (ii) those with failed interferon therapy, (iii) are interferon-intolerant and (iv) stable patients after liver and kidney and liver transplant. For such patient groups it will probably be better to wait until such safer therapies become freely available and affordable in India.

However with the arrival of Sofosbuvir in early 2015 the Indian guidelines are also expected to undergo a dramatic change. Although efforts are on to bring these drugs within the reach of people at an affordable cost, it is not clear as to how much time it will take. Time and more intense studies demonstrating the efficacy of the DAAs in
the Indian patients is needed before a national consensus guideline incorporating regimens at par with international guidelines can be implemented.

Conclusion

Hepatitis C continues to be a major global threat leading to development of chronic hepatitis, cirrhosis and hepatocellular carcinoma. The distribution of HCV has wide geographic variations with multiple genotypes and subtypes. The pattern of distribution in India too is diverse with a distinct geographic heterogeneity in distributions of the different genotypes. Furthermore, an enigmatic presence of HCV genotype 6 in a certain ecologic niche in the north-eastern part of the country makes the management more challenging. As the treatment undergoes dramatic changes with newer drugs revolutionizing the management protocols, India too has to incorporate these newer changes. Nevertheless, keeping in mind the accessibility and affordability of the newer regimens in a lower-middle income country like India, it is perhaps not yet time to completely abandon the conventional modes of therapy which are currently being used.

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