Treatment of HCV in Persons with Substance Use

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Module 6: Treatment of Key Populations and Unique Situations
Lesson 3: Treatment of HCV in Persons with Substance Use

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Background

Substance use disorders are common in the United States, with 2022 data indicating that nearly 1 in 4 (24.9%) of persons 12 years of age or older used an illicit drug in the past year (Figure 1).[1] The availability of highly effective direct-acting antiviral (DAA) medications has radically changed the assessment and consideration of substance use in hepatitis C virus (HCV) treatment decisions. The AASLD-IDSA HCV Guidance states that recent or active injection drug use or alcohol use is not a contraindication to HCV treatment and requirements for pretreatment screening for illicit drug or alcohol use should be discontinued.[2] Nevertheless, for persons with chronic HCV infection, substance use, either past or present, which encompasses the use of opioids, amphetamines, cannabis, cocaine, alcohol, and other drugs, may still be relevant to the individual's overall health and medication access. The following discussion will address the potential impact of substance use on HCV disease and HCV treatment and provide current recommendations for the treatment of HCV in persons with a substance use disorder.

Approach to HCV Treatment in Persons with Substance Use

HCV Treatment Eligibility

The AASLD-IDSA HCV Guidance recommends treatment for all persons with chronic HCV, except those with a short life expectancy that HCV treatment cannot remediate, liver transplantation, or another directed therapy.[2,3] The general approach to considering initiation of HCV treatment for individuals with a prior history of substance use, including injection drug use, should be the same as in persons with no history of drug use. Persons with substance use disorder, including those with injection drug use or alcohol use should have the same HCV pretreatment screening requirements as those without a substance use disorder.[2]

Abstinence Requirements

Although some payers require 6 months or more of abstinence prior to HCV treatment, studies of both injection drug use and alcohol use have found no impact of duration of abstinence on the likelihood of achieving a sustained virologic response 12 weeks (SVR12) after completing DAA-based therapy.[4] Thus, there is no medical reason to ensure abstinence (for any duration) prior to HCV treatment. Current AASLD-IDSA HCV Guidance recommendations state that current or prior substance use is not a contraindication to HCV treatment.[2]

Impact of Comorbidities in Persons with Substance Use Disorder

In clinical practice, treating persons with an active substance use disorder may be complicated by coexisting social stressors, competing survival needs, and barriers erected by payers. However, clinical experience suggests that, with appropriate infrastructure and patient support, including treatment of substance use disorders, HCV treatment is very feasible in this population.[5,6,7,8,9] Examples of patient support include directly-observed therapy and related approaches, patient navigation, and group treatment models, particularly in substance use disorder treatment settings.[10]

HCV Treatment in Persons with Opioid Use

Impact of Opioid Use on Natural History of HCV

Through the sharing of syringes and other injection equipment, injection opioid use is a major driver of HCV transmission, but opioid use itself, either orally or by injection, does not appear to speed the progression of liver disease in persons with chronic HCV.[11] Opioid analgesic use disorder is also a risk factor for HCV acquisition and transmission, as some individuals transition from oral ingestion of prescribed opioids to use of illicit opioids, which can include injection opioid use.[12,13] Nevertheless, opioid use itself, either orally or by injection, does not appear to impact the natural history of liver disease in persons with chronic HCV.[11]

Impact of Treating People with Active Injection Drug Use on HCV Transmission

Mathematical modeling, even assuming a reinfection rate equal to initial infection rates, has demonstrated that HCV treatment among persons with active injection drug use would result in a significant reduction in HCV transmission.[14,15,16,17] Several studies utilizing mathematical modeling based on DAA regimens concluded that scaling up HCV treatment in people who inject drugs (PWID) would have a major impact in reducing HCV incidence and prevalence in this patient population, even more so in the setting of robust access to sterile injection equipment and medications for opioid use disorder (e.g., buprenorphine-naloxone or methadone).[16] Further, scaling up and widespread treatment of HCV in PWID as a prevention tool, akin to treating HIV to reduce community viral load, has become a more realistic goal with the short-course, well-tolerated oral regimens. Indeed, several efforts focusing on HCV micro-elimination (i.e., elimination of HCV as a public health threat within a defined population) have provided real-world evidence for HCV treatment as prevention, demonstrating significant reductions in HCV incidence resulting from widespread treatment.[18,19,20]

Impact of Opioid Use and HCV Treatment Adherence

Multiple studies that have enrolled persons with active or recent injection drug use have shown excellent adherence with DAA-based HCV therapy (Figure 2).[10,21,22,23,24,25] In particular, persons with opioid use disorder who receive opioid agonist maintenance therapy (e.g., methadone, buprenorphine, or buprenorphine-naloxone) during HCV treatment have excellent rates of adherence, treatment completion, and sustained virologic response (SVR) rates, all comparable to results of other study participants.[26,27]

HCV Treatment Outcomes with DAA Therapy in PWID

The following summarizes several key studies that have analyzed HCV treatment responses with DAA-based therapy in persons who inject drugs or who have previously injected drugs and were receiving opioid agonist therapy. Multiple studies clearly show that DAA-based therapy in persons with past or current injection drug use results in high SVR rates, comparable to those seen in persons who do not use drugs.[21,24]

- **Elbasvir-Grazoprevir (C-EDGE CO-STAR)**: In this phase 3 multinational study (C-EDGE CO-STAR), investigators evaluated HCV treatment with elbasvir-grazoprevir in 301 PWID who had also been receiving opioid agonist therapy (e.g. methadone, buprenorphine, or buprenorphine-naloxone maintenance) for at least 3 months prior to enrollment.[23] Participants with chronic HCV genotypes 1, 4, or 6 were randomized to immediately receive a 12-week course of elbasvir-grazoprevir (immediate treatment group) or to receive the 12-week course of elbasvir-grazoprevir after a 16-week delay (deferred treatment group). On day 1 of the study, 58% of the subjects had a positive urine drug screen (excluding opiate agonist therapy). Overall, when excluding participants who discontinued for non-treatment reasons, 92% in the immediate treatment group and 90% in the deferred treatment group achieved an SVR12; the treatment responses were excellent regardless of cirrhosis status and baseline drug screening results (Figure 3).[23]
- **Glecaprevir-Pibrentasvir**: In a pooled analysis of 7 phase 3 studies that involved HCV treatment

with 8 or 12 weeks of glecaprevir-pibrentasvir, investigators compared SVR12 rates among persons with recent drug use (use in the past 12 months), former drug use (use more than 12 months ago), and no drug use.[28] The SVR12 rates were high among all groups: 93% in persons with recent drug use, 97% in those with former drug use, and greater than 99% in persons who did not use drugs.[28] A similar pooled analysis of 8 phase 2 and 3 trials of glecaprevir-pibrentasvir evaluated treatment outcomes among persons receiving opioid substitution therapy.[29] In the intention-to-treat analysis, SVR12 rates were 96% among persons receiving opioid substitution therapy and 98% in those not receiving opioid substitution therapy.[29] In an analysis of 2 prospective cohort studies in Spain, in the intention-to-treat analysis, the SVR12 rates were 97% among persons who never used drugs, 96% among persons with past drug use, and 85% among persons with recent or active drug use.[30] This slightly lower SVR12 rate among persons with recent or active drug use was felt to be secondary to voluntary discontinuation, as no difference in SVR12 rates was seen when comparing persons who never used drugs (99%), persons with past drug use (100%), and persons with recent or active drug use (100%) in the per-protocol analysis.[30]

- **Ledipasvir-Sofosbuvir (ION Trials)**: In the ION trials, participants with chronic HCV genotype 1 infection received 8, 12, or 24 weeks of ledipasvir-sofosbuvir.[31] Investigators performed a pooled data analysis to compare HCV treatment response in participants receiving opioid substitution therapy during treatment (n = 70) with participants not receiving opioid substitution therapy (n = 1882).[31] The two groups had similar treatment completion, adherence, and SVR12 rates (Figure 4).[31] Among those receiving opioid substitution therapy, 94% (66 of 71) achieved an SVR12.[31] In addition, a pilot trial of ledipasvir-sofosbuvir among 31 persons with active injection drug use randomized participants 1:1 to modified directly-observed treatment (mDOT) or unobserved dosing. All but one participant (in the mDOT arm) completed treatment; 97% achieved an end-of-treatment response and 90% achieved SVR12.[32]
- **Sofosbuvir-Based Treatment in Phase 3 Trials**: In a pooled analysis of data from phase 3 trials using sofosbuvir-based regimens, investigators compared HCV treatment responses in participants receiving opioid substitution therapy during treatment (n = 194) with participants not receiving opioid substitution therapy (n = 4,549).[27] There were no significant differences in SVR12 rates (94% in those receiving opioid substitution therapy versus 97% for those not receiving opioid substitution therapy).[27]
- **Sofosbuvir-Velpatasvir (SIMPLIFY)**: In this single-arm, open-label, phase 4 study, 103 participants with chronic HCV genotype 1, 2, 3, 4, 5, or 6 infection and recent (within 6 months) injection drug use were treated with a 12-week course of sofosbuvir-velpatasvir.[22] During the treatment course, 59% (61 of 103) participants were receiving opioid substitution therapy, and 74% (76 of 103) had used injection drugs in the month prior to starting treatment.[22] A total of 97% (100 of 103) participants completed treatment; 2 were lost to follow-up, and one person died from an accidental overdose. Overall, 97 (94%) of participants achieved an SVR12.[22]
- **Sofosbuvir-Velpatasvir (ASTRAL Trials)**: In a subset analysis of the phase 3 ASTRAL trials, investigators compared treatment response to sofosbuvir-velpatasvir in persons receiving opioid substitution therapy during treatment (n = 51) compared with those who were not receiving opioid substitution therapy (n = 984).[26] Receipt of opioid substitution therapy did not impact treatment responses; 96% of participants receiving opioid substitution therapy achieved an SVR12.[26]

Pretreatment Requirements

The AASLD-IDSA HCV Guidance does not recommend requiring abstinence from opioids prior to HCV treatment.[2] Indeed, active injection drug use in an individual with chronic HCV is considered by many to be a direct indication for HCV treatment, due to the potential benefit of reducing secondary HCV transmission.[33,34] In contrast to these expert recommendations, some payers may require abstinence from non-prescribed opioids and other recreational substances, but this type of requirement is not evidence-based and not consistent with recommendations in the AASLD-IDSA HCV Guidance.[3] There are three FDA-approved treatments for opioid use disorder—buprenorphine, methadone, and extended-release naltrexone. Although methadone access is limited to designated outpatient treatment programs, the other medications can be prescribed in the context of routine medical care.

Management Strategies

For persons with a past or current history of opioid use disorder, treatment of HCV infection would ideally be performed in a multidisciplinary setting whereby treatment for HCV and opioid use disorder can be jointly addressed.[2] Multiple treatment options exist for opioid use. Agonist maintenance therapy is the most effective known treatment and has been shown to reduce the risk of new HCV infection.[35] Data on the use of DAAs and medications for opioid use disorder (MOUD) indicate comparable SVR rates when comparing persons on MOUD with those who do not use drugs.[22,23,26,27,29,31] Injectable extended-release naltrexone is also approved for opioid dependence, although access can be challenging and uptake can be limited. There are no known clinically significant interactions between opioid agonist therapies or naltrexone and currently approved DAA medications.[36,37] A detailed discussion of opioid agonist therapy is beyond the scope of this topic review. In addition, several studies have evaluated optimal models for HCV treatment within harm reduction and opioid treatment programs.[10,38,39] One large study (HERO study) conducted in eight opioid treatment programs and 15 community health centers randomized 755 individuals to sofosbuvir-velpatasvir plus patient navigation or sofosbuvir-velpatasvir plus modified directly observed therapy (mDOT).[10] Although there was slightly higher adherence to sofosbuvir-velpatasvir in the mDOT arm, there was no difference in SVR12 rates, with 60% and 91% of patients in the mDOT arm and 62% and 93% of patients in the patient navigation arm achieving an SVR12 in the intention-to-treat analysis and per-protocol analysis, respectively.[10]

Potential Reinfection with HCV among Persons who Inject Drugs

Multiple studies have shown a significant risk of HCV reinfection in persons cured with HCV therapy. Thus, it is essential to counsel persons with past or active injection drug use be counseled that they can become reinfected with HCV after achieving an SVR. This risk is significant in persons who inject drugs, but reinfection can also occur through sexual contact, particularly among men who have sex with men (MSM).[40] In one study that clearly evaluated reinfection among treated PWID, the reinfection rate for those reporting ongoing injection after SVR was 5.3/100 person-years.[41] Similarly, in a recent systematic review of 36 studies of HCV reinfection following successful HCV treatment in persons who inject drugs, the overall rate of HCV reinfection was 6.2 per 100 person-years among those who reported recent injection drug use.[42] A similar systematic review and meta-analysis of 41 single-armed observational studies found a pooled rate of reinfection of 4.1 per 100 person-years), with a rate of reinfection of 2.8 per 100 person-years among PWID, 7.4 per 100 person-years among MSM, and 7.2 per 100 person-years among incarcerated individuals.[43] Thus, providing access to counseling for safe injection practices and MOUD is an important component of HCV treatment programs. Detailed guidance on safer injection techniques, such as ensuring a source of sterile syringes and other injection equipment and reviewing possible sources of HCV transmission, such as cottons, cookers, water, alcohol pads, or any syringes used to divide, prepare, or inject drugs, may lessen the risk of reinfection.[44]

HCV Treatment in Persons with Stimulant Use

Impact of Stimulant Use on Natural History of HCV

Injection of cocaine or methamphetamine is another major driver of HCV transmission.[45,46,47] Other routes of administration of stimulants, such as intranasal, may also be associated with HCV transmission.[48] In addition, prolonged stimulant use may result in cardiac and cerebrovascular toxicity.

Pretreatment Requirements

There is no medical requirement for abstinence from stimulant use prior to HCV treatment. Although some payers may require abstinence from methamphetamine prior to starting treatment, any such requirement is not consistent with recommendations in the AASLD-IDSA HCV Guidance.[3]

HCV Treatment Outcomes in Persons with Stimulant Use

Although there are limited data specific to the impact of stimulant use on outcomes of DAA therapy, several large trials evaluating treatment outcomes among persons with opioid use disorder have included sizable proportions of persons with concurrent stimulant use.[10,22] For example, in the HERO study, which was a randomized trial of sofosbuvir-velpatasvir plus modified directly observed therapy versus sofosbuvir-velpatasvir plus patient navigation in 8 opioid treatment programs and 15 community health centers, 32% of individuals with available urine drug screen results at baseline had a test positive for methamphetamine, and 42% were positive for cocaine,[10] In this study, SVR12 rates in the per-protocol analysis mirrored those of persons without substance use or stimulant use, with greater than 90% of participants with or without substance use achieving an SVR12.[10] Similarly, in the SIMPLIFY study, an open-label, single-arm trial of sofosbuvir-velpatasvir for persons with recent injection drug use, 30% reported injection methamphetamine use, and 13% reported injection cocaine use in the past 30 days.[22] In this study, 94% of individuals achieved an SVR12.[22]

Management Strategies

Stimulant use is often more intermittent than opioid or alcohol use but can also be associated with periods of poor adherence to medical care. Pharmacologic options are limited, with multiple current trials underway for both methamphetamine and cocaine dependence.[49] In a double-blind, placebo-controlled, randomized trial conducted from 2012 to 2015, a 12-week course of extended-release naltrexone did not appear to reduce amphetamine use among dependent persons.[] In a similar randomized, controlled trial, a 12-week course of aripiprazole, when given over 12 weeks, did not significantly reduce methamphetamine use.[51] Mirtazapine has demonstrated efficacy in reducing methamphetamine use in two separate trials.[52] Bupropion and modafinil have also demonstrated benefits in small trials.[53,54] A combination of high-dose bupropion plus high-frequency extended-release naltrexone demonstrated benefit in a trial of 403 participants.[55]

HCV Treatment in Persons who Consume Alcohol

Impact of Alcohol Use on Liver Fibrosis in Persons with Chronic HCV

Several studies have shown that heavy alcohol consumption (at least 60 grams/day in men and 40 grams/day in women) accelerates the progression of HCV-related hepatic fibrosis (Figure 5).[56,57] A typical alcoholic drink (12 ounces of beer, 5 ounces of wine, and 1.5 ounces of whiskey) contains 12 grams of alcohol. An estimated one-third of patients with chronic HCV infection have cirrhosis attributable to heavy alcohol consumption.[58] In a study in Alaska, investigators compared outcomes in persons who recovered from HCV with those who had chronic HCV and found heavy alcohol use (at least 50 grams of alcohol daily) was associated with the highest incidence of end-stage liver disease, regardless of whether the individual had recovered from HCV or had chronic HCV infection.[59] In addition, separate studies have shown that liver disease progression may continue among persons who heavily use alcohol even if SVR is achieved with HCV treatment. Taken together, the available data suggest reducing alcohol use is critical to liver health. The effects of low or moderate alcohol consumption on liver health are not well characterized for persons with chronic HCV infection.

HCV Treatment Outcomes Among Persons with Alcohol Use

Most of the studies that have addressed whether alcohol use impacts treatment outcomes were performed in the pre-DAA treatment era and results from these studies were mixed.[9,60,61] In the DAA treatment era, a large observational study out of the Veteran's Affairs (VA) healthcare system evaluated the impact of alcohol use on HCV DAA-based treatment outcomes.[62] Of the 15,151 persons who initiated DAA therapy and had a documented AUDIT-C score, 68.5% were categorized as abstinent, 22.6% as low-level drinking, and 8.9% as unhealthy drinking. Overall SVR12 rates were high among all persons in the study, regardless of alcohol use, with no statistical difference between HCV genotype or by cirrhosis status (Figure 7).[62] These findings support current recommendations to not exclude persons from HCV treatment based on their alcohol use.

Pretreatment Requirements

Although abstinence from alcohol is strongly encouraged for patients with chronic HCV infection, the AASLD-IDSA HCV Guidance recommends that abstinence from alcohol (or a reduction in alcohol consumption) should not be a requirement prior to HCV DAA treatment.[2] Although some payers may still require abstinence from alcohol prior to HCV treatment, this requirement is not based on guidelines or data.

Management Strategies

Although abstinence from alcohol prior to HCV DAA treatment is no longer required, alcohol consumption is discouraged in patients with chronic HCV infection due to the hepatotoxic effects of alcohol and its acceleration in liver fibrosis. Multiple pharmacologic agents are available for alcohol use disorder, including naltrexone, acamprosate, and topiramate.[63] Among these, the most promising results have been seen with naltrexone, particularly when given as a monthly injection. Brief counseling on alcohol has also shown reductions in use among persons with HCV infection. A multidisciplinary approach, involving personalized addiction care and case management, may provide further benefit in managing alcohol dependence.[9] Following DAA-based therapy, one study that included 123 participants found provider-delivered, alcohol-related counseling during HCV treatment was successful in reducing alcohol consumption patterns both during and after treatment in individuals with harmful alcohol use.[64]

HCV Treatment in Persons who Use Cannabis

There is mixed evidence regarding cannabis use and HCV-related hepatic fibrosis progression.[65,66,67] Two separate longitudinal cohort studies found no association between cannabis use and progression of liver fibrosis among patients coinfected with HCV and HIV.[65,68] In addition, one study found a positive association between cannabis use and good adherence with HCV treatment.[69] Although individuals living with HCV are typically advised to abstain from regular cannabis use, ongoing cannabis use is not considered a contraindication for initiating HCV therapy.

Summary Points

- Active or past substance use, substance use disorder, or injection drug use is not a contraindication to HCV treatment.

- Treatment of HCV in persons with active injection drug use likely has major public health benefits in terms of reducing secondary HCV transmission.

- It is important to talk to patients about their substance use to best understand how to support them through treatment and lower the risk of reinfection.
- Persons with HCV should be aware that heavy use of alcohol may continue to cause damage to the liver.
- Therapeutic approaches to substance use disorders are generally more effective when a pharmacologic agent is included.
- Care should be taken to ensure that PWID are aware of specific drug use techniques to avoid reinfection, particularly in the ways drugs are divided or prepared for injection.

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Figures

Figure 1 Number of Persons Aged 12 or Older with Past Year Illicit Drug Use, United States, 2022

Source: Substance Abuse and Mental Health Services Administration (SAMHSA). (2022). Key substance use and mental health indicators in the United States: Results from the 2022 National Survey on Drug Use and Health (HHS Publication No. PEP23-07-01-006, NSDUH Series H-58). Rockville, MD: Center for Behavioral Health Statistics and Quality, Substance Abuse and Mental Health Services Administration.

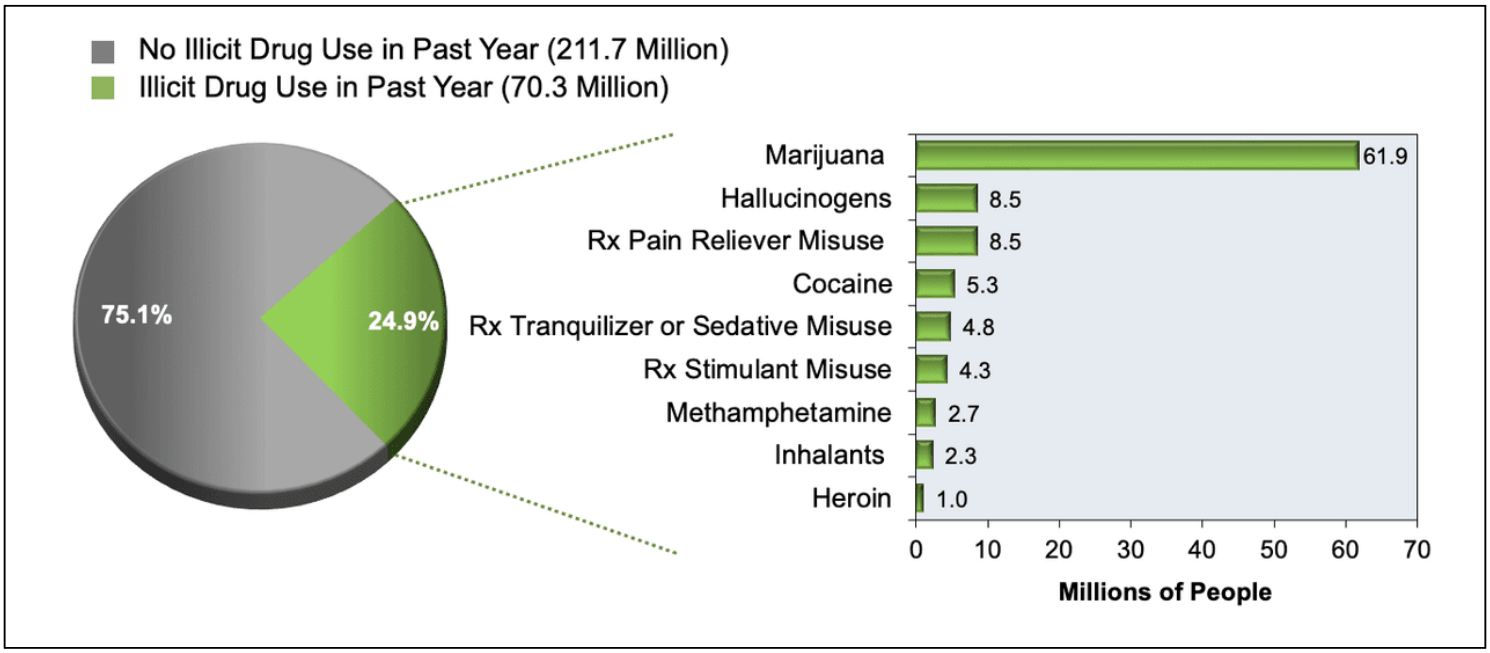

Figure 2 Adherence with HCV Therapy in C-EDGE CO-STAR Trial*

HCV treatment with elbasvir-grazoprevir in 301 PWID who were receiving opioid agonist therapy for at least 3 months prior to enrollment.

Source: Dore GJ, Altice F, Litwin AH, et al. Elbasvir-Grazoprevir to Treat Hepatitis C Virus Infection in Persons Receiving Opioid Agonist Therapy: A Randomized Trial. Ann Intern Med. 2016;165:625-34.

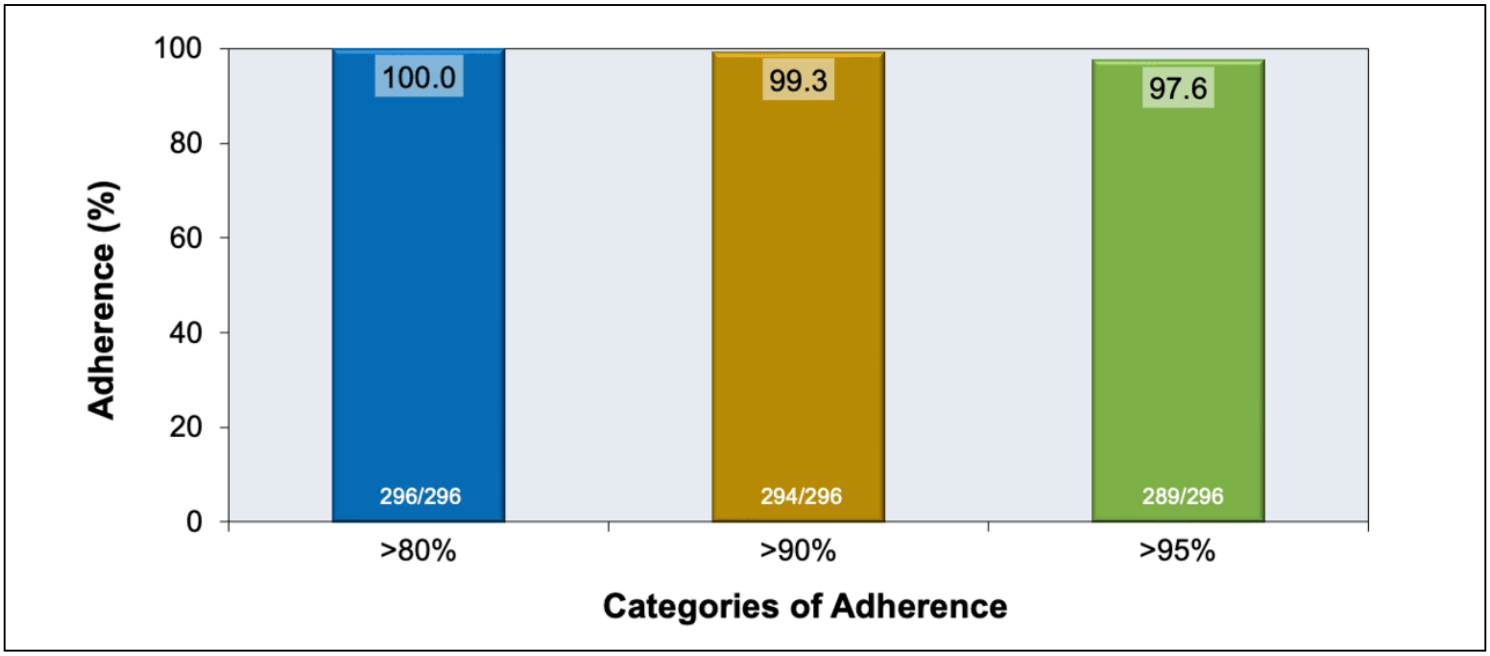

Figure 3 (Image Series) - Elbasvir-Grazoprevir in PWID: C-EDGE CO-STAR (Image Series) - Figure 3 (Image Series) - Elbasvir-Grazoprevir in PWID: C-EDGE CO-STAR
Image 3A: SVR12, by HCV Genotype (Immediate-Treatment Group)

In this analysis, reinfections are considered as failures.

Source: Dore GJ, Altice F, Litwin AH, et al. Elbasvir-Grazoprevir to Treat Hepatitis C Virus Infection in Persons Receiving Opioid Agonist Therapy: A Randomized Trial. Ann Intern Med. 2016;165:625-634.

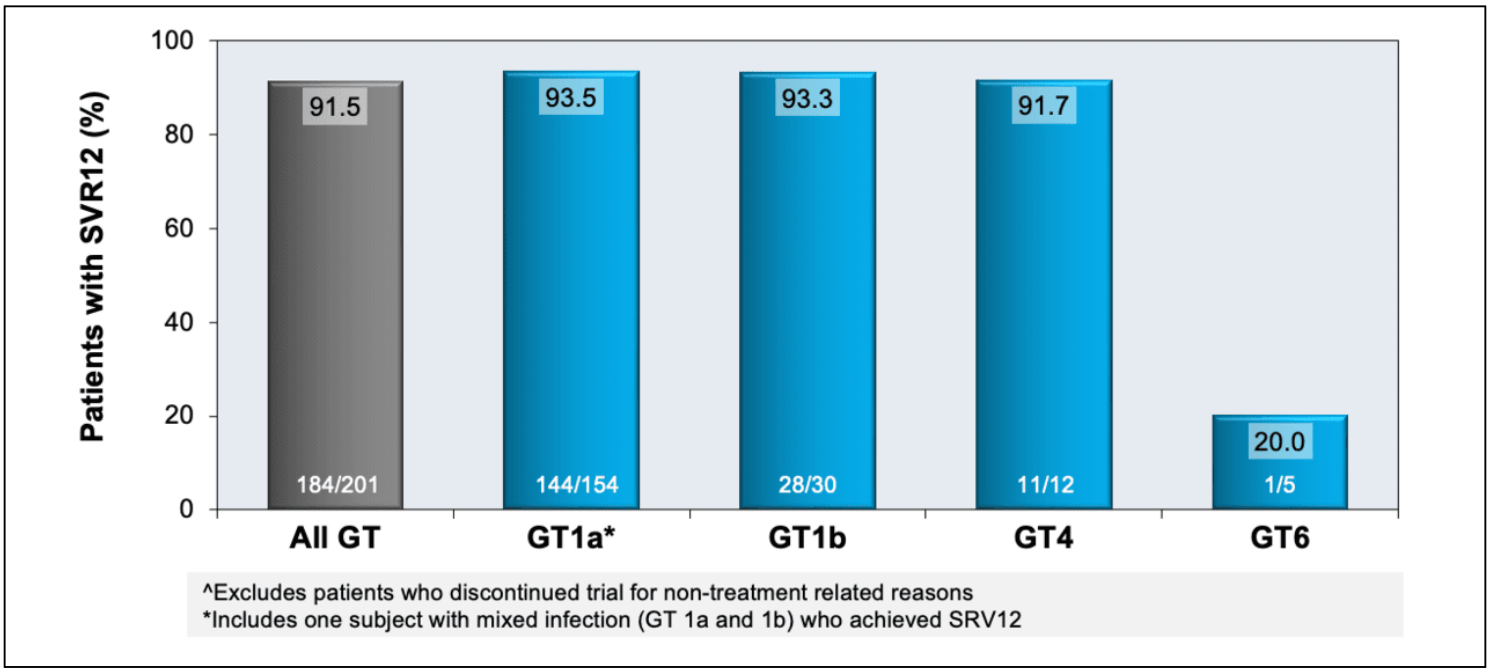

Figure 3 (Image Series) - Elbasvir-Grazoprevir in PWID: C-EDGE CO-STAR
Image 3B: SVR12, by HCV Genotype (Modified Full Analysis Set^)

In this analysis, reinfections are considered as responders.

Source: Dore GJ, Altice F, Litwin AH, et al. Elbasvir-Grazoprevir to Treat Hepatitis C Virus Infection in Persons Receiving Opioid Agonist Therapy: A Randomized Trial. Ann Intern Med. 2016;165:625-634.

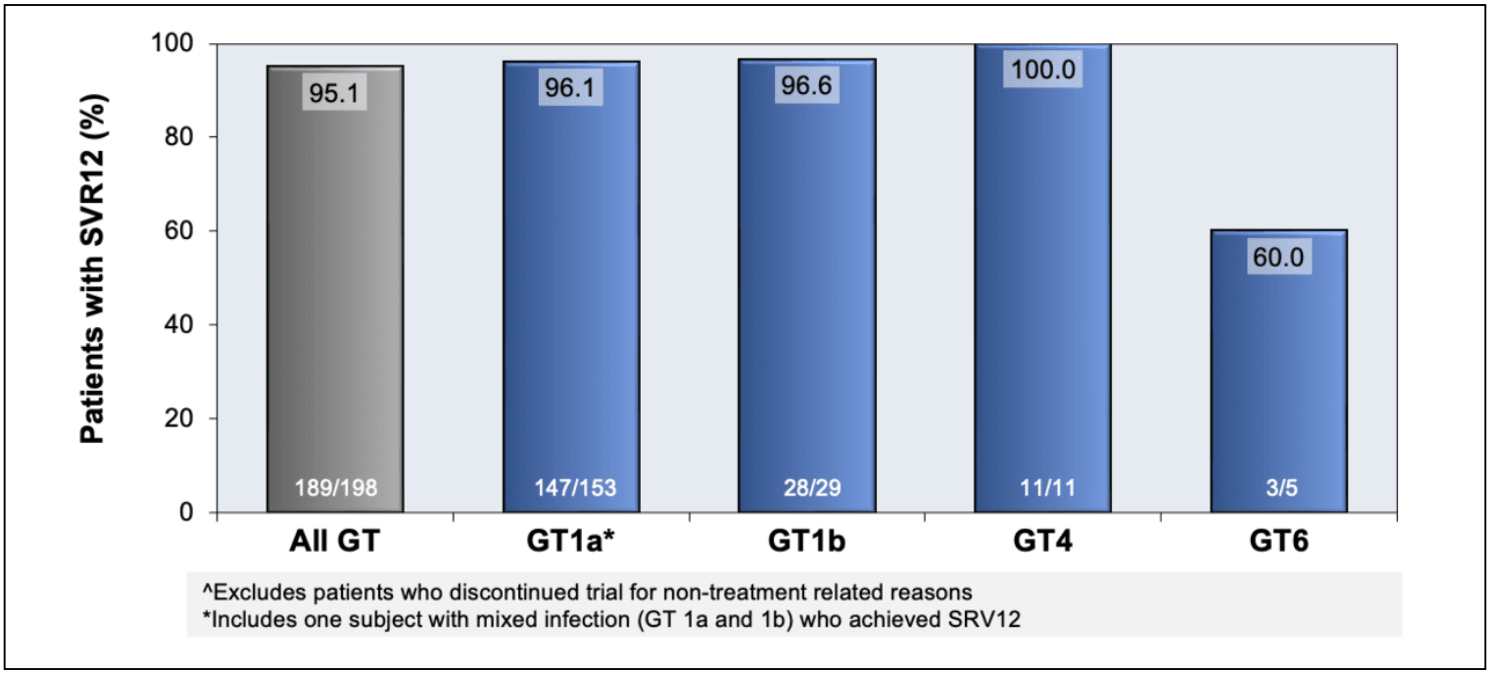

Figure 3 (Image Series) - Elbasvir-Grazoprevir in PWID: C-EDGE CO-STAR
Image 3C: SVR12 by Subgroups Analysis

Dore GJ, Altice F, Litwin AH, et al. Elbasvir-Grazoprevir to Treat Source: Hepatitis C Virus Infection in Persons Receiving Opioid Agonist Therapy: A Randomized Trial. Ann Intern Med. 2016;165:625-634.

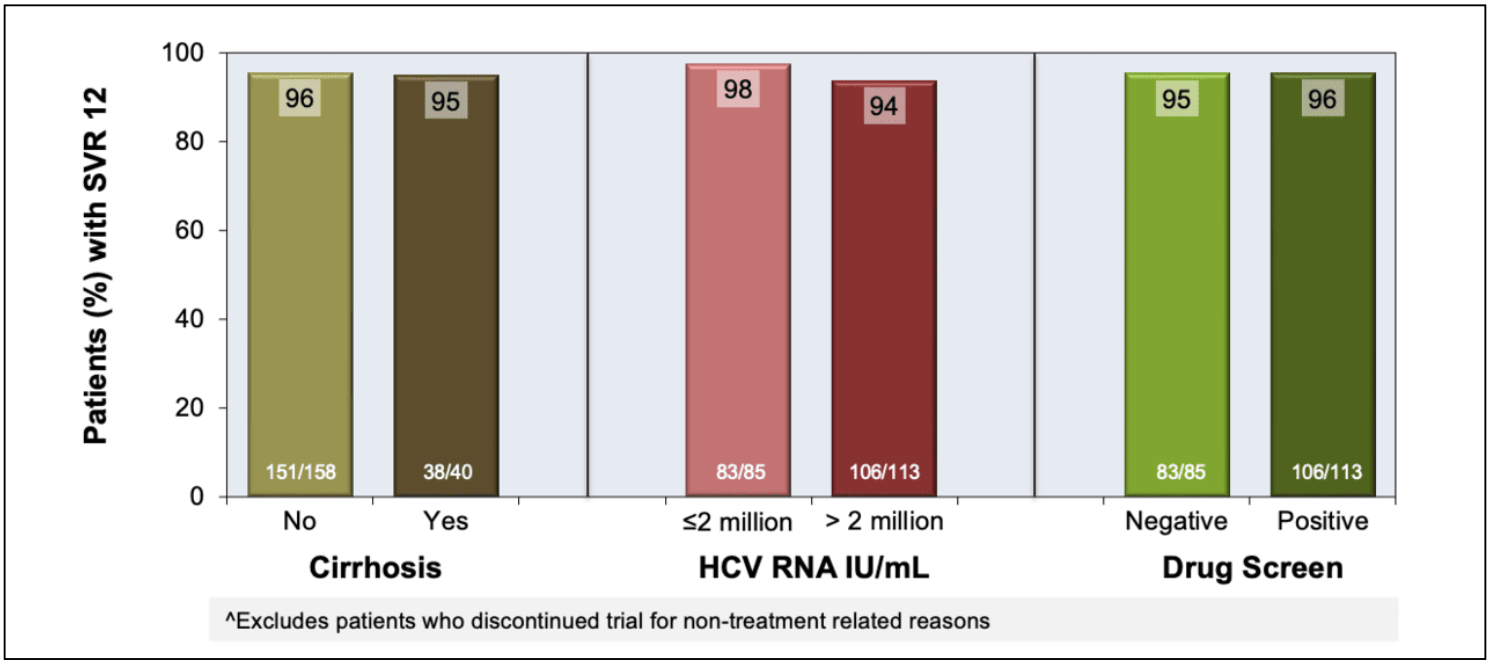

Figure 4 SVR12 Response to Ledipasvir-Sofosbuvir in Persons Receiving Opioid Substitution Therapy: Phase 3 ION Trials

Abbreviations: OST = opioid substitution therapy
Data from phase 3 ION trials for the subset of participants receiving OST versus those not receiving OST.

Source: Grebely J, Mauss S, Brown A, et al. Efficacy and Safety of Ledipasvir/Sofosbuvir With and Without Ribavirin in Patients With Chronic HCV Genotype 1 Infection Receiving Opioid Substitution Therapy: Analysis of Phase 3 ION Trials. Clin Infect Dis. 2016;63:1405-1411.

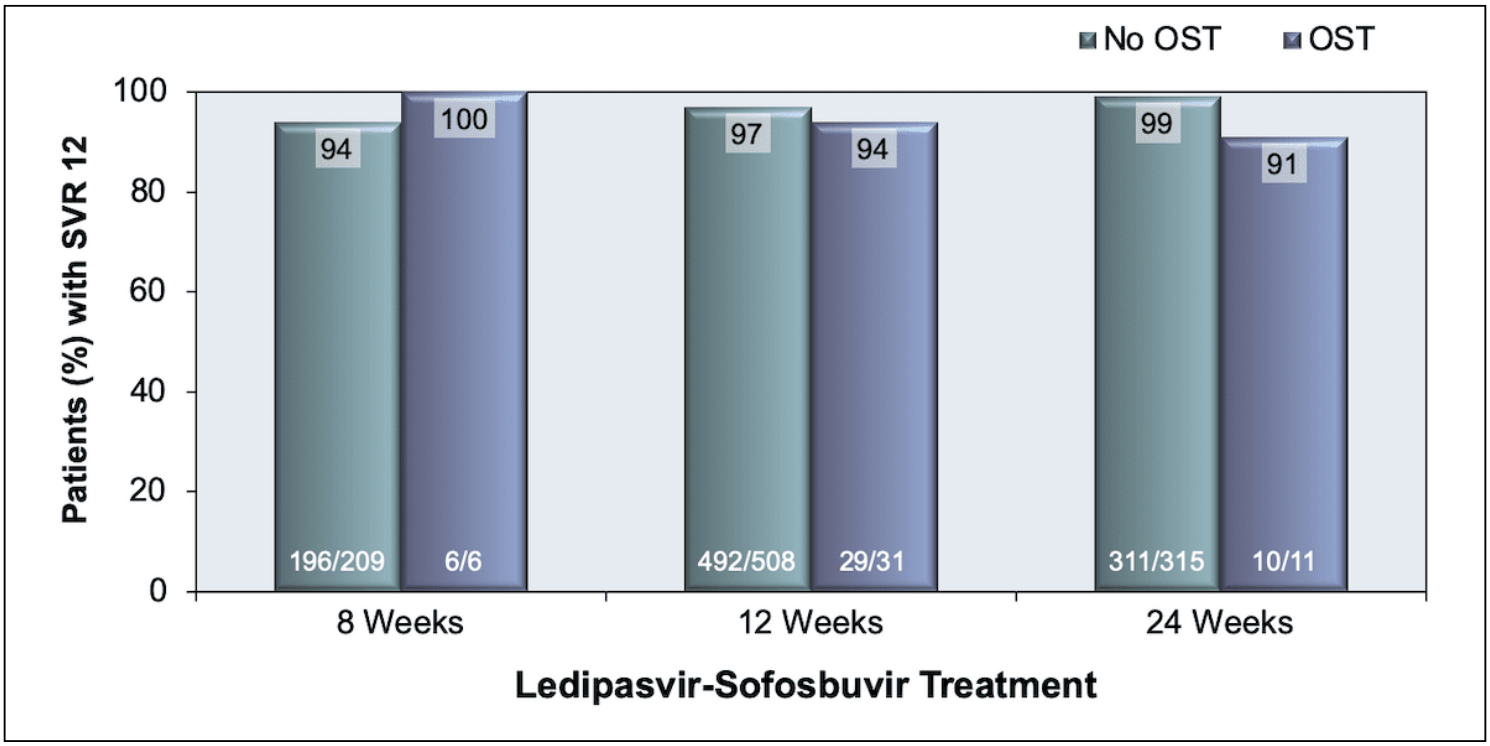

Figure 5 Impact of Alcohol Consumption on HCV Treatment Response

In this study, excessive alcohol consumption was defined as ≥60 g/day for men and ≥40 g/day for women.

Source: Wiley TE, McCarthy M, Breidi L, McCarthy M, Layden TJ. Impact of alcohol on the histological and clinical progression of hepatitis C infection. Hepatology. 1998;28:805-9.

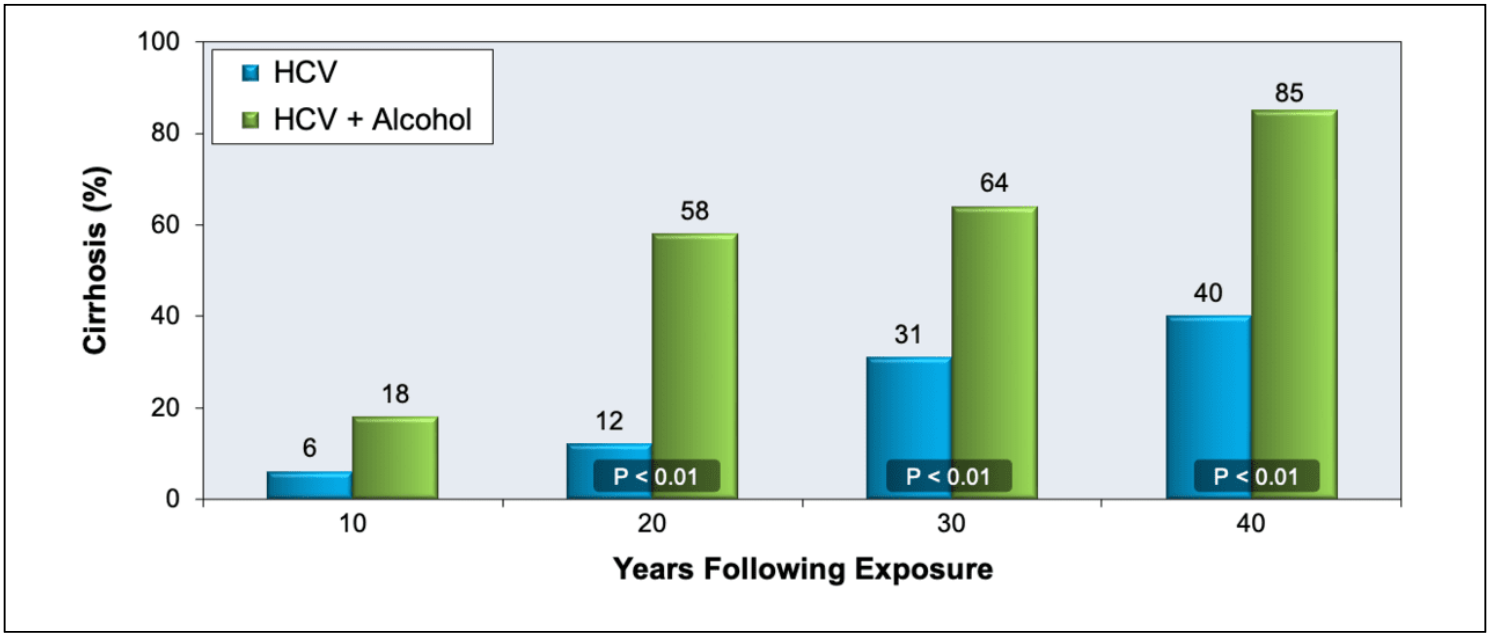

Figure 6 Impact of Alcohol Consumption on HCV Treatment Response

Investigators enrolled 73 patients with chronic hepatitis C (genotypes 1, 2, 3, or 4) who had ongoing alcohol consumption (or abstinence for less than 6 months) and were treated with peginterferon and ribavirin. Abstinence referred to patients off alcohol during the entire treatment period. Low-risk consumption was defined as weekly consumption of no more than 21 standard drinks for men and 14 drinks for women, and no more than 4 by drinking occasion. Excessive consumption was defined as drinking more than the limits defined for low-risk consumption on at least two occasions during the treatment period.

Source: Le Lan C, Guillygomarc'h A, Danielou H, et al. A multi-disciplinary approach to treating hepatitis C with interferon and ribavirin in alcohol-dependent patients with ongoing abuse. J Hepatol. 2012;56:334-40.

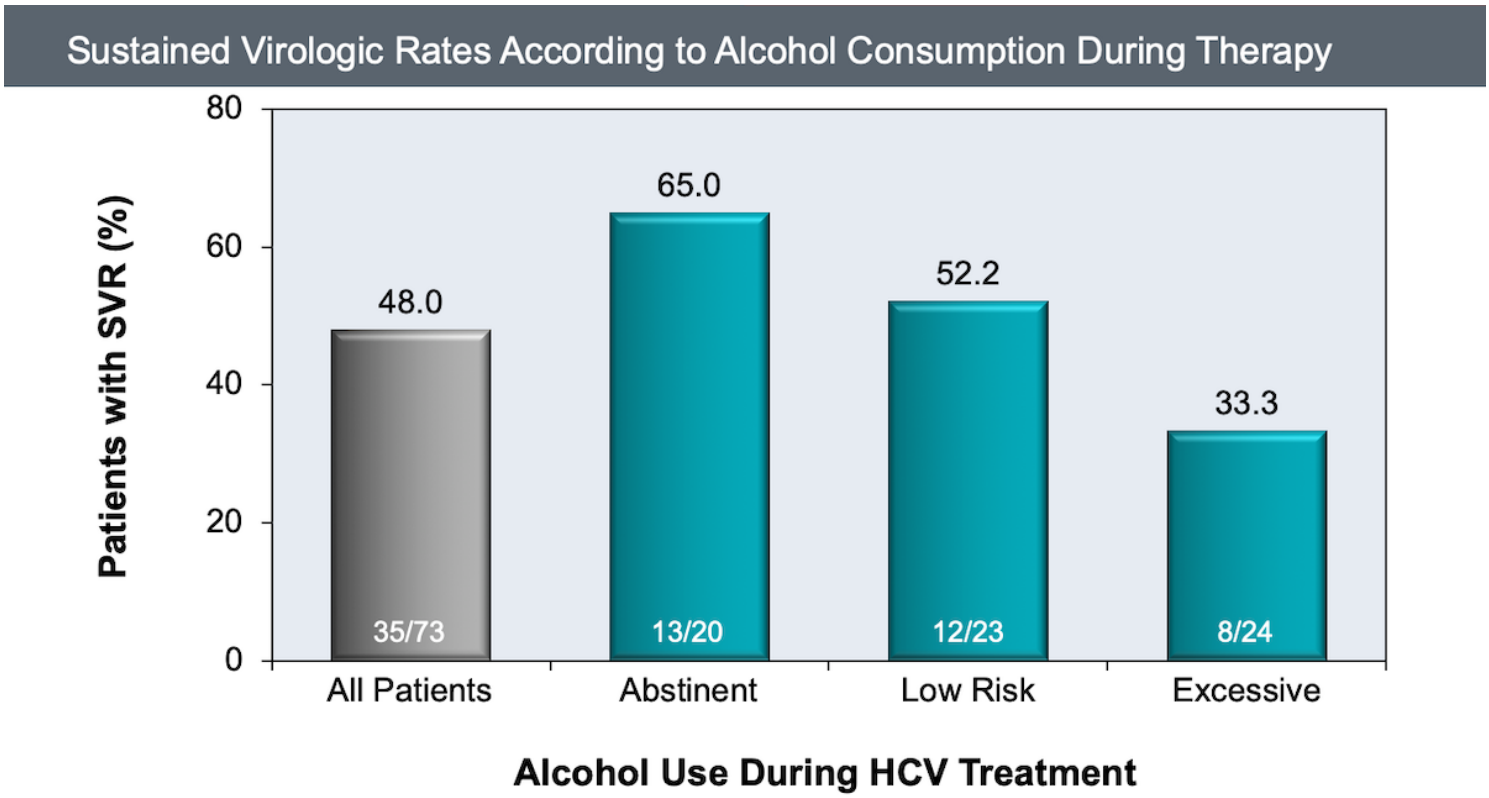

Figure 7 (Image Series) - SVR12 Rates with DAA-Based Treatment, by AUDIT-C Category (Image Series) - Figure 7 (Image Series) - SVR12 Rates with DAA-Based Treatment, by AUDIT-C Category

Image 7A: Response by HCV Genotype

This study was conducted in 2014-2015

Source: Tsui JI, Williams EC, Green PK, Berry K, Su F, Ioannou GN. Alcohol use and hepatitis C virus treatment outcomes among patients receiving direct antiviral agents. Drug Alcohol Depend. 2016;169:101-9.

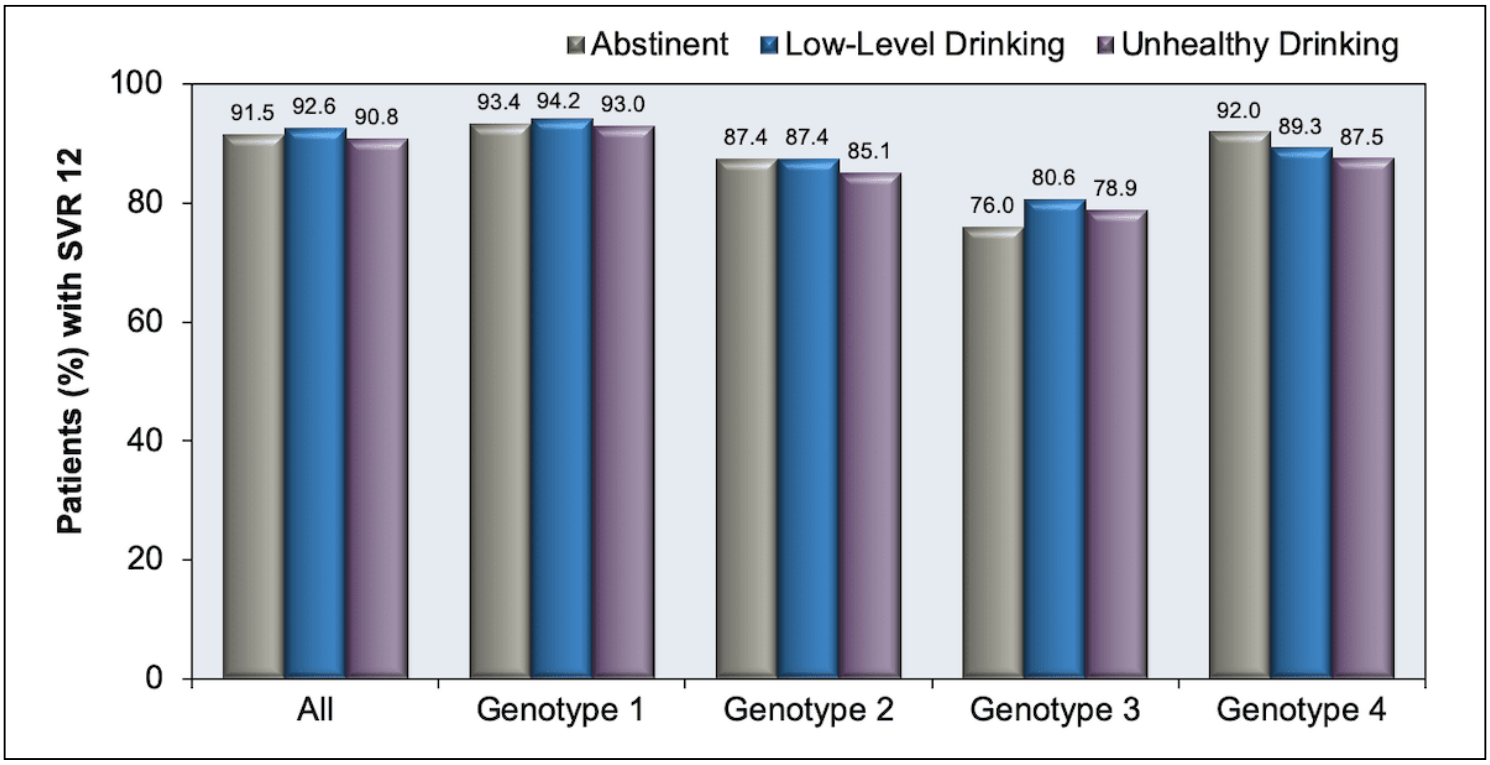

Figure 7 (Image Series) - SVR12 Rates with DAA-Based Treatment, by AUDIT-C Category
Image 7B: Response by Cirrhosis Status

This study was conducted in 2014-2015

Source: Tsui JI, Williams EC, Green PK, Berry K, Su F, Ioannou GN. Alcohol use and hepatitis C virus treatment outcomes among patients receiving direct antiviral agents. Drug Alcohol Depend. 2016;169:101-9.