

# Hepatitis C Guidance 2023 Update: American Association for the Study of Liver Diseases– Infectious Diseases Society of America Recommendations for Testing, Managing, and Treating Hepatitis C Virus Infection

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The Infectious Diseases Society of America and the American Association for the Study of Liver Diseases have collaboratively developed evidence-based guidance regarding the diagnosis, management, and treatment of hepatitis C virus (HCV) infection since 2013. A panel of clinicians and investigators with extensive infectious diseases or hepatology expertise specific to HCV infection periodically reviews evidence from the field and update existing recommendations or introduce new recommendations as evidence warrants.

This update focuses on changes to the guidance since the previous 2020 published update, including ongoing emphasis on recommended universal screening; management recommendations for incomplete treatment adherence; expanded eligibility for simplified chronic HCV infection treatment in adults with minimal monitoring; updated treatment and retreatment recommendations for children as young as 3 years; management and treatment recommendations in the transplantation setting; and screening, treatment, and management recommendations for unique and key populations.

**Keywords.** HCV screening; direct-acting antivirals; HCV guidance; HCV treatment; HCV prevention.

The Infectious Diseases Society of America (IDSA) and the American Association for the Study of Liver Diseases (AASLD) collaboratively initiated the Hepatitis C Virus (HCV) Guidance Project in 2013 to provide clinicians with evidence-based, unbiased, timely guidance regarding diagnosis, treatment, and management of HCV infection. The project includes the web-based HCV guidance platform ([www.hcvguidelines.org](http://www.hcvguidelines.org)) to enable rapid, accessible dissemination of new and/or updated information and recommendations in response to the latest data from the field. The HCV guidance

website (hereafter, the HCV guidance) has been highly successful. From the launch of the HCV guidance in January 2014 through April 2022, the site has been accessed by more than 2 million unique users generating more than 4 million page views. In 2021, the site had more the 194 000 unique users from 201 countries and territories, with most visits originating from the United States, India, Russia, Canada, and Pakistan. Under the umbrella of the HCV guidance, the AASLD-IDSA HCV Guidance Panel (hereafter, the Guidance Panel) also issues regular, periodic published updates to review new or updated data and recommendations as well as an overview of the ever-changing landscape of the HCV epidemic.

Recognizing that viral hepatitis poses a public health threat on par with human immunodeficiency virus (HIV), malaria, and tuberculosis, in June 2016, the World Health Organization (WHO) published its first global health sector strategy and set forth the goal of elimination of viral hepatitis as a major public health threat by 2030 [1]. Specific HCV elimination targets include a 90% reduction in incidence and prevalence, treatment of 80% of eligible persons with chronic infection, a 65% reduction in HCV-related deaths, and universal access to key prevention and treatment services [1]. In response to the WHO's call to action, the National Academies

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- Universal HCV screening is recommended.
- The simplified HCV treatment algorithm now includes persons living with HIV.
- A new algorithm for incomplete treatment adherence is included with a key recommendation for persons who have missed  $\geq 7$  days of DAA therapy.
- HCV treatment is recommended for infected persons residing in jail or prison.
- Emerging data highlight the safety and efficacy of HCV DAA treatment in persons who have undergone solid organ transplantation.

**Figure 1.** Key points in HCV guidance summary. Abbreviations: DAA, direct-acting antiviral; HCV, hepatitis C virus; HIV, human immunodeficiency virus.

of Sciences, Engineering, and Medicine developed a US strategic plan for viral hepatitis elimination [2]. The US Centers for Disease Control and Prevention (CDC) [3] and the US Department of Health and Human Services (DHHS) [4] subsequently developed national implementation strategies and targets commensurate with those set forth by the WHO. Notably, the new and updated recommendations highlighted and discussed in this update both independently and collectively support, promote, and advance accomplishment of HCV elimination.

Major changes in the HCV guidance since the previous 2020 publication [5] featured in this update include an ongoing emphasis on universal HCV screening; new recommendations that address the management of incomplete treatment adherence; updated recommendations regarding simplified treatment with minimal monitoring and expanded eligibility; management and treatment recommendations for solid organ transplant recipients; newly expanded treatment and retreatment recommendations for children and adolescents; and screening, management, and treatment recommendations for unique and key populations. In addition, we highlight key issues critical to HCV management with the mission of HCV elimination in mind. See Figure 1 for key points in this HCV guidance update.

## PROCESS

The HCV guidance was developed and is regularly updated by a volunteer panel of more than 30 infectious diseases and hepatology clinicians and investigators with HCV expertise representing IDSA and AASLD, respectively. Four co-chairs (2 from each society) oversee the work of the Guidance Panel. The HCV guidance undergoes major biannual updates based on a rigorous literature review that encompasses peer-reviewed, published literature and relevant abstracts from national and international scientific conferences. The data are reviewed by section leads, with points of discussion resolved during section and full panel remote meetings.

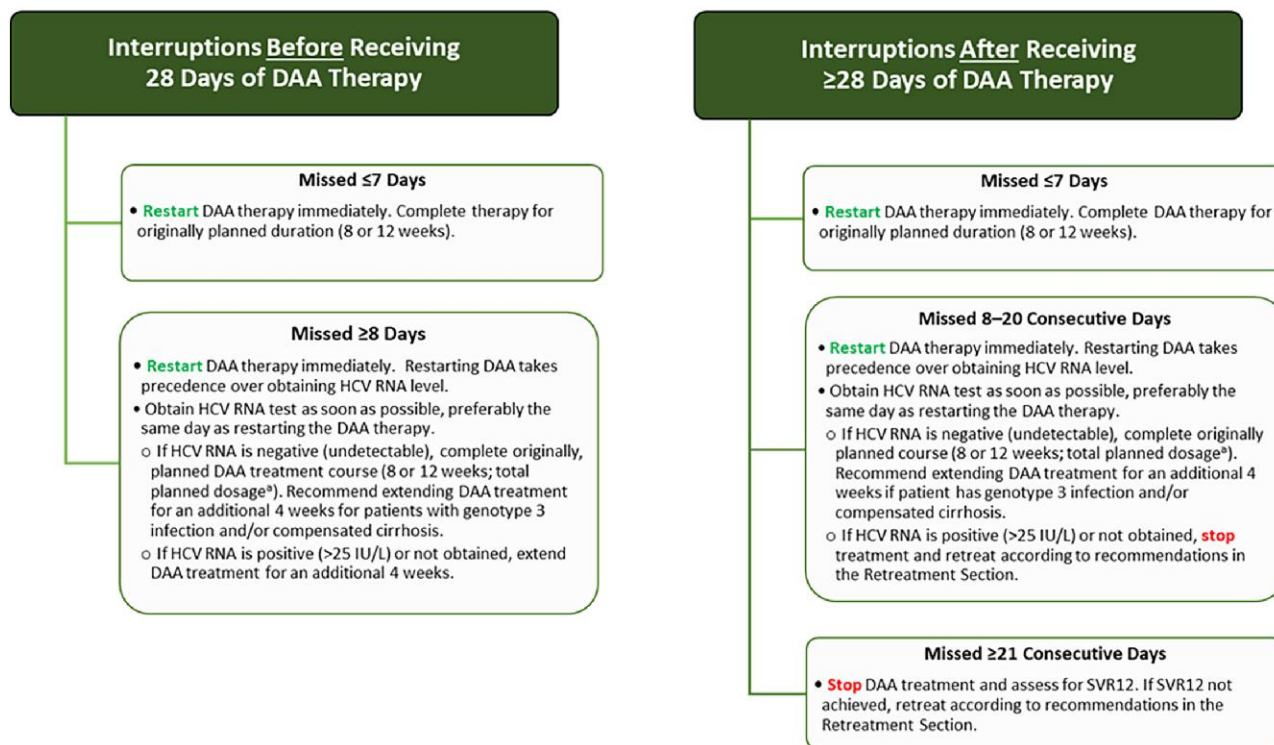
New or updated recommendations are evaluated using a modified scale adapted from the American College of Cardiology and the American Heart Association practice guidelines [6, 7] (see the HCV guidance for further details). All new and updated recommendations are reviewed and approved by the IDSA and AASLD governing boards prior to on-line release or print publication.

## TESTING, EVALUATION, AND MONITORING

### Implementation of Universal HCV Screening

The Guidance Panel first recommended universal HCV screening for all adults aged  $\geq 18$  years in 2019 [5], concomitant with congruous draft recommendations from the US Preventive Services Task Force (USPSTF) and the CDC. The USPSTF subsequently recommended universal HCV screening for adults aged 18 to 79 years in March 2020 [8]. In April 2020, the CDC recommended HCV screening at least once in all adults aged  $\geq 18$  years and for all pregnant persons during each pregnancy, except in settings where HCV prevalence is  $< 0.1\%$  [9]. The rationale for universal HCV screening includes cost-effectiveness [10–13]; improved HCV case finding [8, 9]; shifting epidemiology of HCV infection with incident infections occurring primarily in young adults [14–16]; and the availability of safe, cost-effective direct-acting antiviral (DAA) treatment [17]. Universal screening is a crucial and necessary component of any HCV elimination strategy [1–4] because it is the entry point into the HCV continuum of care [18, 19]. For initial HCV testing, the Guidance Panel recommends HCV antibody screening with reflex HCV RNA testing to establish the presence of active infection (as opposed to spontaneous or treatment-induced viral clearance).

Recommendations without rigorous implementation, however, are ineffectual. HCV screening, diagnosis, and treatment were significantly adversely affected by the coronavirus disease 2019 (COVID-19) pandemic [20]. The number of HCV antibody and HCV RNA tests processed by a large US, multicenter, commercial clinical laboratory decreased precipitously beginning in mid-March 2020 [21], coincident with the US federal government declaring a national state of emergency due to COVID-19 [22]. HCV RNA–positive test results decreased 62% in March 2020 and remained 39% below baseline in July 2020, with a concomitant decline in the number of DAA prescriptions dispensed [21]. Investigators who conducted a similar study in Ontario, Canada, reported comparable decreases in HCV antibody screening and confirmative HCV RNA testing during each of the first 3 waves of the COVID-19 pandemic [23]. The reduced level of HCV testing negatively affecting initiation of HCV treatment appears corroborated by findings from a US national, retrospective study wherein only 23% of people on Medicaid with a positive HCV RNA test between 30 January 2019 and 31 October 2020 initiated DAA treatment



**Figure 2.** Recommended management of DAA treatment interruptions for treatment-naïve patients without cirrhosis or with compensated cirrhosis receiving glecaprevir/pibrentasvir or sofosbuvir/velpatasvir. Abbreviations: DAA, direct-acting antiviral; HCV, hepatitis C virus; SVR12, sustained virologic response 12 weeks after completion of therapy. <sup>a</sup>Extend duration of therapy such that the patient receives the total planned dosage (ie, the total number of daily pills). For example, if a patient missed 10 days of a planned 8-week course of therapy, treatment would be extended to 8 weeks plus 10 days.

within 360 days of diagnosis [24]. A survey conducted among European Association for the Study of the Liver members representing 48 clinical centers also demonstrated decreased HCV testing, diagnosis, and treatment in 2020 compared with 2019 (prepandemic) [25]. Collectively, these findings underscore the critical importance of ongoing, rigorous, universal HCV screening for case identification and linkage to care. In addition, monitoring the proportion of persons who meet steps in the HCV cascade of care will be critical to assessing the quality of HCV care.

#### Management of Incomplete DAA Adherence

Incomplete medication adherence is well known, even in the highly structured clinical trial setting [26, 27]. Recognizing that incomplete DAA treatment may occur in clinical practice and potentially contribute to treatment failure, the HCV guidance includes a new algorithm for the management of incomplete adherence as part of DAA treatment monitoring (Figure 2). The algorithm is applicable only to DAA treatment-naïve persons and, generally, the same patient populations who are eligible for the simplified treatment algorithms described in the following section. Excluded persons with incomplete adherence should be managed in consultation with a specialist in HCV management.

Although few studies have examined incomplete medication adherence in the DAA era, data suggest that it is relatively common, occurring in 11% to 40% of persons on treatment [28–31]. Most episodes of nonadherence appear short-lived. One study demonstrated that 61% of nonadherent episodes lasted 1 to 2 days [31]. These short periods of nonadherence were not associated with virologic failure. Sustained virologic response (SVR) 12 weeks after the completion of treatment (SVR12) was 94% among both adherent and nonadherent participants, where nonadherence was defined as taking <90% of the total prescribed dosage [31]. Longer periods of nonadherence, however, may adversely affect SVR. Investigators who examined the relationship between premature discontinuation of DAA therapy and SVR found that among study participants with F0 to F3 liver disease, SVR12 was 50% in persons who received <4 weeks of DAA treatment compared with 99% SVR12 in those who received ≥4 weeks of treatment [32]. Among participants with compensated cirrhosis, SVR12 rates were 83% and 95% in those who completed <8 weeks of DAA therapy compared with ≥8 weeks of treatment, respectively [32].

Based on these limited findings and the expert consensus of the Guidance Panel, a management algorithm that considers the timing and duration of the nonadherence, as well as specific patient factors (ie, genotype 3 infection and presence of

compensated cirrhosis), is recommended (see [Figure 2](#)). Additional large-scale studies in clinical practice settings that examine the relationship between DAA adherence and SVR12, including the threshold level of adherence below which SVR12 is adversely affected, are sorely needed.

## INITIAL TREATMENT

### Simplified HCV Treatment for Treatment-Naïve Adults

The Guidance Panel continues to strongly recommend universal DAA treatment for all people with acute or chronic HCV infection (except those with a short life expectancy that cannot be remediated by HCV therapy, liver transplantation, or another directed therapy). A key aspect of facilitating the implementation of this recommendation/goal is expanding the pool of clinicians who provide HCV treatment, thereby boosting accessibility and delivery of care. Accordingly and coincident with the accumulation of real-world data and experience with the pangenotypic DAA regimens, the HCV guidance first introduced the simplified treatment algorithms for treatment-naïve persons (without cirrhosis or with compensated cirrhosis) in 2019 [5]. The current update to the simplified treatment algorithms features reduced pretreatment and on-treatment clinician intervention and expanded eligibility of persons who can be treated using these approaches.

Recent data from a global sample of persons undergoing DAA treatment for chronic HCV infection suggest that a minimal on-treatment monitoring approach is safe and effective and leads to an SVR rate that is comparable to that realized with standard monitoring [33]. The minimal monitoring (MINMON) approach was examined in an international, phase 4, open-label,

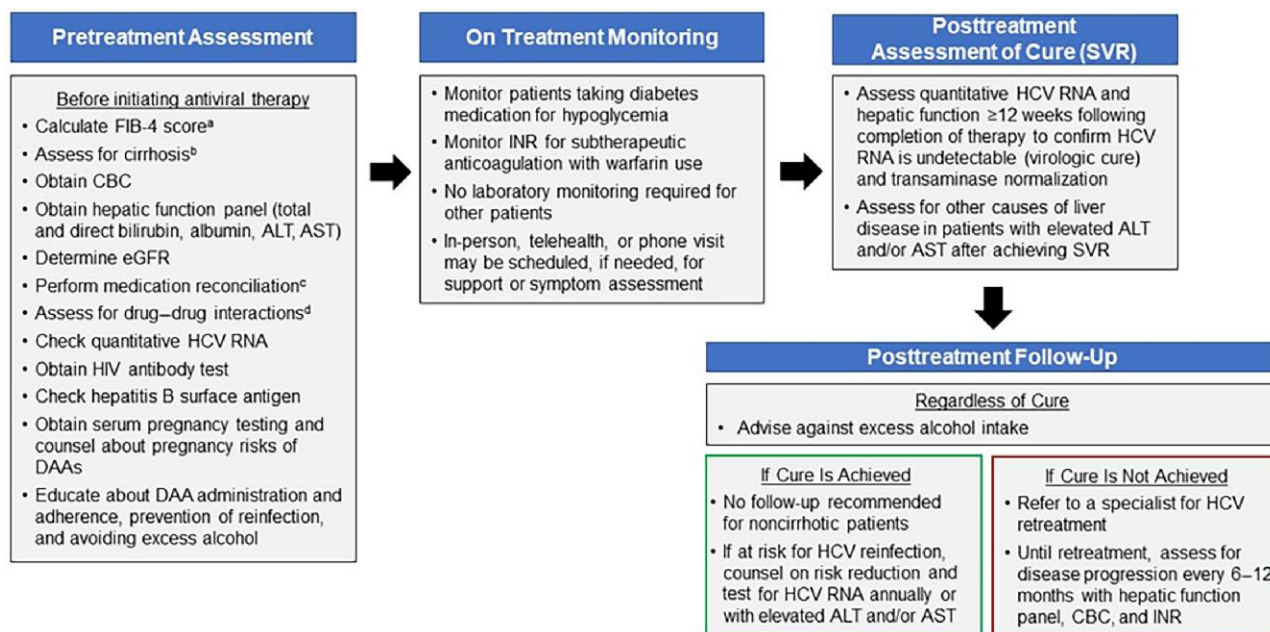
single-arm trial. Four hundred treatment-naïve participants aged  $\geq 18$  years with active HCV infection were enrolled from 38 sites in Brazil, South Africa, Thailand, Uganda, and the United States. Participants included persons with compensated cirrhosis and HIV coinfection. Key exclusion criteria were pregnancy, breastfeeding, and chronic hepatitis B virus (HBV) infection (hepatitis B surface antigen [HBsAg] positive; due to possible risk of HBV reactivation). However, participants with resolved HBV infection (hepatitis B core antibody [anti-HBc] positive, with or without hepatitis B surface antibodies [anti-HBs]) were eligible. Of the 400 enrolled participants, 399 initiated a planned 12-week course of once-daily sofosbuvir (400 mg)/velpatasvir (100 mg). At entry, 42% (166) were living with HIV, 9% (34) had compensated cirrhosis, and 32% (121 of 374) with an available HBV panel had resolved HBV infection. The 4 components of minimal monitoring included no pretreatment genotyping, dispensing the entire treatment course at entry, no scheduled on-treatment visits or laboratory monitoring, and remote contact at week 4 to assess DAA adherence and at week 22 to schedule SVR assessment at week 24. SVR was achieved by 95% (379 of 399) of those who initiated treatment. Fourteen participants experienced a serious adverse event between treatment initiation and week 28; none were treatment-related or led to treatment discontinuation or death [33].

Given the findings of this minimal monitoring study, treatment-naïve persons with HIV/HCV coinfection are newly eligible for a simplified HCV treatment approach. [Figure 3](#) shows the eligibility and exclusion criteria for the simplified HCV treatment approaches. [Figure 4](#) provides an overview of the simplified HCV treatment algorithm for treatment-naïve adults without cirrhosis. [Figure 5](#) reviews the simplified treat-

Who is <u>Eligible</u> for Simplified HCV Treatment Algorithm	Who is <u>Excluded</u> from Simplified HCV Treatment Algorithm
<p>Adults with chronic HCV infection, including persons living with HIV:</p> <ul style="list-style-type: none"> <li>• Infected with any genotype</li> <li>• Have <u>not</u> previously received HCV treatment</li> <li>• Without cirrhosis <u>or</u> with compensated cirrhosis (Child-Pugh A) as determined by: <ul style="list-style-type: none"> <li>– Liver stiffness <math>&gt;12.5</math> kPa by FibroScan</li> <li>– FIB-4 <math>&gt;3.25</math></li> <li>– Noninvasive serologic test<sup>a</sup></li> <li>– Liver biopsy</li> <li>– Liver nodularity or splenomegaly on imaging</li> <li>– Platelet count <math>&lt;150,000/\text{mm}^3</math></li> </ul> </li> </ul>	<p>Adults with chronic HCV infection:</p> <ul style="list-style-type: none"> <li>• Previously received HCV treatment</li> <li>• Hepatitis B surface antigen-positive</li> <li>• Compensated cirrhosis (Child-Pugh A) <u>with</u> end-stage renal disease (eGFR <math>&lt;30</math> mL/min/m<sup>2</sup>)</li> <li>• Current or prior decompensated cirrhosis, defined by Child-Pugh score <math>\geq 7^b</math></li> <li>• Current pregnancy</li> <li>• Known or suspected hepatocellular carcinoma</li> <li>• Prior liver transplantation</li> </ul>

**Figure 3.** Inclusion and exclusion criteria for simplified HCV treatment algorithm. Abbreviations: eGFR, estimated glomerular filtration rate; FIB-4, fibrosis-4 index for liver fibrosis; HCV, hepatitis C virus. <sup>a</sup>Noninvasive serologic tests include HCV FibroSure or enhanced liver fibrosis test. <sup>b</sup>Child-Pugh score based on presence of ascites, hepatic encephalopathy, total bilirubin  $>2.0$  mg/dL, albumin  $\leq 3.5$  g/dL, or international normalized ratio  $\geq 1.7$ .





**Figure 4.** Simplified algorithm for HCV treatment among HCV treatment-naïve adults without cirrhosis. Recommended DAA regimens for this simplified treatment approach include either 8 weeks of glecaprevir (300 mg)/pibrentasvir (120 mg) taken with food or 12 weeks of sofosbuvir (400 mg)/velpatasvir (100 mg). More detailed descriptions of the patient evaluation process and antivirals used for HCV treatment can be found on the HCV guidance website. Abbreviations: ALT, alanine aminotransferase; AST, aspartate aminotransferase; CBC, complete blood count; DAA, direct-acting antiviral; eGFR, estimated glomerular filtration rate; FIB-4, fibrosis-4 index for liver fibrosis; HCV, hepatitis C virus; HIV, human immunodeficiency virus; INR, international normalized ratio; SVR, sustained virologic response. <sup>a</sup>FIB-4 is a noninvasive measure of hepatic fibrosis that is calculated by: (age [years] × AST [U/L]) ÷ (platelet count [109/L] × ALT1/2 [U/L]). <sup>b</sup>A patient is presumed to have cirrhosis if they have a FIB-4 score >3.25 or if they have any of the following from a previously performed test: transient elastography indicating cirrhosis (ie, liver stiffness >12.5 kPa), noninvasive serologic test above the proprietary cutoff indicating cirrhosis (eg, FibroSure, enhanced liver fibrosis test), clinical evidence of cirrhosis (eg, liver nodularity and/or splenomegaly on imaging, platelet count <150 000/mm<sup>3</sup>), or prior liver biopsy showing cirrhosis. <sup>c</sup>Medication reconciliation should record currently prescribed medications, over-the-counter drugs, and herbal/dietary supplements. <sup>d</sup>Drug–drug interaction assessment should be performed using the table in the Monitoring Section of the HCV Guidance website or the University of Liverpool drug interaction checker.

ment algorithm for HCV treatment-naïve adults with compensated cirrhosis.

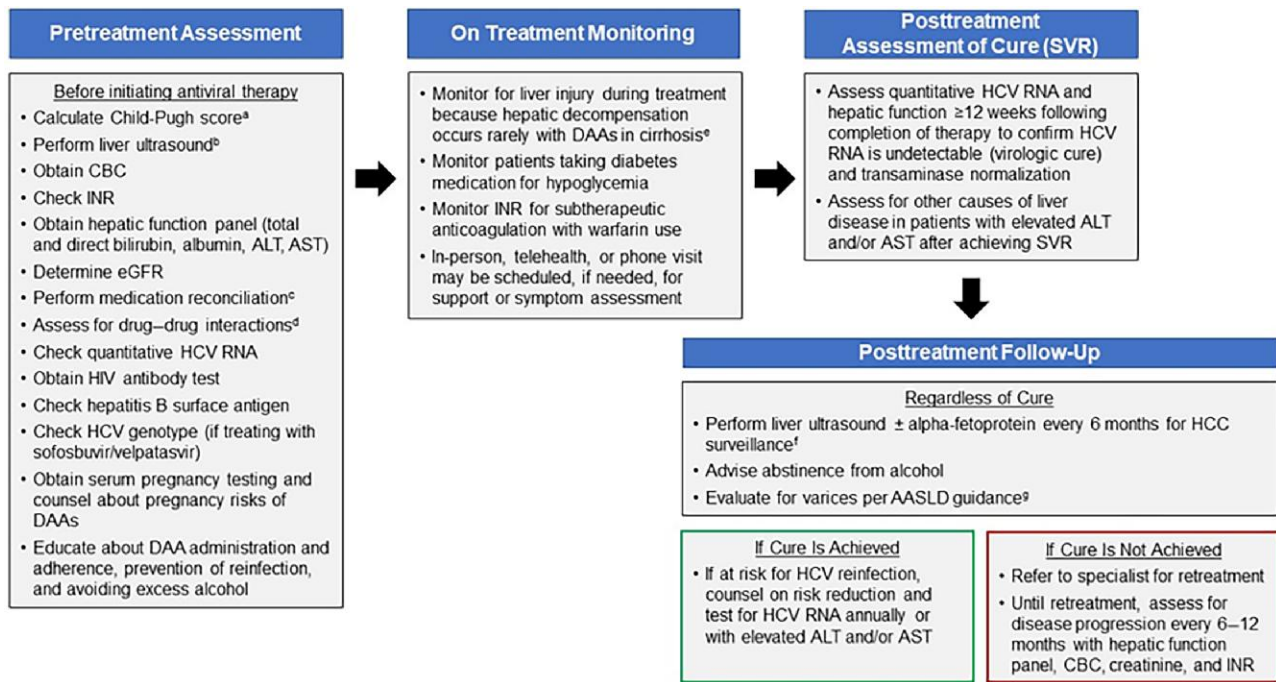
The inclusion of persons living with HIV in the simplified HCV treatment algorithm is consistent with the DHHS Guidelines for the Prevention and Treatment of Opportunistic Infections in Adults and Adolescents with HIV [34]. In this guidance, the decision to expand eligibility to include persons living with HIV was informed by the comparable SVR12 rates in those with and without HIV coinfection in the MINMON study [33], the availability of integrase strand transfer inhibitor–based antiretroviral regimens that mitigate concerns of drug–drug interactions between HIV and HCV medications, and the need to expand treatment access, particularly in the COVID-19 pandemic era.

#### Initial Treatment Regimens

In the current DAA era of hepatitis C treatment, therapy is safe, effective, of relatively short duration, and curative in most people [1, 17]. Widespread use of recommended initial treatment regimens has the potential to substantially reduce hepatitis C prevalence. Given the many benefits of virologic cure, including reduced risk of cirrhosis, hepatocellular carcinoma, liver-

related mortality [35], and all-cause mortality [35–37], expanded use of DAA treatment and the associated probable cure has the capacity to reduce HCV-related disease burden at individual, national, and potentially global levels.

Since the last published update [5], genotypic activity has been added to the hierarchical ranking of treatment regimens (in addition to recommended or alternative, evidence level, and alphabetical order). Table 1 presents a summary of initial treatment recommendations for treatment-naïve adults. Shortening the duration of glecaprevir/pibrentasvir therapy to 8 weeks for persons with compensated cirrhosis is a notable change. The updated recommendation is supported by the findings from the international, single-arm, open-label, phase 3b EXPEDITION-8 trial, titled “Glecaprevir/pibrentasvir for 8 weeks in treatment-naïve patients with chronic HCV genotypes 1–6 and compensated cirrhosis” [38]. Investigators enrolled 343 treatment-naïve participants aged ≥18 years with chronic HCV infection (genotypes 1 through 6) and compensated cirrhosis. Key exclusion criteria included coinfection with HIV and/or HBV or a history of hepatic decompensation. Participants received an 8-week course of once-daily



**Figure 5.** Simplified algorithm for HCV treatment among HCV treatment-naïve adults with compensated cirrhosis. Recommended DAA regimens for this simplified treatment approach include either 8 weeks of glecaprevir (300 mg)/pibrentasvir (120 mg) taken with food for genotypes 1 through 6 or 12 weeks of sofosbuvir (400 mg)/velpatasvir (100 mg) for genotypes 1, 2, 4, 5, or 6. More detailed descriptions of the patient evaluation process and antivirals used for HCV treatment can be found on the HCV Guidance website. Abbreviations: AASLD, American Association for the Study of Liver Diseases; ALT, alanine aminotransferase; AST, aspartate aminotransferase; CBC, complete blood count; DAA, direct-acting antiviral; eGFR, estimated glomerular filtration rate; HCC, hepatocellular carcinoma; HCV, hepatitis C virus; HIV, human immunodeficiency virus; INR, international normalized ratio; SVR, sustained virologic response. <sup>a</sup>Child–Pugh score based on presence of ascites, hepatic encephalopathy, total bilirubin >2.0 mg/dL, albumin ≤3.5 g/dL, or INR ≥1.7. Patients with a Child–Pugh score ≥7 (ie, Child–Pugh B or C) have decompensated cirrhosis; this simplified treatment approach is not recommended for patients with decompensated cirrhosis. <sup>b</sup>Obtain liver ultrasound within 6 months prior to initiating antiviral treatment to exclude hepatocellular carcinoma and subclinical ascites. This simplified treatment approach is not recommended for patients with hepatocellular carcinoma and/or decompensated cirrhosis. <sup>c</sup>Medication reconciliation should record currently prescribed medications, over-the-counter drugs, and herbal/dietary supplements. <sup>d</sup>Drug–drug interaction assessment should be performed using the table in the Monitoring Section of the HCV Guidance website or the University of Liverpool drug interaction checker. <sup>e</sup>Development of jaundice, ascites, spontaneous bacterial peritonitis, variceal hemorrhage, or hepatic encephalopathy may suggest hepatic decompensation. Patients should be referred to a specialist if they develop worsening liver blood tests (eg, total bilirubin, AST, ALT, INR), jaundice, ascites, encephalopathy, or new liver-related symptoms. <sup>f</sup>Ultrasound surveillance for hepatocellular carcinoma (with or without alpha-fetoprotein testing) every 6 months is recommended for patients with cirrhosis, in accordance with AASLD guidance. <sup>g</sup>See AASLD guidance for recommendations regarding the evaluation and management of varices.

glecaprevir (300 mg)/pibrentasvir (120 mg). SVR12 was 98% (335 of 343) in the intention-to-treat (ITT) population. Seven participants experienced a serious adverse event, only 1 of which was treatment-related. One participant who had low baseline leukocyte and neutrophil counts experienced grade 3 leukopenia and neutropenia that presented on posttreatment day 29, which the investigator considered treatment-related. No adverse event led to treatment discontinuation or death [38].

Another significant change is the recommendation that sofosbuvir/velpatasvir/voxilaprevir may be used as an alternative regimen for persons with genotype 3 infection and compensated cirrhosis. This new recommendation is based on findings from the international, open-label, randomized, phase 3 POLARIS-3 clinical trial, titled “Efficacy of 8 weeks of sofosbuvir, velpatasvir, and voxilaprevir in patients with chronic HCV infection: 2 phase

3 randomized trials” and acknowledges limited access to resistance-associated substitution (RAS) testing in some settings [39]. Investigators enrolled 220 DAA treatment-naïve participants with genotype 3 infection and compensated cirrhosis who were randomized to 8 weeks of once-daily sofosbuvir (400 mg)/velpatasvir (100 mg)/voxilaprevir (100 mg) or 12 weeks of once-daily sofosbuvir (400 mg)/velpatasvir (100 mg). SVR12 was 96% in both treatment arms [39].

Initial treatment using elbasvir/grazoprevir for genotype 1a infection was changed from a recommended to an alternative regimen because of the need for baseline RAS testing. Additionally, several regimens are no longer recommended because the therapeutics are either no longer available in the United States and/or the regimens have inferior SVR rates compared with currently recommended DAA regimens. These include sofosbuvir and daclatasvir; sofosbuvir and ribavirin;

**Table 1. Recommendations for Initial Treatment of Hepatitis C Virus–Infected Adults**

Regimen	Genotype	Classification	Duration	Rating	Caveats and Other Considerations
Treatment-naïve without cirrhosis or with compensated cirrhosis Glecaprevir/pibrentasvir	1–6	Recommended	8 wk	I, A <sup>a</sup>	
Sofosbuvir/velpatasvir	1–6	Recommended	12 wk	I, A <sup>b</sup>	For genotype 3 infection with compensated cirrhosis, NS5A RAS testing is recommended. If baseline NS5A RAS Y93H is present, add weight-based ribavirin or choose another recommended regimen.
Ledipasvir/sofosbuvir	1, 4, 5, 6	Recommended	12 wk	I, A <sup>c</sup>	Not recommended for genotype 6e infection if subtype is known.
	1 without cirrhosis	Recommended	8 wk	I, B	Applicable to patients without cirrhosis who are not living with human immunodeficiency virus and whose HCV RNA is <6 million IU/mL.
Elbasvir/grazoprevir	1b, 4	Recommended	12 wk	I, A <sup>d</sup>	
	1a	Alternative	12 wk	I, A	For genotype 1a infection, NS5A RAS testing is recommended. If baseline RASs are present (ie, substitutions at amino acid positions 28, 30, 31, or 93), another recommended regimen should be used.
Sofosbuvir/velpatasvir + weight-based ribavirin	3	Alternative	12 wk	Ila, A	Applicable to genotype 3 infection with compensated cirrhosis and baseline NS5a Y93 RAS.
Sofosbuvir/velpatasvir/voxilaprevir		Alternative	12 wk	Ila, B	Applicable to genotype 3 infection with compensated cirrhosis and baseline NS5a Y93 RAS.
Treatment-naïve with decompensated cirrhosis					
Sofosbuvir/velpatasvir + weight-based ribavirin	1–6	Recommended	12 wk	I, A <sup>e</sup>	Low initial dose of ribavirin (600 mg) is recommended for patients with CTP class C cirrhosis; increase as tolerated.
Sofosbuvir/velpatasvir	1–6	Recommended	24 wk	I, A <sup>e</sup>	Applicable to patients who are ribavirin ineligible.
Ledipasvir/sofosbuvir + weight-based ribavirin	1, 4, 5, 6	Recommended	12 wk	I, A <sup>f</sup>	Low initial dose of ribavirin (600 mg) is recommended for patients with CTP class C cirrhosis; increase as tolerated.
Ledipasvir/sofosbuvir	1, 4, 5, 6	Recommended	24 wk	I, A <sup>f</sup>	Applicable to patients who are ribavirin ineligible.

Recommendations are listed by recommended vs alternative and by genotypic activity, evidence level, and alphabetically.

Abbreviations: CTP, Child–Turcotte–Pugh score; HCV, hepatitis C virus; NS5A, hepatitis C virus nonstructural protein 5A; RAS, resistance-associated substitution.

<sup>a</sup>The level of evidence rating is I, B for persons with compensated cirrhosis.

<sup>b</sup>The level of evidence rating is I, B for persons with genotype 5 or 6 infection.

<sup>c</sup>The level of evidence rating is Ila, B for persons with genotype 5 or 6 infection and those with genotype 4 infection and compensated cirrhosis.

<sup>d</sup>The level of evidence rating is Ila, B for persons with genotype 4 infection and compensated cirrhosis.

<sup>e</sup>Only available data for genotype 6 infection are in persons with compensated cirrhosis.

<sup>f</sup>Only available data for genotypes 5 or 6 infection are in a small number of persons with compensated cirrhosis.

paritaprevir/ritonavir/ombitasvir/dasabuvir; and sofosbuvir, telaprevir, or boceprevir with pegylated interferon and ribavirin.

## RETREATMENT

Although DAA therapy is curative for most persons [1, 17], the small percentage of those in whom treatment fails to result in SVR12 require retreatment. Updated retreatment recommendations focus on DAA treatment failures, specifically, sofosbuvir-based regimen failure; glecaprevir/pibrentasvir failure; and multiple DAA failure, including sofosbuvir/velpatasvir/voxilaprevir or sofosbuvir plus glecaprevir/pibrentasvir (Table 2). Retreatment recommendations for sofosbuvir-based or HCV nonstructural protein 5A (NS5A) inhibitor-based treatment failures in persons with decompensated cirrhosis are also noted in Table 2.

### Sofosbuvir-based Regimen Failure

Generally, persons who have experienced treatment failure with a sofosbuvir-based regimen should be retreated with 12 weeks of sofosbuvir/velpatasvir/voxilaprevir. The exception is

persons with genotype 3 infection and compensated cirrhosis for whom the addition of weight-based ribavirin to the regimen is recommended. This recommendation is supported by data from clinical trials [40, 41] and real-world cohorts [42–45]. Glecaprevir/pibrentasvir for 16 weeks can be used as an alternative retreatment regimen [46–48]. This regimen, however, has not been evaluated in persons with genotype 3 infection and prior sofosbuvir/NS5A inhibitor exposure and is therefore not recommended for these individuals.

### Glecaprevir/Pibrentasvir Failure

For persons with a prior glecaprevir/pibrentasvir treatment failure, retreatment with glecaprevir/pibrentasvir plus ribavirin and sofosbuvir is a recommended retreatment option. This recommendation is supported by findings from the MAGELLAN-3 clinical trial, titled “Retreatment of patients who failed glecaprevir/pibrentasvir treatment for hepatitis C virus infection” [49]. This open-label, phase 3b study evaluated the efficacy and safety of once-daily glecaprevir (300 mg)/pibrentasvir (120 mg) plus sofosbuvir (400 mg) and twice daily weight-based ribavirin for retreatment of persons with a prior

**Table 2. Recommendations for Retreatment of Hepatitis C Virus–Infected Adults by Prior Exposure**

Regimen	Genotype	Classification	Duration	Rating	Caveats and Other Considerations
<b>Sofosbuvir-based treatment failure without cirrhosis or with compensated cirrhosis</b>					
Sofosbuvir/velpatasvir/voxilaprevir	1–6	Recommended	12 wk	I, A	For genotype 3 infection with compensated cirrhosis, add weight-based ribavirin if there are no contraindications.
Glecaprevir/pibrentasvir	1, 2, 4, 5, 6	Alternative	16 wk	I, A	Not recommended for patients with prior exposure to an NS5A inhibitor plus NS3/4A protease inhibitor regimen (eg, elbasvir/grazoprevir).
<b>Glecaprevir/pibrentasvir treatment failure without cirrhosis or with compensated cirrhosis</b>					
Glecaprevir/pibrentasvir + sofosbuvir + weight-based ribavirin	1–6	Recommended	16 wk	IIa, B	
Sofosbuvir/velpatasvir/voxilaprevir	1–6	Recommended	12 wk	IIa, B	For patients with compensated cirrhosis, addition of weight-based ribavirin is recommended (rating IIa, C).
<b>Sofosbuvir/velpatasvir/voxilaprevir or sofosbuvir + glecaprevir/pibrentasvir treatment failure without cirrhosis or with compensated cirrhosis</b>					
Glecaprevir/pibrentasvir + sofosbuvir + weight-based ribavirin	1–6	Recommended	16 wk	IIa, B	Extension to 24 wk should be considered in extremely difficult cases (eg, genotype 3 infection with compensated cirrhosis) or failure following sofosbuvir + glecaprevir/pibrentasvir therapy.
Sofosbuvir/velpatasvir/voxilaprevir + weight-based ribavirin	1–6	Recommended	24 wk	IIa, B	
<b>Sofosbuvir- or NS5A inhibitor–based treatment failure with decompensated cirrhosis</b>					
Sofosbuvir/velpatasvir + weight-based ribavirin	1–6	Recommended	24 wk	II, C <sup>a</sup>	Low initial dose of ribavirin (600 mg) is recommended for patients with CTP class C cirrhosis; increase as tolerated.
Ledipasvir/sofosbuvir + weight-based ribavirin	1, 4, 5, 6	Recommended	24 wk	II, C <sup>b</sup>	Low initial dose of ribavirin (600 mg) is recommended for patients with CTP class C cirrhosis; increase as tolerated.

Recommendations are listed by recommended vs alternative and by genotypic activity, evidence level, and alphabetically.

Abbreviations: CTP, Child–Turcotte–Pugh score; NS3/4A, hepatitis C virus nonstructural protein 3–4A; NS5A, hepatitis C virus nonstructural protein 5A.

<sup>a</sup>Only available data for genotypes 5 or 6 infection are in a small number of persons with compensated cirrhosis.

<sup>b</sup>Only available data for genotype 6 infection are in persons with compensated cirrhosis.

glecaprevir/pibrentasvir treatment failure. Participants with non–genotype 3 infection without cirrhosis and naive to HCV nonstructural protein 3–4A (NS3/4A) protease inhibitors and NS5A inhibitors received 12 weeks of treatment. Those with genotype 3 infection and/or compensated cirrhosis, and/or prior exposure to NS3/4A protease inhibitors and/or NS5A inhibitors received 16 weeks of treatment. SVR12 was 96% (22 of 23) in the ITT population. One patient experienced a serious adverse event unrelated to treatment. No treatment discontinuations or deaths occurred [49].

Treatment with sofosbuvir/velpatasvir/voxilaprevir for 12 weeks is another recommended option in the setting of prior glecaprevir/pibrentasvir treatment failure. Findings from a prospective, nonrandomized, observational study support this recommendation. Investigators enrolled 31 participants with a history of virologic failure with glecaprevir/pibrentasvir therapy. Participants with compensated cirrhosis were included; those with HBV and/or HIV coinfection were excluded. SVR12 was 94% (29 of 31) with 12 weeks of once-daily sofosbuvir (400 mg)/velpatasvir (100 mg)/voxilaprevir (100 mg). Two participants relapsed at week 4 after completion of therapy. No serious adverse events, treatment discontinuations, or deaths occurred [50]. Although the addition of ribavirin was not evaluated in this study, based on prior studies of DAA failures, addition of weight-based ribavirin to the regimen is recommended for persons with compensated cirrhosis.

#### **Multiple DAA Failures, Including Sofosbuvir/Velpatasvir/Voxilaprevir or Sofosbuvir Plus Glecaprevir/Pibrentasvir**

The MAGELLAN-3 clinical trial demonstrated the efficacy (96% SVR12; 22 of 23) of glecaprevir/pibrentasvir plus sofosbuvir and weight-based ribavirin for heavily DAA-experienced patients, although no sofosbuvir/velpatasvir/voxilaprevir failures were included [49]. Among patients with a prior sofosbuvir/velpatasvir/voxilaprevir treatment failure, 16 weeks of glecaprevir/pibrentasvir plus sofosbuvir and weight-based ribavirin is recommended based on the improved resistance profile of pibrentasvir and high response rate seen with this duration of therapy among genotype 3–infected participants in the MAGELLAN-3 trial [49]. Extension to 24 weeks or longer with this regimen should be considered for persons with factors that may reduce the likelihood of achieving SVR (eg, genotype 3 infection with cirrhosis or prior treatment failure with glecaprevir/pibrentasvir plus sofosbuvir). While there are case report data that use this treatment duration [51–54], no clinical trial data are available to support such an approach.

A 24-week course of sofosbuvir/velpatasvir/voxilaprevir plus weight-based ribavirin is also recommended for persons with a prior sofosbuvir/velpatasvir/voxilaprevir treatment failure. Although there are currently no published clinical trial data that examine retreatment with sofosbuvir/velpatasvir/voxilaprevir for patients in whom initial therapy with the same regimen failed, a small retrospective, observational study of persons



with an initial DAA treatment failure and a subsequent retreatment failure with sofosbuvir/velpatasvir/voxilaprevir included 4 persons who received 24 weeks of sofosbuvir/velpatasvir/voxilaprevir rescue therapy (1 with the addition of ribavirin). SVR12 was 100% (4 of 4) in this small group of extensively DAA-experienced patients [53]. The recommendation to extend duration of therapy to 24 weeks in conjunction with weight-based ribavirin when retreating with the same DAA regimen (sofosbuvir/velpatasvir/voxilaprevir) is predominantly based on extrapolation from prior studies that have shown benefit with this strategy in different populations [55].

#### **Retreatment in Patients With Decompensated Cirrhosis**

Retreatment of persons with decompensated cirrhosis and a history of DAA-based treatment failure is limited by the inability to use an NS3/4A protease inhibitor (eg, glecaprevir, grazoprevir, voxilaprevir) in the setting of decompensated cirrhosis. Recommendations to retreat with a 24-week course of either sofosbuvir/velpatasvir plus weight-based ribavirin or ledipasvir/sofosbuvir plus weight-based ribavirin are based on the relatively favorable SVR rates (91% to 100%) with these regimens among patients with compensated cirrhosis and prior DAA failure [55–57].

### **MANAGEMENT OF UNIQUE AND KEY POPULATIONS**

The HCV guidance stresses the importance of addressing the special considerations and unmet needs of unique and key populations to achieve significant reductions in the burden of HCV-related disease. This approach aligns with the WHO strategy for achieving hepatitis C elimination targets, which also emphasizes the importance of focusing efforts on populations disproportionately affected by HCV infection, specifically HIV/HCV-coinfected persons, people who inject drugs (PWID), men who have sex with men (MSM), and incarcerated persons [1]. The HCV guidance additionally focuses on the special considerations and unmet needs of other unique or key populations, namely, individuals with acute HCV infection, pregnant persons, children and adolescents, and solid organ transplant recipients. Recommendations for these populations aim to maximize the potential benefits of often missed opportunities to reduce hepatitis C infection incidence and prevalence, personal and societal disease burden, and HCV-related morbidity and mortality.

#### **HIV/HCV Coinfection**

Treatment-naïve persons living with HIV and HCV (without cirrhosis or with compensated cirrhosis) are newly eligible for DAA therapy using a simplified treatment algorithm (see [Figures 4 and 5](#)). This recommendation is supported by findings from the MINMON clinical trial, titled “A minimal monitoring approach for the treatment of hepatitis C virus infection

(ACTG A5360 [MINMON]): a phase 4, open-label, single-arm trial”. Among the 166 study participants living with HIV and HCV, 95% (157 of 166) achieved SVR12 [33]. Given that people living with HIV are disproportionately affected by HCV infection [58], the reduction in treatment barriers benefits the affected individuals while furthering the goal of HCV elimination.

#### **Acute HCV Infection**

The Guidance Panel reiterates the recommendation that persons with confirmed acute HCV infection (HCV RNA-positive) should be treated the same as those with chronic HCV infection without awaiting possible spontaneous clearance (ie, a test-and-treat approach). Given that the incidence of acute hepatitis C in the United States increased 124% from 2013 through 2020 [59], treatment of this key population is critical to both HCV prevention and elimination.

Findings from studies that evaluated the efficacy of an abbreviated 6 weeks of therapy for acute HCV infection with various DAA regimens, including ledipasvir/sofosbuvir [60, 61], glecaprevir/pibrentasvir [62], and sofosbuvir/velpatasvir [63], have demonstrated largely inferior response rates compared with the standard of care. As such, an abbreviated course of DAA therapy is not recommended for acute HCV infection.

#### **HCV in Pregnancy**

Following the 2018 HCV guidance recommendation for universal hepatitis C screening during pregnancy [64], the USPSTF and CDC issued largely concurrent recommendations in 2020 [8, 9]. In May 2021, the American College of Obstetricians and Gynecologists issued a practice advisory recommending hepatitis screening for all pregnant persons during each pregnancy [65]. The Society for Maternal-Fetal Medicine endorsed that practice advisory and published a concurring recommendation in September 2021 [66]. Given that the DHHS viral hepatitis national strategic plan specifies expanded implementation of universal hepatitis C screening during pregnancy as an important strategy for actualizing HCV elimination [4], the coalescence of screening recommendations for this key population is an important step toward achieving that goal. Treatment recommendations during pregnancy are largely unchanged from the previous update [5]. Although there have been no published large-scale clinical trials to evaluate the safety of DAA therapy during pregnancy, smaller studies and case series have not demonstrated any safety concerns [67–71]. The Guidance Panel suggests that DAA treatment may be considered during pregnancy on a case-by-case basis after a discussion of potential risks and benefits.

#### **HCV in Children**

Strategies to reduce the burden of HCV-related disease have historically focused on the adult population [72]. Data from a

recent modeling study indicate that at least 3.26 million children and adolescents (aged  $\leq 18$  years) are living with HCV infection worldwide [73]. National hepatitis C incidence and prevalence data among children and adolescents in the United States are sparse and/or outdated [73]. However, with the recent increase in HCV infection among women of child-bearing age [15, 74–78] comes a coincident risk of increased cases of mother-to-child transmission [76], the primary route of HCV transmission in children [79, 80].

Treatment for HCV infection in children has been revolutionized in recent years, beginning with the US Food and Drug Administration approval of the first DAAs for adolescents in April 2017 [81, 82] to the June 2021 approval of 2 pan-genotypic regimens (glecaprevir/pibrentasvir and sofosbuvir/velpatasvir) for children as young as 3 years [83, 84]. Efficacy and safety data from therapeutic DAA clinical trials conducted in children are largely comparable to those from studies conducted in adults [85–90]. As such, the Guidance Panel reaffirms its recommendation to treat all HCV-infected children and adolescents aged  $\geq 3$  years with an approved DAA regimen regardless of disease severity. Treatment and retreatment recommendations for children are shown in Tables 3 and 4, respectively.

**Table 3. Recommendations for Initial Treatment of Hepatitis C Virus–Infected Pediatric Patients Without Cirrhosis or With Compensated Cirrhosis**

Regimen	Genotype	Classification	Duration	Rating
Glecaprevir/pibrentasvir	1–6	Recommended	8 wk	I, B
Sofosbuvir/velpatasvir	1–6	Recommended	12 wk	I, B
Ledipasvir/sofosbuvir	1, 4, 5, 6	Recommended	12 wk	I, B

Recommendations are listed by genotypic activity, evidence level, and alphabetically.

## Management of HCV After Solid Organ Transplantation

Clinical trial and real-world data provide robust evidence supporting the safety and efficacy of HCV DAA treatment in patients who have undergone solid organ transplantation [91–95]. Discussion of specific clinical scenarios follows. Table 5 shows HCV treatment recommendations posttransplantation.

### Treatment of Recurrent HCV Infection Post Liver and Kidney Transplantation

The phase 3, single-arm, open-label MAGELLAN-2 trial, titled “Glecaprevir/pibrentasvir treatment in liver or kidney transplant patients with hepatitis C virus infection” evaluated a 12-week course of once-daily glecaprevir (300 mg)/pibrentasvir (120 mg) for the treatment of HCV infection (genotypes 1 through 6) among patients without cirrhosis who had undergone liver or kidney transplantation and were  $\geq 3$  months posttransplantation. Those whose immunosuppressive regimen included cyclosporine  $> 100$  mg/d or prednisone  $> 10$  mg/d were excluded. Treatment-naïve and treatment-experienced (genotypes 1, 2, 4, 5, 6; prior treatment with interferon-based therapy or sofosbuvir plus ribavirin with or without pegylated interferon) participants were included. Treatment-experienced persons with genotype 3 infection were excluded. Overall, SVR12 was 98% (98 of 100). No treatment-related serious adverse events were reported [91].

Sofosbuvir-based regimens have also shown efficacy in persons who have undergone liver or kidney transplantation [93–96]. Investigators who conducted a real-world observational study to evaluate the efficacy and safety of DAA therapy in 179 liver, kidney, or dual liver and kidney transplant recipients reported an SVR12 of 94% (169 of 179) among participants treated with ledipasvir/sofosbuvir. Adverse events, including acute cellular rejection, were rare [93]. A phase 2, open-label study that evaluated 12 weeks of daily sofosbuvir (400 mg)/velpatasvir (100 mg) in 79 HCV-infected (genotypes 1, 2, 3, 4)

**Table 4. Recommendations for Retreatment of Hepatitis C Virus–Infected Pediatric Patients by Prior Exposure and Cirrhosis Status**

Regimen	Genotypes	Classification	Duration	Rating	Cirrhosis Status
Interferon-based regimen ( $\pm$ ribavirin) and/or sofosbuvir treatment failure without NS3/4A protease inhibitor or NS5A inhibitor exposure					
Glecaprevir/pibrentasvir	1, 2, 4, 5, 6	Recommended	8 wk	I, C	No cirrhosis
Glecaprevir/pibrentasvir	1, 2, 4, 5, 6	Recommended	12 wk	I, C	Compensated cirrhosis
Glecaprevir/pibrentasvir	3	Recommended	16 wk	I, C	Without cirrhosis or with compensated cirrhosis
Sofosbuvir/velpatasvir	1–6	Recommended	12 wk	I, C	Without cirrhosis or with compensated cirrhosis
Sofosbuvir/velpatasvir + weight-based ribavirin	1–6	Recommended	12 wk	I, C	Decompensated cirrhosis
NS3/4A protease inhibitor treatment failure without NS5A inhibitor exposure					
Glecaprevir/pibrentasvir	1–6	Recommended	12 wk	I, C	Without cirrhosis or with compensated cirrhosis
NS5A inhibitor treatment failure without NS3/4A protease inhibitor exposure					
Glecaprevir/pibrentasvir	1–6	Recommended	16 wk	I, C	Without cirrhosis or with compensated cirrhosis
Interferon ( $\pm$ ribavirin) plus a hepatitis C virus protease inhibitor treatment failure					
Ledipasvir/sofosbuvir	4, 5, 6	Recommended	12 wk	I, C	Without cirrhosis or with compensated cirrhosis
Ledipasvir/sofosbuvir	1	Recommended	12 wk	I, C	No cirrhosis
Ledipasvir/sofosbuvir	1	Recommended	24 wk	I, C	Compensated cirrhosis

Recommendations are listed by genotypic activity, evidence level, and alphabetically.

Abbreviations: NS3/4A, hepatitis C virus nonstructural protein 3–4A; NS5A, hepatitis C virus nonstructural protein 5A.

**Table 5. Recommendations for Hepatitis C Virus Treatment Posttransplantation**

Regimen	Genotypes	Classification	Duration	Rating	Caveats and Other Considerations
<b>Recurrent HCV post liver transplant without cirrhosis</b>					
Glecaprevir/ pibrentasvir	1–6	Recommended	12 wk	I, B	
Sofosbuvir/ velpatasvir	1–6	Recommended	12 wk	I, B	
Ledipasvir/ sofosbuvir	1, 4, 5, 6	Recommended	12 wk	I, B	
<b>Recurrent HCV post liver transplant with compensated cirrhosis</b>					
Sofosbuvir/ velpatasvir	1–6	Recommended	12 wk	I, B	
Glecaprevir/ pibrentasvir	1–6	Recommended	12 wk	I, C	
Ledipasvir/ sofosbuvir	1, 4, 5, 6	Recommended	12 wk	I, A	
<b>Recurrent HCV post kidney transplant without cirrhosis or with compensated cirrhosis</b>					
Glecaprevir/ pibrentasvir	1–6	Recommended	12 wk	I, A <sup>a</sup> IIa, C <sup>b</sup>	
Sofosbuvir/ velpatasvir	1–6	Recommended	12 wk	IIa, C	
Ledipasvir/ sofosbuvir	1, 4, 5, 6	Recommended	12 wk	I, A	
Elbasvir/ grazoprevir	1, 4	Alternative	12 wk	I, B	Limited to patients without baseline NS5A RASs for elbasvir.
<b>HCV-uninfected recipients of liver grafts from HCV-viremic donors</b>					
Glecaprevir/ pibrentasvir	1–6	Recommended	12 wk	I, C	Timing: initiate treatment within the first 2 wk posttransplant, preferably within the first week.
Sofosbuvir/ velpatasvir	1–6	Recommended	12 wk	I, C	Timing: initiate treatment within the first 2 wk posttransplant, preferably within the first week.
<b>HCV-uninfected recipients of non-liver solid organs from HCV-viremic donors</b>					
Glecaprevir/ pibrentasvir	1–6	Recommended	8 wk <sup>c</sup>	I, C	Timing: initiate treatment prior to HCV RNA results, immediately pretransplant or day 0 posttransplant, if possible. Otherwise, begin on day 0 to within the first week posttransplant when clinically stable.
Sofosbuvir/ velpatasvir	1–6	Recommended	12 wk	I, C	Timing: initiate treatment prior to HCV RNA results, immediately pretransplant or day 0 posttransplant, if possible. Otherwise, begin on day 0 to within the first week posttransplant when clinically stable.

Recommendations are listed by genotypic activity, evidence level, and alphabetically.

Abbreviations: HCV, hepatitis C virus; NS5A, hepatitis C virus nonstructural protein 5A; RAS, resistance associated substitution.

<sup>a</sup>Rating is based on evidence for persons without cirrhosis.

<sup>b</sup>Rating is based on evidence for persons with compensated cirrhosis.

<sup>c</sup>If treatment initiation is delayed beyond the first week after transplant, treatment should be extended to 12 weeks.

liver transplant recipients demonstrated a similar response rate with an SVR12 of 96% (76 of 79). No treatment-related serious adverse events, transplant rejection episodes, or deaths occurred during the study period [94].

Important drug–drug interactions unique to the posttransplant setting should be addressed prior to initiation of DAA therapy. Cyclosporine significantly increases the area under the curve of elbasvir/grazoprevir [97, 98] as well as sofosbuvir/velpatasvir/voxilaprevir [99] and should not be coadministered with these regimens. Coadministration of glecaprevir/pibrentasvir and cyclosporine >100 mg/d is also not recommended [83].

#### ***Treatment of HCV-Uninfected Transplant Recipients Receiving Organs From HCV-Viremic Donors***

A large disparity persists among people in need of solid organ transplantation and available deceased donor organs [100].

Given that available data support the safety and efficacy of DAA therapy in the posttransplant setting, many transplant centers have begun using solid organs from HCV-positive donors for HCV-negative recipients to increase the pool of available organs [101–106]. The pool of HCV-positive donors includes both HCV-viremic donors (ie, HCV RNA-positive) and HCV-seropositive donors (ie, HCV antibody-positive, HCV RNA-negative [nonviremic]). The use of HCV-positive organs has been shown to be an effective strategy for increasing access to transplantation and reducing wait-list time and overall mortality [107–110].

***Timing and Treatment of HCV-Viremic Liver Grafts in Nonviremic Recipients.*** Emerging data support HCV treatment as early as possible when transplanting an HCV-viremic liver graft into an HCV-seronegative recipient [111]. In a recent multicenter

prospective study, 34 HCV-seronegative liver transplant patients underwent transplantation using organs from HCV-positive donors (20 viremic, 14 nonviremic). All recipients of grafts from HCV-viremic donors became viremic by day 3 posttransplantation. DAA treatment was initiated in these graft recipients a median of 27.5 days after transplantation. SVR12 was 100% (20 of 20). One patient developed acute HCV-related membranous nephropathy on postoperative day 18 (prior to initiation of DAA therapy), ultimately resulting in end-stage renal disease requiring dialysis despite achieving SVR12 [112]. This case highlights the importance of early initiation of DAA therapy posttransplantation to avoid HCV-related complications. The Guidance Panel recommends initiating therapy at least within 2 weeks after transplantation but preferably within 1 week when the patient is clinically stable.

An abbreviated duration of DAA therapy is currently not recommended for recipients of organs from HCV-viremic donors due to lack of data demonstrating efficacy. The large reservoir of HCV in a transplanted liver graft may be responsible for the lack of efficacy.

**Timing and Treatment of HCV-Viremic Non-Liver Grafts in Nonviremic Recipients.** HCV treatment should occur as early as possible in HCV-seronegative patients who undergo transplantation with a non-liver graft from an HCV-viremic donor. This strategy reduces the likelihood of hepatic and extrahepatic HCV-related complications in the immediate posttransplant period. The phase 4, open-label, multicenter MYTHIC clinical trial evaluated the efficacy and safety of 8 weeks of once-daily glecaprevir (300 mg)/pibrentasvir (120 mg) in 30 HCV-negative kidney transplant recipients who underwent transplantation using a graft from an HCV-viremic donor [113]. Treatment initiation occurred 2 days to 5 days posttransplantation (target was 3 days). All 30 participants achieved SVR12; no HCV-related serious adverse events were reported [113]. Based on this study and others showing benefit(s) associated with early HCV treatment [113–116], use of a prophylactic (immediately prior to transplantation or day 0 posttransplantation) or preemptive (day 0 to day 7 posttransplantation; as soon as the patient is clinically stable) strategy for initiation of DAA treatment is recommended for HCV-negative recipients of a non-liver solid organ graft from an HCV-viremic donor. Note that neither approach requires demonstration of HCV viremia in the transplant recipient.

Shorter durations of DAA-based therapy in this setting are currently under investigation with promising results. These practices, however, are currently not recommended outside of a clinical trial [115, 117, 118].

**Outcomes and Process in Transplantation Using HCV-Viremic Donor Grafts in HCV-Seronegative Recipients.** Data evaluating longer-

term patient outcomes after transplantation with an HCV-viremic donor organ have shown encouraging results. Among 51 dual heart/kidney transplant recipients undergoing transplantation with organs from HCV-viremic donors, 1-year survival was comparable to survival in those who received organs from nonviremic donors [119]. Another study that evaluated outcomes among multiorgan transplant recipients (heart/kidney, heart/lung, heart/liver) demonstrated similar 1-year survival among recipients of organs from HCV-viremic donors compared with those who received organs from HCV-negative donors [106].

In an analysis of the United Network for Organ Sharing database, HCV-negative liver transplant patients who received the graft from an HCV-positive donor (viremic and nonviremic) were shown to have superior 1-year graft survival rates compared with those who received a graft from an HCV-negative donor [120]. Notably, HCV-positive donors were statistically significantly younger than their HCV-negative counterparts. Multivariate analysis demonstrated that donor age, but not donor HCV status, was an independent predictor of 1-year graft survival [120].

Extensive informed consent, as recommended by the American Society of Transplantation Consensus Panel [121], and shared decision-making between the patient and clinical team should occur prior to transplantation of an HCV-viremic organ into an HCV-negative recipient. Patients should understand the risk of HCV infection, risk to caregivers from needlestick exposures, as well as success rates and risks of DAA-based therapy [115, 121–125]. Given the breadth of safety and efficacy data now available, institutional review board–approved protocols are no longer required. However, based on the unique factors noted, transplant centers should have a specific HCV consent and follow-up process in place.

### People Who Inject Drugs

Injection drug use (IDU) is the most common risk factor for HCV infection in North America and Europe. The HCV seroprevalence among PWID ranges from 18% to 88%, depending on geographic location [126] and duration of IDU exposure [127, 128]. IDU accounts for approximately 70% of new HCV infections [59]. Thus, the growing opioid epidemic has become an important force in the perpetuation of the HCV epidemic [1, 2, 4, 14, 16, 59]. Consequently, achieving the goal of HCV elimination depends heavily on diagnosing and treating HCV infection in PWID and on implementing harm reduction strategies to prevent future infections [1, 2, 4, 122, 129–132]. Data from Australia support the efficacy of the treatment-as-prevention approach among PWID. After implementation of unrestricted access to DAA therapy in 2016, the proportion of PWID diagnosed with active HCV infection who were treated increased from 3% to 47%, while the



proportion of those with HCV viremia declined from 44% to 17% [133].

Annual HCV testing is recommended for PWID with ongoing IDU regardless of either no prior testing or past negative testing. Substance use disorder treatment programs and needle/syringe exchange programs should offer routine, opt-out HCV antibody testing with confirmatory HCV RNA testing and linkage to care for those determined to be HCV-infected [132, 134]. PWID with HCV infection should be counseled about measures to reduce the risk of transmission to others and offered linkage to harm reduction services, including intranasal naloxone, needle/syringe service programs, medications for opioid use disorder, and other substance use disorder treatment programs.

Clinical trials and observational studies of PWID reporting current IDU at the start of HCV treatment and/or continued use during therapy demonstrate SVR12 rates approaching 95% [135–140]. The Guidance Panel strongly asserts that active or recent drug use or concern for reinfection is not a contraindication to HCV treatment. At least annual HCV RNA testing is recommended for PWID with recent IDU after they have spontaneously cleared HCV infection or have been successfully treated [141–144].

#### **Men Who Have Sex With Men Not Living With HIV**

While the increased risk of HCV infection among MSM living with HIV is well known [145], acute HCV infections have also been reported among MSM not living with HIV who present for HIV preexposure prophylaxis (PrEP) [146, 147]. HCV testing at HIV PrEP initiation and at least annually thereafter (while on PrEP) is recommended for MSM not living with HIV. All MSM should be counseled about the risk of sexual HCV transmission with high-risk sexual and drug use practices and educated about measures to prevent HCV infection or transmission [148, 149].

Antiviral treatment for HCV-infected MSM should be coupled with ongoing counseling about the risk of HCV reinfection and education about methods to reduce HCV reinfection risk after cure [150]. At least annual (and risk-based, if indicated) HCV RNA testing is recommended for all high-risk sexually active MSM after successful treatment or spontaneous clearance of HCV infection [151, 152].

#### **Persons in Correctional Settings**

Recent cross-sectional surveys suggest that the HCV seroprevalence among incarcerated populations in the United States ranges from 3.0% to 34.6% [153], which exceeds the 1.7% HCV seroprevalence in the general population [154]. More than 90% of these persons are eventually released and reenter the general population where they can contribute to HCV spread in the community [155, 156] and may have little contact with the healthcare system [157, 158]. Given the high HCV

prevalence among persons in the US correctional system, the success of the US HCV elimination effort depends on identifying infected individuals in jails and prisons, linking these persons to medical care for HCV management, and providing access to antiviral treatment [2, 159]. Jails and prisons should therefore implement opt-out HCV testing that consists of HCV antibody testing followed by confirmatory HCV RNA testing if antibody-positive. Universal opt-out testing of incarcerated persons for chronic HCV is highly cost-effective and has been shown to reduce ongoing HCV transmission and the incidence of advanced liver disease [160].

DAA treatment for chronic HCV infection is feasible within jail and prison settings and would aid the HCV elimination effort [161, 162]. Chronically infected persons residing in jails should receive counseling about HCV infection and be provided linkage to follow-up community healthcare for evaluation of liver disease and treatment upon release [163–166]. Those whose jail sentence is sufficiently long to complete a recommended course of DAA therapy should receive that treatment while incarcerated [161]. Chronically infected individuals in prison should receive DAA therapy according to AASLD–IDSA guidance while incarcerated [162, 167]. Jails and prisons should facilitate continuation of HCV therapy for persons on HCV treatment at the time of incarceration. HCV treatment in correctional settings is cost-effective because DAAs halt progression of HCV-related liver disease and decrease the risk of cirrhosis, hepatic decompensation, and hepatocellular carcinoma, offsetting future healthcare costs from liver and non-liver complications [168].

Upon release from a correctional facility, HCV-infected persons with advanced hepatic fibrosis or cirrhosis should be provided linkage to community healthcare for surveillance for HCV-related complications. To prevent HCV reinfection and reduce the risk of progression of HCV-associated liver disease, correctional facilities should provide harm reduction and evidence-based treatment for underlying substance use disorders [169]. Addressing hazardous alcohol use among persons with chronic HCV in a correctional setting may help slow liver disease progression, decrease HCV transmission, and might reduce recidivism.

#### **Notes**

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