

Counseling Persons with Chronic HCV Infection

This is a PDF version of the following document:

Module 2: [Evaluation, Staging, and Monitoring of Chronic Hepatitis C](#)

Lesson 3: [Counseling Persons with Chronic HCV Infection](#)

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<https://www.hepatitisC.uw.edu/go/evaluation-staging-monitoring/counseling-liver-health/core-concept/all>.

Background

The following lesson provides guidance on how to counsel persons with chronic hepatitis C virus (HCV) infection on the use of over-the-counter pain medications, iron and vitamin supplements, and complementary and alternative therapies. This section also provides guidance on diet and weight management, as well as the use of alcohol, cannabis, and tobacco.

Over-the-Counter Pain Medications

Because many medications are metabolized through the liver, it is important for the medical provider to know all the medications a person with chronic HCV is taking, including over-the-counter medications. A current medication list should be solicited frequently.

Acetaminophen

Acetaminophen can cause clinically important hepatotoxicity, either through an acute overdose or when taken on a regular basis (even at lower doses).[\[1,2,3,4\]](#) In one large study that examined acetaminophen-related acute liver failure, individuals taking less than 4 grams per day of acetaminophen accounted for 7% of the cases.[\[3\]](#) Among healthy adults taking 4 grams per day for 14 days, 38% developed alanine aminotransferase (ALT) values in excess of 3 times the upper limit of normal.[\[2\]](#) In contrast, healthy adults who took 1 gram of acetaminophen twice daily for 12 weeks had only minor elevations in aminotransferase levels.[\[5\]](#) Studies involving persons with chronic HCV have shown an increased risk of acute liver injury among individuals with chronic HCV following acetaminophen overdose.[\[6,7\]](#) Concurrent alcohol use further increases the chance of acute or chronic acetaminophen-induced hepatotoxicity in persons with chronic HCV infection.[\[6,7\]](#) Formal guidelines for the safe use of acetaminophen in persons with HCV infection do not exist, but some experts have issued general recommendations.[\[8,9,10,11,12\]](#)

Recommendations

- Low dosages of acetaminophen (up to 2 grams per day) can safely be used in most persons with chronic HCV infection without cirrhosis.
- Individuals with chronic HCV infection and cirrhosis should limit their intake of acetaminophen to 1-2 grams per day.
- Persons with chronic HCV who drink excess alcohol should avoid taking acetaminophen.
- Clinicians should inform persons with chronic HCV infection that many narcotic combination pills and over-the-counter cold and flu medications may contain acetaminophen.
- Individuals with chronic HCV who frequently take acetaminophen should have laboratory monitoring for hepatotoxicity every 3 to 6 months.

Aspirin and Nonsteroidal Anti-inflammatory Medications

Aspirin and nonsteroidal anti-inflammatory drugs (NSAIDs) are generally safe for persons who have chronic HCV without cirrhosis, when taken at standard doses. For persons with cirrhosis, NSAIDs and aspirin are best avoided, especially for those with decompensated cirrhosis.[\[12,13\]](#) The American Association for the Study of Liver Diseases (AASLD) recommends persons with cirrhosis and ascites to avoid taking NSAIDs, except in special circumstances.[\[14\]](#) In persons with cirrhosis, particularly decompensated cirrhosis, the use of NSAIDs and aspirin may further increase the inherent risk these patients have of developing nephrotoxicity and gastrointestinal bleeding.[\[12,15\]](#) There are limited data on the use of topical NSAIDs in patients with cirrhosis, but these have limited systemic absorption and are likely safe to use in patients with underlying cirrhosis.[\[12\]](#)

Recommendation

- Individuals with chronic HCV who do not have cirrhosis can take aspirin or NSAIDs at low or standard recommended dosages.
- Persons with chronic HCV infection and cirrhosis should, in general, avoid taking NSAIDs or aspirin.
- Individuals with cirrhosis who have short-term, minor pain should take acetaminophen in this setting as long as the acetaminophen dose does not exceed 2 grams per day. If a person with compensated cirrhosis has joint or musculoskeletal pain unresponsive to acetaminophen, NSAIDs can be used for a very brief period of time (less than 3 days), if given at the lowest daily dose possible. In this situation, topical NSAIDs can also be considered.

- Persons with chronic HCV and decompensated cirrhosis should not take aspirin or NSAIDs.

Iron and Vitamin Supplements

Iron

In persons with chronic hepatitis C infection, mild to moderate hepatic iron overload is common.[[16,17](#)] In some studies, excess hepatic iron in persons with chronic HCV has been associated with accelerated fibrosis, whereas other studies have not shown a correlation of iron and hepatic fibrosis progression in persons with chronic HCV.[[18,19,20,21](#)]

Recommendation

- Based on the potential negative impact of iron in persons with chronic HCV, many experts have recommended these individuals avoid taking iron supplements or a daily multivitamin that contains iron, unless a compelling reason exists to regularly take iron, such as iron deficiency anemia from gastrointestinal bleeding.[[15](#)]

Vitamin A

Vitamin A is a fat-soluble vitamin that can be obtained in the diet as provitamin A (mostly plant sources) and preformed vitamin A (animal sources or supplements). Intake of vitamin A at levels contained in a multivitamin does not cause hepatotoxicity. Vitamin A deficiency is often made on a clinical basis, but is supported by either a serum retinol level less than 20 micrograms/dL or a molar ratio of retinol to retinol-binding protein less than 0.8.[[22](#)] Vitamin A deficiency is uncommon in the United States. Chronic ingestion of mega-doses of vitamin A, especially in excess of 25,000 international units per day, can potentially cause severe hepatotoxicity.[[23,24,25,26](#)]

Recommendations

- Vitamin A is a fat-soluble vitamin and should only be taken at standard doses of less than 5,000 international units per day.
- Intake of high doses of vitamin A, such as a dose greater than 25,000 international units per day, is not recommended.

Vitamin D

Vitamin D deficiency is common in persons with chronic HCV infection, particularly those individuals with cirrhosis.[[27,28](#)] Some experts have suggested that vitamin D has anti-inflammatory and antifibrotic properties in persons with chronic HCV infection, but results from these studies are mixed.[[29,30](#)] In one study that examined 25-hydroxyvitamin D levels in 218 persons receiving direct-acting antiviral (DAA) therapy, investigators found no correlation of vitamin D levels and sustained virologic response (SVR) rates.[[28](#)] The Institute of Medicine defines vitamin D deficiency as a serum 25-hydroxyvitamin D level less than 20 ng/mL, but other societies, including the Endocrine Society, the National Osteoporosis Foundation, the International Osteoporosis Foundation, and the American Geriatric Society recommend using less than 30 ng/mL as the cutoff for vitamin D deficiency.[[31,32,33,34](#)]

Recommendations

- There are no liver-specific reasons to take vitamin D supplementation, but persons with chronic HCV infection who are deficient in vitamin D should have vitamin D replacement therapy as per standard of care.
- For otherwise healthy adults, the Institute of Medicine has a recommended vitamin D dietary allowance of 600 IU per day, with an upper level of intake of 4,000 IU/day.[[31](#)] Some experts recommend that persons who do not have regular sun exposure take 800 IU per day of vitamin D.

There are no known hepatotoxic effects of excess vitamin D dosing.[\[35\]](#)

- For individuals with more severe vitamin deficiency (e.g., serum 25-hydroxyvitamin D level less than 10 ng/mL), many experts recommend initial replacement therapy consisting of 50,000 IU of vitamin D once weekly for 8 weeks, followed by a maintenance dose of 800 to 1000 IU per day. For milder vitamin D deficiency, a daily dose of 1000 IU per day may be sufficient.

Complementary and Alternative Therapies

Complementary and alternative therapies are frequently used among adults in the United States, including persons with chronic HCV.[36,37,38] Some of these complementary and alternative therapies, however, may have harmful effects on the liver or cause serious interactions with medications used for HCV therapy.[39,40] At the initial visit for HCV care and at regular intervals thereafter, clinicians should obtain a complete list of the individual's prescription medications, over-the-counter medications, complementary and alternative therapies, and dietary and herbal supplements. For patients taking complementary or alternative therapies, clinicians should discuss whether it is wise for them to continue taking these alternative and complementary therapies. The list of complementary and alternative therapies should be reviewed again during HCV treatment so as not to miss potential drug interactions. The [National Center for Complementary and Alternative Medicine](#) (NCCAM) resource [LiverTox](#) has information on the safety and efficacy of dietary and herbal supplements.[41] The following summarizes several common complementary and alternative therapies that have been used by persons living with chronic HCV infection.

General Recommendation

- Given the extremely high SVR rates with current DAA therapy, it is unlikely that any of the complementary and alternative therapies listed below would provide any significant benefit for HCV treatment. In addition, complementary and alternative medications may cause drug interactions with DAA medications. Accordingly, we do not recommend the use of any complementary or alternative therapies for persons with HCV infection.

Ginseng

Ginseng includes Asian ginseng (*Panax ginseng*) and American ginseng (*Panax quinquefolius*). Asian ginseng is purported to have hepatoprotective effects.[42,43] There are insufficient data on the use of ginseng in persons with HCV to make any recommendations on its use. Ginseng can cause varying degrees of herb-drug interactions.[42] In addition, Asian ginseng may lower blood glucose, so it should be used cautiously in persons with a history of hypoglycemia.[42]

Lactoferrin

Lactoferrin, also known as apolactoferrin or lactotransferrin, is a protein found in milk and other body fluids that binds and transports iron. The highest concentration of lactoferrin is in colostrum. Bovine lactoferrin is the formulation of lactoferrin most often used when taken as a dietary supplement, with a typical dose of 1.8 to 3.6 grams per day; lactoferrin can also be produced via recombinant technology. Several small studies suggest that lactoferrin may lower HCV RNA levels.[44,45,46,47] In contrast, in a placebo-controlled trial among persons with chronic HCV, orally administered bovine lactoferrin at a dose of 1.8 g daily for 12 weeks had no greater impact on HCV RNA levels than placebo.[48] In a randomized placebo-controlled trial of interferon alpha-2b plus ribavirin, with and without lactoferrin, for treatment of chronic HCV, lactoferrin did not increase SVR rates.[49] Although lactoferrin appears to be safe, there is no compelling reason to use it in persons with HCV infection.[50]

Licorice Root (*Glycyrrhiza glabra*)

Licorice root contains a compound called glycyrrhizin (or glycyrrhizic acid). Several preliminary studies suggested intravenous glycyrrhizin had some beneficial effects for persons with HCV.[51,52] The intravenous formulation, however, is not available outside a research setting. Since there are minimal reliable data on oral licorice root for treatment of HCV, no specific recommendations can be made regarding its use. When taken in large amounts, however, licorice root containing glycyrrhizin can cause high blood pressure, salt and water retention, low potassium levels, and alterations in serum cortisol levels.[53,54] Because of the lack of convincing benefit and potential adverse effects, it is not recommended for individuals taking diuretics or

those with cirrhosis or cardiovascular problems. There are no reports of liver injury due to licorice or licorice root.[55]

Red Yeast Rice Extract

Red yeast rice extract is a dietary supplement often used to lower blood cholesterol.[56,57,58] It contains a compound (monacolin K) that is biochemically similar to the HMG-CoA-reductase inhibitor lovastatin. Some red yeast rice extract contains citrinin, a chemical that can cause nephrotoxicity. In addition, red yeast rice extract can potentially cause the same adverse effects that occur with statins, such as acute hepatitis, myopathy, and rhabdomyolysis.[58] For these reasons, we strongly recommend against using red yeast rice extracts in persons with chronic HCV infection.

S-Adenosyl-L-Methionine (SAME)

S-adenosyl-L-methionine (also called S-adenosyl methionine, S-adenosylmethionine, SAME, or SAM-e) is a molecule involved in multiple cellular reactions, acting as the principal methyl donor.[59] Animal models have shown that SAME depletion may favorably impact liver function.[60] In vitro models of HCV demonstrated that SAME enhances the antiviral effect of interferon, although SAME does not have any direct antiviral activity. Persons with HCV who took SAME in conjunction with peginterferon and ribavirin had improved viral kinetics (improved early response), but this did not translate into better sustained virologic response rates.[61,62] Most often, SAME is taken at a dose of 1200 mg per day. There are no large, high-quality studies to date demonstrating a treatment outcome benefit of taking SAME for persons with chronic HCV.[63]

Sho-Saiko-To/Dai-Saiko-To/TJ-9/Xiao-Chai-Hu-Tang

Sho-saiko-to, a mixture of at least seven different herbs, is widely used in parts of Asia, including China and Japan, to treat persons with hepatitis.[64] Sho-saiko-to is often referred to interchangeably as TJ-9 and Dai-saiko-to and Xiao-chai-hu-tang, but these products may have slightly different herbal mixtures.[64] Although the mechanism for Sho-saiko-to remains unclear, there are limited data suggesting it lowers serum aminotransferase levels in persons with chronic hepatitis and may reduce the risk of hepatocellular carcinoma in patients with cirrhosis.[65,66,67] Sho-saiko-to is generally described as having little to no side effects, but there have been several case reports suggesting Sho-saiko-to and Dai-saiko-to can cause acute liver injury (without liver failure).[64,68,69,70] The mechanism of acute liver injury and the responsible ingredients are not well understood, but Skullcap (*Scutellaria lateriflora*), which is often used with Sho-saiko-to, is felt to be a likely culprit.[64] Persons taking Sho-saiko-to or Dai-saiko-to should be counseled on the rare but real risk of acute liver injury with this herbal supplement.

Silymarin

The alternative medication, silymarin, is an extract produced from the seeds of the flowering milk thistle plant (*Silybum marianum*).[71] In vitro and animal studies suggest silymarin and its derivatives protect liver cells from injury and have antiviral activity.[72,73,74] Although oral silymarin is a frequently taken supplement by persons living with chronic HCV infection, clinical studies have not shown a convincing benefit.[72,75,76,77] A randomized study with high-dose oral silymarin found no significant improvement in ALT or decreases in HCV RNA levels.[75] There are data that suggest intravenous silymarin may produce antiviral effects in persons with chronic HCV who receive a liver transplant, but the intravenous form of silymarin is available only in a research setting.[72] Oral milk thistle can be purchased in health food stores without a prescription; the most frequently studied dose is 420 mg per day. The most common side effects with oral silymarin are gastrointestinal (laxative effect, nausea, and epigastric discomfort) and arthralgias.[73] Silymarin may also rarely cause hypoglycemia. Individuals who have allergies to ragweed, chrysanthemum, marigold, or daisy may have a similar reaction to silymarin. Oral preparations of silymarin do not appear to cause hepatotoxicity, but it can decrease bilirubin conjugation and inhibit the cytochrome P450 enzyme system.[78,79] In summary, milk thistle taken orally does not appear to have any beneficial or toxic effects on the liver, and it does not significantly alter HCV RNA levels.

St. John's Wort (*Hypericum perforatum*)

The St. John's wort plant (*Hypericum perforatum*) and its derivatives, hypericin and hyperforin, are commonly used herbal medicines for the treatment of depression.[80] The evidence for its effectiveness in depression is mixed, with several large studies showing no benefit over placebo for major depression.[81,82] Although sometimes taken by persons living with chronic HCV infection, St. John's wort does not have anti-HCV activity. In a small phase 1 study that examined the safety and efficacy of oral St. John's wort in persons with chronic HCV, investigators reported no detectable antiviral activity, but significant problems with phototoxicity.[83] St. John's wort is a strong CYP3A inducer and thus can significantly lower the levels of medications that are substrates of CYP3A.[82,84,85] This effect on CYP3A can potentially impact all recommended DAA regimens used to treat HCV. Thus, it is very important that persons receiving HCV DAA therapy avoid concomitantly taking St. John's wort. Other significant drug interactions with St. John's wort include several cardiovascular drugs, anticoagulants, immunosuppressants, lipid-lowering agents, oral contraceptives, and anti-epileptics.[85] Due to the lack of antiviral activity, potential adverse effects, and major problematic drug interactions, persons with chronic HCV infection should not take St. John's wort.

Thymus Extract

Thymus extract is produced from the thymus gland of cows and contains lymphocytopoietic factors, referred to as thymosins, some of which have pleiotropic immunomodulatory effects, including augmentation of T-cell activity.[86] One member in the family of thymosins, thymosin alpha-1, is a 28-amino acid peptide that stimulates T-cell maturation, antigen recognition, and stimulation of native interferons.[87] Studies involving thymosin alpha-1 in persons with chronic HCV have primarily focused on its use as an adjunctive immune modulatory with interferon-based therapy.[87,88,89] Overall, these studies suggested that thymosin alpha-1 might improve SVR rates in interferon-based therapy, but thymosin alpha-1 was never recommended as part of the standard of care for HCV treatment.[90] Since current HCV therapy has moved completely away from interferon-based therapy, there is no longer significant need or interest in using thymosin alpha-1 to augment HCV treatment. Further, some have raised concerns of contamination of thymus extract products as well as the potential for zoonotic disease transmission, given its bovine source.

Coffee, Diet, and Sodium Intake

Coffee

Coffee consumption may provide a benefit to persons with chronic HCV infection by slowing fibrosis progression and decreasing the risk of developing hepatocellular carcinoma.[[91,92,93,94,95](#)] A meta-analysis of 16 studies found that coffee consumption was associated with a decreased risk of advanced hepatic fibrosis (OR 0.73 [0.58-0.92]) and cirrhosis (OR 0.61 [0.45-0.84]).[[96](#)] Similarly, a large, NIH-sponsored study (referred to as the HALT-C Trial) found that individuals who consumed 3 or more cups of coffee per day were half as likely to have progression of their liver disease and twice as likely to respond to peginterferon and ribavirin therapy.[[97](#)] Despite these findings suggesting an improved response to peginterferon and ribavirin among coffee drinkers, similar data do not exist in the DAA era.

Recommendation

- Persons with chronic HCV infection should be advised that moderate to high coffee consumption may have a beneficial effect on their liver.

Diet

In general, persons with HCV should eat a well-balanced diet that emphasizes fresh fruits, vegetables and protein. A well-balanced diet can help persons receive appropriate amounts of all the vitamins, minerals, and other nutrients their body requires. All persons receiving HCV therapy should drink an adequate amount of water (at least 2 liters per day) to prevent dehydration. It is important to emphasize that protein restriction has not been shown to reduce the risk of hepatic encephalopathy in persons with advanced liver disease and protein restriction has been associated with increased mortality.[[98,99,100](#)] Individuals living with chronic HCV, especially those with cirrhosis, should consume enough protein (1.2 to 1.5 g/kg/day) to avoid muscle wasting and to promote tissue healing.[[101](#)] In particular, persons with cirrhosis are often malnourished and should be encouraged to optimize their protein intake. Examples of high-quality protein include chicken, fish, lean beef, pork, tofu, nuts, beans, milk, yogurt, and eggs.

Recommendations

- Persons living with chronic HCV infection should have a balanced diet and choose nutritious foods from each major food group.
- Individuals with cirrhosis should not have protein restriction; a protein intake of approximately 1.2 to 1.5 g/kg/day is recommended.
- For persons with cirrhosis and hepatic encephalopathy, major guidelines recommend a protein intake of 1.2 to 1.5 g/kg/day.[[102](#)]

Sodium Intake

Individuals with cirrhosis and ascites should limit sodium intake to less than 2,000 mg per day (88 mEq per day) because excessive sodium intake can lead to fluid retention in the form of lower extremity edema and ascites.[[14,101](#)] As a rough guide, one teaspoon of table salt contains about 2,000 mg of sodium. Persons with cirrhosis should not add salt to their food, but instead should replace salt with herbs or spices that can add flavor to the food, without the sodium content. Advise persons with cirrhosis to choose fresh, unprocessed foods instead of salted, smoked, cured, canned, or dried meats, since these processed meats often contain large amounts of sodium.

Recommendation

- Persons with chronic HCV, cirrhosis, and ascites should limit sodium intake to less than 2,000 mg per

day.

Modifying Obesity

In the United States and globally, obesity is a growing epidemic. Obesity is defined as a body mass index (BMI) of 30 or greater and overweight as a BMI of 25 or greater (see [BMI Calculator](#)).^[103] Concurrent with the obesity epidemic, the prevalence of metabolic dysfunction-associated steatotic liver disease (MASLD) has risen substantially, as has the more severe form of MASLD, known as metabolic dysfunction-associated steatohepatitis (MASH).^[104,105,106,107] Even in the absence of HCV infection, MASH can cause cirrhosis, liver cancer, and end-stage liver disease.^[108] In persons with chronic HCV, hepatic fibrosis is accelerated by steatotic liver disease.^[109] In addition, among persons with chronic HCV and cirrhosis, the presence of hepatic steatosis is an independent risk factor for developing hepatocellular carcinoma.^[110,111,112,113] In several older HCV treatment clinical trials involving interferon-based therapy, obesity, insulin resistance, and hepatic steatosis consistently predicted a poorer response to therapy.^[114,115,116] In contrast, with DAA therapy for HCV, individuals with obesity and hepatic steatosis appear to have responses similar to non-obese persons who receive the same therapy.^[117] There are, however, concerns that persons successfully treated for HCV, but who have continued MASLD or MASH, could develop further liver disease and liver complications.

Recommendations

- Persons with steatosis who undergo HCV treatment and achieve an SVR should have their liver monitored with liver function tests every 6 to 12 months to ascertain ongoing liver inflammation.
- Persons with an elevated BMI should be offered nutritional counseling to help promote weight loss and liver health. A combination of exercise and diet often produces the best long-term results.

Alcohol, Cannabis, and Tobacco

Alcohol

In the United States, excessive alcohol consumption is one of the most common causes of cirrhosis and liver failure. The combination of excessive alcohol use with HCV infection causes accelerated fibrosis progression, thereby significantly increasing the risk of developing cirrhosis and liver complications, including hepatocellular carcinoma.[118,119,120,121,122] Available data show that consumption in excess of 30 grams of alcohol per day is hazardous to liver health.[123,124] Note that one alcoholic beverage typically contains about 14 grams of alcohol. In addition, an exact “safe” level of alcohol consumption has never been clearly established for persons with chronic HCV. The Alcohol Use Disorders Identification Test Consumption questionnaire ([AUDIT-C](#)) is a useful screening test for estimating a person's alcohol consumption.[125] Data on the impact of alcohol on DAA HCV treatment are limited, but a large study from the Veterans Affairs health care system suggested excellent SVR rates with DAA therapy, regardless of alcohol use.[126] Individuals who clear HCV with treatment, but continue to drink excessive amounts of alcohol, will continue to be at risk of long-term hepatic complications related to alcohol consumption.[127]

Recommendations

- Ideally, persons with chronic HCV infection should abstain from alcohol. The highest priority is in persons with a history of excessive alcohol use (including alcohol use associated with legal, employment, or relationship problems); these individuals should abstain completely from alcohol.
- According to United States guidelines, women who have never had an alcohol problem should have no more than one alcoholic drink per day, and men with no history of alcohol problems should have no more than two alcoholic drinks per day.
- Individuals with past or present alcohol use should not be excluded from consideration of HCV treatment. Nevertheless, those with ongoing alcohol use should be encouraged to discontinue alcohol prior to, during, and after therapy for HCV, since continued alcohol use will place them at ongoing risk for long-term hepatic complications and liver-induced mortality. In addition, these individuals should be considered for medical therapy, such as naltrexone, to treat alcohol use disorder.

Cannabis

Data evaluating the impact of cannabis on liver fibrosis are mixed. Several studies in persons with chronic HCV mono-infection have shown that individuals who frequently smoked cannabis had an increased risk of developing cirrhosis, even after controlling for other factors, such as alcohol use and obesity.[128,129,130] A more recent meta-analysis showed that marijuana use in persons with chronic HCV was not associated with increased prevalence of hepatic fibrosis.[131] Several studies involving persons with HIV and HCV coinfection have not shown any adverse effect of cannabis or cannabidiol (CBD) on hepatic fibrosis.[131,132,133] There are insufficient data on the impact of cannabis or cannabidiol (CBD) on HCV treatment outcomes.

Recommendations

- Clinicians should inform all persons with chronic HCV about potential hazards of cannabis, but abstinence is not a requirement for HCV treatment.
- Past or present cannabis use should not exclude a person with chronic HCV from receiving HCV treatment.

Tobacco

The effects of tobacco smoking on the liver are controversial.[134,135,136,137] In a study involving 244 adults with chronic HCV infection, persons who smoke had increased hepatic inflammation when compared with persons who did not smoke, but the two groups had similar rates of hepatic fibrosis.[136] In addition,

other studies involving persons with chronic liver disease have shown increased rates of steatosis and advanced fibrosis, but those studies were retrospective and difficult to control for concurrent alcohol use and obesity.[[135](#),[138](#),[139](#)] One study involving adults with HIV and HCV coinfection did not show any association between smoking and accelerated liver disease.[[134](#)] It is clear, however, that tobacco use significantly increases the risk of hepatocellular carcinoma, irrespective of HCV status.[[140](#),[141](#),[142](#)]

Recommendation

- Clinicians should counsel all tobacco smokers with chronic HCV infection to quit tobacco completely for the following three reasons: (1) smoking may potentially accelerate hepatic fibrosis, (2) smoking clearly increases the risk of developing hepatocellular cancer, and (3) smoking is associated with numerous serious adverse health outcomes that are not related to liver disease.

Summary Points

- Persons with chronic HCV infection without cirrhosis may take up to 2 grams per day of acetaminophen. Individuals with cirrhosis should limit their daily intake of acetaminophen to 1 to 2 grams per day. Those with excessive alcohol intake should not take acetaminophen.
- In general, NSAIDs are safe for persons with chronic HCV, except for those with cirrhosis, in whom they should be avoided.
- Individuals with chronic HCV can take a multivitamin without iron, but excess iron intake in the absence of iron deficiency can promote hepatic injury.
- There are no liver-specific reasons to take vitamin D, but individuals with vitamin D deficiency (commonly defined as serum levels less than 20 ng/mL) should receive vitamin D supplementation.
- No complementary or alternative medications have shown a definite benefit for persons with HCV, and St. John's wort should be avoided in persons receiving treatment for HCV, given its potential to interact with DAA medications.
- Drinking moderate to high amounts of coffee (approximately 3 or more cups per day) may have beneficial effects for the liver.
- A balanced, low-fat (less than 30% of total calories) diet is recommended. Individuals with cirrhosis should limit sodium intake to less than 2 grams per day and consume at least 1.2 to 1.5 grams of protein per kilogram per day.
- Metabolic dysfunction-associated steatotic liver disease (MASLD) can accelerate the progression of fibrosis in patients with chronic HCV and increase the risk for HCC. Persons who are overweight should receive counseling on diet and exercise modifications that can help promote weight loss.
- Ideally, persons with chronic HCV infection should abstain from alcohol for liver health, but persons with past or present alcohol use should not be excluded from consideration of HCV treatment.

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