

## Hepatitis Testing

Policy Number: AHS – G2036 – Hepatitis Testing	Prior Policy Name and Number, as applicable:  AHS-G2036-Hepatitis C
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[POLICY DESCRIPTION](#) | [INDICATIONS AND/OR LIMITATIONS OF COVERAGE](#) | [TABLE OF TERMINOLOGY](#) | [SCIENTIFIC BACKGROUND](#) | [GUIDELINES AND RECOMMENDATIONS](#) | [APPLICABLE STATE AND FEDERAL REGULATIONS](#) | [APPLICABLE CPT/HCPCS PROCEDURE CODES](#) | [EVIDENCE-BASED SCIENTIFIC REFERENCES](#) | [REVISION HISTORY](#)

### I. Policy Description

Infectious hepatitis is an inflammation of the liver caused by the hepatitis viruses. Hepatitis C is a blood-borne virus that can be spread via sharing needles or other equipment to inject drugs as well as in inadequate infection control in healthcare settings (CDC, 2018). Hepatitis C causes liver disease and inflammation. A chronic HCV infection can lead to hepatic damage, including cirrhosis and hepatocellular carcinoma, and is the most common cause of liver transplantation in the United States (AASLD-IDSA, 2015).

Hepatitis B is spread by the “Percutaneous, mucosal, or nonintact skin exposure to infectious blood, semen, and other body fluids.” As the hepatitis B virus is concentrated most highly in blood, “percutaneous exposure is an efficient mode of transmission”, though HBV can also be transmitted through birth to an infected mother and sexual contact with an infected person and less commonly through needle-sticks or other sharp instrument injuries, organ transplantation and dialysis, and interpersonal contact through sharing items, such as razors or toothbrushes or contact with open sores of an infected person. Similar to HCV infection, 15% to 25% of people with chronic HBV infection develop chronic liver disease (CDC, 2020a).

The general route of transmission for the hepatitis A virus is through the fecal-oral route by close person-to-person contact with an infected person, sexual contact with an infected person, or the ingestion of contaminated food or water, with the bloodborne transmission of HAV being uncommon (CDC, 2020a). Though death is uncommon and most people with acute HAV infection recover with no lasting liver damage, HAV remains a worldwide public health issue and is endemic in many low- to middle-income countries (CDC, 2020a; Keles et al., 2021).

For HCV and HBV screening in pregnant individuals, please see AHS-G2035-Prenatal Screening (Nongenetic).

### II. Indications and/or Limitations of Coverage

Application of coverage criteria is dependent upon an individual’s benefit coverage at the time of the request.

- 1) For asymptomatic, non-pregnant individuals, the following annual Hepatitis B virus (HBV) infection screening **MEETS COVERAGE CRITERIA**:
  - a) Hepatitis B surface antigen (HBsAg) when **one** of the following high-risk situations is met:
    - i) For individuals born in geographic regions with a HBsAg prevalence 2% or higher (see Note 1).
    - ii) For U.S.-born individuals not vaccinated as infants whose parents were born in geographic regions with a HBsAg prevalence 8% or higher (see Note 1).
    - iii) For individuals with elevated alanine aminotransferase (ALT)/aspartate transaminase (AST) of unknown etiology.
  - b) HBsAg and hepatitis B surface antibody (anti-HBs) for infants born from an HBsAg-positive individual.
  - c) HBsAg, anti-HBs, and total hepatitis B core antibody (anti-HBc), when **one** of the following high-risk situations is met:
    - i) For injection-drug users.
    - ii) For men who have sex with men.
    - iii) For individuals with select medical conditions who require cytotoxic or immunosuppressive therapy.
    - iv) For individuals who are on long-term hemodialysis treatment.
    - v) For household contacts, needle-sharing contacts, and sex partners of HBV-infected individuals.
    - vi) For individuals infected with HIV.
    - vii) For healthcare and public safety workers exposed to blood or body fluids.
- 2) For individuals who test positive for HBV by initial antibody screening, follow up serologic testing (anti-HBc, anti-HBs, IgM antibody to anti-HBc [IgM anti-HBc]) **MEETS COVERAGE CRITERIA**.
- 3) For the confirmation of seroconversion after hepatitis B vaccination, anti-HBs testing **MEETS COVERAGE CRITERIA**.
- 4) For individuals who test positive for HBV by initial antibody screening and who will undergo immunosuppressive drug therapy, HBV DNA testing **MEETS COVERAGE CRITERIA**.
- 5) For all individuals 18 years of age and older, antibody testing for Hepatitis C (HCV) infection once per lifetime **MEETS COVERAGE CRITERIA**.
- 6) For any individual with the following recognized conditions or exposures, one-time, post-exposure antibody testing for HCV infection **MEETS COVERAGE CRITERIA**:
  - a) For individuals who have used illicit intranasal or injectable drugs.
  - b) For individuals who have received clotting factor concentrates produced before 1987.

- c) For individuals with a history of hemodialysis.
  - d) For individuals with evidence of liver disease (based on clinical presentation, persistently abnormal ALT levels, or abnormal liver function studies).
  - e) For individuals infected with HIV.
  - f) For individuals who received an organ transplant before July 1992.
  - g) For individuals who received a blood transfusion or blood component before July 1992.
  - h) For individuals notified that they received blood from a donor who later tested positive for an HCV infection.
  - i) For individuals with a history of incarceration.
  - j) For individuals who received a tattoo in an unregulated setting.
  - k) For healthcare, emergency medical, and public safety workers after needle sticks, sharps, or mucosal exposures to HCV-positive blood.
  - l) For children born from an HCV-positive individual.
  - m) For current sexual partners of HCV-infected persons.
- 7) Routine periodic antibody testing for HCV **MEETS COVERAGE CRITERIA** for individuals with **any** of the following ongoing risk factors (while risk factors persist):
- a) For individuals who currently inject drugs and share needles, syringes, or other drug preparation equipment.
  - b) For individuals who are receiving ongoing hemodialysis.
  - c) For individuals engaging in high-risk sexual behavior.
- 8) For individuals who test positive for HCV by initial antibody screening, follow up nucleic acid testing for HCV (to differentiate between active infection and resolved infection) **MEETS COVERAGE CRITERIA.**
- 9) Prior to the initiation of direct acting anti-viral (DAA) treatment, one time testing for HCV genotype to guide selection of the most appropriate antiviral regimen **MEETS COVERAGE CRITERIA.**
- 10) Testing for HCV viral load with a quantitative nucleic acid test **MEETS COVERAGE CRITERIA** in **any** of the following situations:
- a) Prior to the initiation of DAA therapy.
  - b) After four weeks of DAA therapy.
  - c) At the end of treatment.
  - d) Twelve, twenty-four, and forty-eight weeks after completion of treatment.
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**NOTES:**

**Note 1:** The CDC defines HBsAg prevalence by geographic region:

<https://wwwnc.cdc.gov/travel/yellowbook/2020/travel-related-infectious-diseases/hepatitis-b>.

### III. Table of Terminology

Term	Definition
AASLD	American Association for the Study of Liver Diseases
Ab	Antibody
AFP	Alpha fetoprotein
Ag	Antigen
AGA	American Gastroenterological Association
ALT	Alanine aminotransferase
anti-HBc	Total hepatitis B core antibody
anti-HBs	Hepatitis B surface antibody
aPTT	Activate partial thromboplastin time
AST	Aspartate aminotransferase
CBC	Complete blood count
CDC	Centers for Disease Control and Prevention
CHC	Chronic hepatitis C
CIA	Chemiluminescence immunoassays
CLIA '88	Clinical Laboratory Improvement Amendments of 1988
CMS	Centers for Medicare and Medicaid
DAA	Direct acting anti-viral
DBS	Dried blood spot
EASL	European Association for the Study of the Liver
EIA	Enzyme immunoassays
ELISA	Enzyme-linked immunosorbent assay
FBC	Full blood count
FDA	Food and Drug Administration
HAV	Hepatitis A virus
HBsAg	Hepatitis B surface antigen
HBV	Hepatitis B virus
HCR	Hepatitis C Screening
HCV	Hepatitis C virus
HIV	Human immunodeficiency virus
ICP	Intrahepatic cholestasis of pregnancy
IDSA	Infectious Disease Society of America
IHS	Indian Health Services
INR	International normalized ratio
LDH	Lactic acid dehydrogenase

Term	Definition
LDTs	Laboratory-developed tests
MTCT	Mother-to-child transmission
NAAT	Nucleic acid amplification test
NAT	Nucleic acid test
NCD	National coverage determinations
PALF	Pediatric acute liver failure
PCR	Polymerase chain reaction
POC	Point of care
PrEP	Pre-exposure prophylaxis
PT	Prothrombin time
PWID	People who inject drugs
Rdts	Rapid diagnostic tests
RNA	Ribonucleic acid
SVR	Sustained virologic response
T2DM	Type 2 diabetes mellitus
TB	Tuberculosis
USPSTF	United States Preventive Services Task Force
WHO	World Health Organization

## IV. Scientific Background

### *Hepatitis C*

The Centers for Disease Control and Prevention estimate that 2.4 million people in the United States have chronic hepatitis C (CDC, 2020c). Prevalence of the infection is highest in individuals born between 1945 and 1965. This rate is approximately six times higher than that seen in other adult age groups, and the CDC estimated approximately 50,300 new infections occurring each year (CDC, 2018). Hepatitis C virus (HCV) infection is the most common reason for liver transplantation in adults in the U.S. and may lead to hepatocellular carcinoma (Chopra, 2021).

It is estimated that 20% of people with HCV infection will develop cirrhosis, and nearly 5% will die from liver disease resulting from the HCV infection. The number of deaths from hepatitis is increasing and is projected to continue to increase for several more decades unless treatment is scaled up considerably (Razavi et al., 2014). Although HCV infection is common, it is estimated that 50-75% of individuals who are infected are unaware of their infection as symptoms are absent or nonspecific until much later, and therefore do not receive the care and treatment that can mitigate progression to severe liver disease and possibly death (Hagan et al., 2006; Rein et al., 2012).

HCV is spread through exposure to blood of infected individuals. Such exposure includes injection drug use, blood transfusions (prior to 1992), and to a lesser extent, high-risk sexual behaviors. Additionally, being born to an HCV-infected mother, hemodialysis, intranasal drug use, tattoos, incarceration, needle sticks, and invasive procedures (prior to implementation of universal precautions) are also associated with increased risk of HCV infection. Some countries

are experiencing a recent resurgence of HCV infection among young intravenous drug users and HIV-infected homosexual men (CDC, 2015; Wandeler et al., 2015).

HCV is a small, positive-stranded RNA-enveloped virus with a highly variable genome (Simmonds, 2001). Assessment of the HCV genotype is crucial for management of the HCV infection. There are currently six major genotypes of HCV, and major treatment decisions (regimen, dosing, duration) vary from genotype to genotype (Chopra & Arora, 2020). Some regimens for one genotype (such as ledipasvir-sofosbuvir [“Harvoni”] for genotype 1) may not be effective for another (in this case, Harvoni may be used for genotypes 1, 4, 5, and 6 but not 2 or 3) (Chopra & Muir, 2022; Lexicomp, 2022).

HCV is frequently asymptomatic, necessitating the need of strong screening procedures. As many as 50% of HCV-infected individuals are unaware of their diagnosis, and risk factors such as drug use or blood transfusions may increase risk of acquiring an HCV infection. Several expert groups, such as the CDC, have delineated screening recommendations in order to provide better care against the virus (Chopra & Arora, 2022).

Hepatitis C can be diagnosed with either serologic antibody assays or molecular RNA tests. A serologic assay can detect an active infection and a resolved HCV infection, but cannot differentiate whether the infection is acute, chronic, or no longer present. Various serologic assays include enzyme immunoassays (EIA), chemiluminescence immunoassays (CIA), and point-of-care rapid immunoassays (Spach, 2020).

Molecular RNA tests detect Hepatitis C RNA, and the process includes nucleic acid test (NAT) or nucleic acid amplification test (NAAT). The NAT test becomes positive 1-2 weeks after initial infection and it has become the gold standard test for patients who have a positive EIA screening test. The NAT can detect whether a patient has a current active infection or a resolved infection (Spach, 2020).

### *Hepatitis B*

The hepatitis B virus (HBV) is a double-stranded DNA virus belonging to the hepadnavirus family. The diagnosis of its acute infection is characterized by the detection of hepatitis B surface antigen (HBsAg) and immunoglobulin M (IgM) antibody to hepatitis B core antigen (anti-HBc), and chronic conditions develop in 90% of infants after acute infection at birth, 25%–50% of children newly infected at ages 1–5 years, and 5% of people newly infected as adults (CDC, 2020a; Lok, 2022).

HBV is transmitted from infected patients to those who are not immune (i.e., hepatitis B surface antibody [anti-HBs]-negative). Methods of transmission include mother-to-child (whether in utero, at birth, or after birth), breastfeeding, paternal transmission (i.e., close contact with infected blood or fluid of fathers), transfusion, sexual transmission, nosocomial infection, percutaneous inoculation, transplantation, and blood exposure via minor breaks in skin or mucous membranes (Teo & Lok, 2022).

In the United States, an estimated 862,000 people were living with chronic hepatitis B infection in 2016, with 21,600 new infections in 2018. Though most people with acute disease recover with no lasting liver damage, 15% to 25% of those with chronic disease develop chronic liver

disease, including cirrhosis, liver failure, or liver cancer. It is believed that there are more than 250 million HBV carriers in the world, 600,000 of whom die annually from HBV-related liver diseases. As many as 60% of HBV-infected persons are unaware of their infection, and many remain asymptomatic until the presentation of cirrhosis or late-stage liver disease (CDC, 2020a; Lok, 2022; US Preventive Services Task Force, 2020).

The initial evaluation of chronic HBV infection should include a history and physical examination focusing on “risk factors for coinfection with hepatitis C virus (HCV), hepatitis delta virus (HDV), and/or HIV; use of alcohol; family history of HBV infection and hepatocellular carcinoma (HCC); and signs and symptoms of cirrhosis.” Furthermore, it should employ laboratory tests, such as “a complete blood count with platelets, liver chemistry tests (aspartate aminotransferase [AST], alanine aminotransferase [ALT], total bilirubin, alkaline phosphatase, albumin), international normalized ratio (INR), and tests for HBV replication (HBeAg, antibody to HBeAg [anti-HBe], HBV DNA”, and testing for hepatitis A virus (HAV) immunity with HAV immunoglobulin G (IgG) antibody in those who are not immune. Other considerations include evaluation for other causes of liver disease, screening for HIV infection, screening for hepatocellular carcinoma (HCC), screening for fibrosis, and, in rare cases, a liver biopsy (Lok, 2022).

### *Hepatitis A*

Hepatitis A infection is caused by the hepatitis A virus, of which humans are the only known reservoir. The HAV virus is member of the genus *Hepatovirus* in the family Picornaviridae, and other previously used names for HAV infection include epidemic jaundice, acute catarrhal jaundice, and campaign jaundice (Lai & Chopra, 2022).

The hepatitis A virus is generally transmitted through the fecal-oral route, either via person-to-person contact (e.g., transmission within households, within residential institutions, within daycare centers, among military personnel, or sexually) or consumption of contaminated food or water (consumption of undercooked foods or foods infected by food handlers). Additional modes of transmission include blood transfusion and illicit drug use, and it should be noted that maternal-fetal transmission has not yet been described (Lai & Chopra, 2022).

Globally, approximately 1.4 million new cases of HAV infection occur each year—in the United States alone, an estimated 24,900 new infections were detected in 2018. Acute infection by HAV is usually a self-limited disease, with fulminant manifestations of hepatic failure occurring in fewer than 1 percent of cases. However, symptomatic illness due to HAV still presents itself in seventy percent of adults. Consequently, “diagnosis of acute HAV infection should be suspected in patients with abrupt onset of prodromal symptoms (nausea, anorexia, fever, malaise, or abdominal pain) and jaundice or elevated serum aminotransferase levels, particularly in the setting of known risk factors for hepatitis A transmission” through detection of serum IgM anti-HAV antibodies due to its persistence throughout the duration of the disease (CDC, 2020a; Lai & Chopra, 2022).

### ***Proprietary Testing***

Many point-of-care tests have been developed to diagnose hepatitis C efficiently. These point-of-care tests are particularly important for diagnoses in economically impoverished areas.

Examples of these tests include OraQuick, TriDot and SDBioline. The OraQuick HCV test is an FDA approved point-of-care test which utilizes a fingerstick and a small whole blood sample to detect the virus. This test is reportedly more than 98% accurate and provides results in 20 minutes (OraSure, 2013). The 4<sup>th</sup> Generation HCV Tri-Dot is a rapid test which can detect all subtypes of HCV with 100% sensitivity and 98.9% specificity (JMitra&Co, 2015). This test uses human serum or plasma and can provide results in three minutes. Finally, the SDBioline is an immunochromatographic rapid test that can identify HCV antibodies in human serum, plasma, or whole blood (Abbott, 2023). This test uses a safe fingerstick procedure to obtain a sample.

Hepatitis panel tests have also been developed. For example, the VIDAS® Hepatitis panel by BioMérieux tests for hepatitis A, B and C in less than two hours (BioMérieux, 2018). This panel includes nine automated assays and is a rapid, reliable and simple testing method. Legacy Health's Hepatitis Chronic Panel detects Hepatitis B and C within 24-48 hours through a CIA method (Legacy\_Health, 2021).

A hepatitis C vaccine is currently not available although many vaccines are under development; barriers to the development of such a vaccine include virus diversity, a lack of knowledge of the immune responses when an infection occurs, and limited models for the testing of new vaccines (Ansaldi et al., 2014; Bailey et al., 2019). The World Health Organization hopes for a 90% reduction in new hepatitis C cases by the year 2030 (Bailey et al., 2019).

Management of HCV infection typically involves monitoring the effect of treatment. The goal of treatment is to achieve a “sustained virologic response” (SVR), which is defined as “an undetectable RNA level 12 weeks following the completion of therapy” (Chopra & Pockros, 2020). This measure is a proxy for elimination of HCV RNA. The assessment schedule may vary regimen to regimen, but the viral load is generally evaluated every few weeks (Chopra & Pockros, 2020).

### ***Clinical Utility and Validity***

In order to determine the link between hepatitis A infection and its rare complication of acute liver failure in children in Somalia, a retrospective study was conducted on children aged 0 to 18 who were admitted to the pediatric outpatient clinic and pediatric emergency departments of the Somalia Mogadishu-Turkey Training and Research Hospital, Somali, from June 2019 and December 2019, and who were tested for HAV and had complete study data available (Keles et al., 2021). The authors found that of the 219 hepatitis A cases analyzed, 25 (11%) were diagnosed with pediatric acute liver failure (PALF) while the remaining 194 were not. It was found that children with PALF had “significantly had more prolonged PT and aPPT, and higher INR values in coagulation assays; and had higher levels of albumin in biochemical tests than the group without liver failure (for all,  $p \leq 0.05$ )”, though no other significant differences were found based on the other laboratory parameters tested. Moreover, “Hepatic encephalopathy was observed in individuals with hepatitis A disease (12/219; 15.4%), in which PALF positive group (5/25;40%) was significantly higher compared to the non-PALF group (7/194; 4%) ( $p = < 0,001$ ). The length of stay in the hospital or intensive care unit was significantly higher in children with acute liver failure ( $p = 0.001$ ) .” As such, Keles et al. (2021) astutely notes that though “death rates of Hepatitis A infection seem to be low”, HAV infection may potentially “require long-term hospitalization of patients due to the complication of acute liver failure, which causes loss of



workforce, constitutes a socio-economic burden on individuals and healthcare systems, and leads to mortality in settings where referral pediatric liver transplantation centers are not available .”

Spenatto et al. (2013) screened 6194 asymptomatic patients who were requesting an STI screening for hepatitis B infection. The authors found that only “male gender, lack of employment, and birth, in medium or high endemic country, were independently associated with HBsAg positivity in multivariate analysis”, and neither sexual behavior nor vaccination status are needed to target high-risk populations (Spenatto et al., 2013).

Su et al. (2022) evaluated the cost-effectiveness of implementing universal HBV screening in China to identify optimal screening strategies. By using a Markov cohort model, the researchers “simulated universal screening scenarios in 15 adult age groups between 18 and 70 years, with different years of screening implementation (2021, 2026, and 2031) and compared to the status quo (ie, no universal screening)”, investigating a total of 180 different scenarios. Their work found suggested that “with a willingness-to-pay level of three times the Chinese gross domestic product (GDP) per capita (US\$30 828), all universal screening scenarios in 2021 were cost-effective compared with the status quo”, with the “serum HBsAg/HBsAb/HBeAg/HBeAb/HBcAb (five-test) screening strategy in people aged 18-70 years was the most cost-effective strategy in 2021” and “the two-test strategy for people aged 18-70 years became more cost-effective at lower willingness-to-pay levels .” Most importantly, they claimed that the “five-test strategy could prevent 3.46 million liver-related deaths in China over the lifetime of the cohort” and that delaying strategic intervention will reduce overall cost-effectiveness (Su et al., 2022).

Messina et al. (2015) performed a meta-analysis on the prevalence of HCV genotypes worldwide. The authors evaluated 1217 studies encompassing approximately 90% of the global population. They calculated genotype 1 to comprise 83.4 million cases (46.2% of all HCV cases), genotype 3 to comprise 54.3 million cases (30.1%), and genotypes 2, 4, and 6 to comprise a combined 22.8% cases. Genotype 5 comprised less than 1% of HCV cases. The diversity of genotypes also varied; the highest diversity is observed in China and South-East Asia, while in some countries, such as Egypt and Mongolia, almost all HCV infections are caused by a single genotype (Messina et al., 2015).

Inoue et al. (2017) described four HCV patients whose treatment failed. These four HCV patients had received a treatment regimen of daclatasvir plus asunaprevir, which is used for genotype 1b. However, these four patients were re-tested and found to have a different genotype; 3 patients had genotype 2 and the 4<sup>th</sup> patient had genotype 1a. The authors suggested that the daclatasvir plus asunaprevir regimen was ineffective for patients without genotype 1b (Inoue et al., 2017).

Moreno et al. (2016) performed a cost analysis of expanded HCV coverage. Two scenarios were simulated, one with expanded fibrosis coverage to stage 2 fibrosis, and the other to all fibrosis cases. Over a 20-year simulation, treatment costs increased, but private payers experienced overall savings of \$10 billion to \$14 billion after treatment costs. A positive “spillover” benefit of \$400 million to Medicare was seen in the 5-year model, and a benefit of \$7 billion to Medicare was seen in the 20-year model (Moreno et al., 2016).

Linthicum et al. (2016) assessed the cost-effectiveness of expanding screening and treatment coverage over a 20-year horizon. The authors investigated three scenarios, each of which

expanded coverage to a different stage of fibrosis. “Net social value” was the primary outcome evaluated, and it was calculated by the “value of benefits from improved quality-adjusted survival and reduced transmission minus screening, treatment, and medical costs.” Overall, the scenario with only fibrosis stage 3 and fibrosis stage 4 covered generated \$0.68 billion in social value, but the scenario with all fibrosis patients (stages 0-4) treated produced \$824 billion in social value. The authors also noted that the scenario with all fibrosis stages covered created net social value by year 9 whereas the scenario with only stages 3 and 4 covered needed all 20 years to break even (Linthicum et al., 2016).

Chen et al. (2019) completed a meta-analysis to research the relationship between type 2 diabetes mellitus development and patients with a HCV infection. Studies were included from 2010 to 2019. Five types of HCV individuals were incorporated in this study including those who were “non-HCV controls, HCV-cleared patients, chronic HCV patients without cirrhosis, patients with HCV cirrhosis and patients with decompensated HCV cirrhosis” (Chen et al., 2019). HCV infection was found to be a significant risk factor for type 2 diabetes mellitus development. Further, “HCV clearance spontaneously or through clinical treatment may immediately reduce the risk of the onset and development of T2DM [type 2 diabetes mellitus] (Chen et al., 2019).”

Saeed et al. (2020) completed a systematic review and meta-analysis of health utilities for patients diagnosed with a chronic hepatitis C infection. Health utility can be defined as a measure of health-related quality or general health status. A total of 51 studies comprised of 15,053 patients were included in this study. The researchers have found that “Patients receiving interferon-based treatment had lower utilities than those on interferon-free treatment (0.647 vs 0.733). Patients who achieved sustained virologic response (0.786) had higher utilities than those with mild to moderate CHC [chronic hepatitis C]. Utilities were substantially higher for patients in experimental studies compared to observational studies (Saeed et al., 2020).” Overall, these results show that chronic hepatitis C infections are significantly harming global health status based on the measurements provided by health utility instruments.

Vetter et al. (2020) conducted a retrospective study to assess the performance of rapid diagnostic tests (RDTs) for Hepatitis C virus (HCV) infection. 13 RDTs were studied including the Standard Q HCV Ab by SD Biosensor, HCV Hepatitis Virus Antibody Test by Antron Laboratories, HCV-Ab Rapid Test by Beijing Wantal Biological Pharmacy Enterprise, Rapid Anti-HCV Test by InTec, First Response HCV Card Test by Premier Medical Corporation, Signal HCV Version 3.0 by Arkray Healthcare, TRI DOT HCV by J. Mitra & Co, Modified HCV-only Ab Test by Biosynex SA, SD Bioline HCV by Abbott Diagnostics, OraQuick HCV by OraSure, Prototype HCV Ab Test by BioLytical Laboratories, Prototype DPP HCV by Chembio Diagnostic Systems, and Prototype Care Start HCV by Access Bio. 1,710 samples were evaluated in which 648 samples were HCV positive and 264 samples were also HIV positive. In the samples from HIV negative patients, most RDTs showed high sensitivity of > 98% and specificity of >99%. In HIV positive patients, sensitivity was lower with only 1 RDT reaching >95%. However, specificity was higher, with only 4 RDTs showing a specificity of <97%. The authors concluded that these tests are compliant with the World Health Organization (WHO) guidance which recommends an HCV RDT to have a sensitivity of >98% and specificity >97%. However, in HIV positive patients, the specificity remained high, but none of the tests met the WHO sensitivity criteria. The authors conclude that “these findings serve as a valuable baseline to investigate RDT

performance in prospectively collected whole blood samples in the intended use settings (Vetter et al., 2020)."

In a prospective study, Chevaliez et al. (2020) evaluated the use of molecular point of care (POC) testing and dried blood spot (DBS) for HCV screening in people who inject drugs (PWID). 89 HCV-seropositive PWID were further assessed with a liver assessment, blood tests, POC HCV RNA testing, and fingerstick DBS sampling. 77 patients had paired fingerstick capillary whole blood for POC HCV RNA testing and fingerstick sampling with interpretable results, while the other 12 samples had no valid result due to low sample volume. The POC HCV RNA test detected 30 HCV-seropositive PWID and DBS sampling detected 27 HCV-seropositive PWID. The rate of invalid results using the POC test was below 10%, so it may be performed by staff without extensive clinical training in decentralizing testing location. This study also showed high concordance for detection of active HCV infection from DBS compared to the POC test. The authors conclude that the use of POC diagnostic testing and DBS sampling should be recommended as a one-step screening strategy to increase diagnosis, increase treatment, and reduce the number of visits.

In an Australian observational study, Catlett et al. (2021) evaluated the Aptima HCV Quant Dx Assay to see how well it could detect HCV RNA from fingerstick capillary dried blood spot (DBS) and venipuncture-collected samples. DBS collection would benefit marginalized populations in areas that may not have access to phlebotomy services or who may have difficult venous access. DBS has also been shown to “enhance HCV testing and linkage to care,” be easy for transport and storage, and can be used for other purposes like HCV sequencing and testing for HIV or hepatitis B simultaneously, which is useful in more resource-limited settings. From 164 participants, they found HCV RNA in 45 patients. The Aptima assay rendered a sensitivity and specificity of 100% from plasma, and a sensitivity of 95.6% and specificity of 94.1% from DBS. This demonstrated the comparable diagnostic performance of this assay when it comes to detecting active HCV infection from DBS samples and plasma samples, and hopefully the eventual use of other similar assays with similar performances.

## V. Guidelines and Recommendations

### Centers for Disease Control and Prevention (CDC)

#### *Hepatitis C*

The CDC recommends universal hepatitis C screening for

- “Hepatitis C screening at least once in a lifetime for all adults aged 18 years and older, except in settings where the prevalence of HCV infection (HCV RNA-positivity) is less than 0.1%”
- “Hepatitis C screening for all pregnant women during each pregnancy, except in settings where the prevalence of HCV infection (HCV RNA-positivity) is less than 0.1%”

Moreover, one-time hepatitis C testing regardless of age or setting prevalence among people with recognized conditions or exposures is recommended for the following groups:

- “People with HIV

- “People who ever injected drugs and shared needles, syringes, or other drug preparation equipment, including those who injected once or a few times many years ago”
- “People with selected medical conditions, including:
  - people who ever received maintenance hemodialysis
  - people with persistently abnormal ALT levels”
- “Prior recipients of transfusions or organ transplants, including:
  - people who received clotting factor concentrates produced before 1987
  - people who received a transfusion of blood or blood components before July 1992
  - people who received an organ transplant before July 1992
  - people who were notified that they received blood from a donor who later tested positive for HCV infection.”
- “Health care, emergency medical, and public safety personnel after needle sticks, sharps, or mucosal exposure to HCV-positive blood”
- “Children born to mothers with HCV infection.”

“Routine periodic testing is recommended for people with ongoing risk factors, while risk factors persist:

- People who currently inject drugs and share needles, syringes, or other drug preparation equipment
- People with selected medical conditions, including:
  - people who ever received maintenance hemodialysis.”

CDC also recommends that “Any person who requests hepatitis C testing should receive it, regardless of disclosure of risk, because many persons may be reluctant to disclose stigmatizing risks” (CDC, 2020e).

The CDC also notes that the initial HCV test should be “with an FDA-approved test for antibody to HCV.” A positive result for the HCV antibody indicates either a current infection or previous infection that has resolved. For those individuals, the CDC recommends testing by an FDA-approved HCV nucleic acid test (NAT) to differentiate between active infection and resolved infection. For the identification of chronic hepatitis C virus infection among persons born between 1945 and 1965, the CDC states that “Persons who test anti-HCV positive or have indeterminate antibody test results who are also positive by HCV NAT should be considered to have active HCV infection; these persons need referral for further medical evaluation and care.” Finally, the CDC also recommends that repeat testing should be considered for individuals with ongoing risk behaviors (CDC, 2012).

The CDC published guidance for healthcare personnel with potential exposure to HCV. CDC recommends testing the source patient and the healthcare personnel. When testing the source patient, baseline testing should be performed within 48 hours after exposure by testing for HCV RNA or HCV antibodies. All HCV RNA testing should be performed with a nucleic acid test. If the source patient was HCV RNA positive or if source patient testing was not performed, baseline testing for healthcare personnel should follow the same steps through nucleic acid testing 3-6 weeks post-exposure. A final HCV antibody test should be performed at 4-6 months post-exposure to ensure a negative HCV RNA test result (CDC, 2020d).

No serologic marker for acute infection is available, but for chronic infections, CDC propounds the use of “Assay for anti-HCV” and “Qualitative and quantitative nucleic acid tests (NAT) to detect and quantify presence of virus (HCV RNA)” (CDC, 2020a).

### *Hepatitis B*

Regarding routine testing and follow-up for chronic hepatitis B virus, CDC organized their recommendations in the following table:

Population	Recommendation	
	Testing	Vaccination/Follow-up
Persons born in regions of high and intermediate HBV endemicity (HBsAg prevalence $\geq 2\%$ )	Test for HBsAg, regardless of vaccination status in their country of origin, including <ul style="list-style-type: none"> <li>– immigrants</li> <li>– refugees</li> <li>– asylum seekers</li> <li>– internationally adopted children</li> </ul>	If HBsAg-positive, refer for medical management. If negative, vaccinate. Susceptible persons should complete a hepatitis B vaccine series.
U.S.-born persons not vaccinated as infants whose parents were born in regions with high HBV endemicity ( $\geq 8\%$ ). <a href="#">Learn more about prevalence of HBV infection around the world.</a>	Test for HBsAg regardless of maternal HBsAg status if not vaccinated as infants in the United States.	If HBsAg-positive, refer for medical management. If negative, vaccinate. Susceptible persons should complete a hepatitis B vaccine series.
Persons who inject drugs	Test for HBsAg, as well as for anti-HBc or anti-HBs to identify susceptible persons.	If HBsAg-positive, refer for medical management. If negative, vaccinate. Susceptible persons should complete a hepatitis B vaccine series to prevent infection from ongoing exposure. First vaccine dose should be given at the same visit as testing.
Men who have sex with men	Test for HBsAg, as well as for anti-HBc or anti-HBs to identify susceptible persons.	If HBsAg-positive, refer for medical management. If negative, vaccinate. Susceptible persons should complete a hepatitis B vaccine series to prevent infection from ongoing exposure. First vaccine dose should be given at the same visit as testing.

Population	Recommendation	
	Testing	Vaccination/Follow-up
Persons needing immunosuppressive therapy, including chemotherapy, immunosuppression related to organ transplantation, and immunosuppression for rheumatologic or gastroenterologic disorders	Test for all markers of HBV infection (HBsAg, anti-HBc, and anti-HBs).	Treat persons who test positive for HBsAg. Monitor closely persons who test positive for anti-HBc for signs of liver disease. If negative, vaccinate. Susceptible persons should complete a hepatitis B vaccine series.
Persons with elevated ALT/AST of unknown etiology	Test for HBsAg along with other appropriate medical evaluation.	Follow-up as indicated.
Donors of blood, plasma, organs, tissues, or semen	Test for HBsAg, anti-HBc, and HBV-DNA as required.	
Persons on maintenance dialysis, including in-center or home hemodialysis and peritoneal dialysis, and persons who are predialysis	Test for all markers of HBV infection (HBsAg, anti-HBc, and anti-HBs). Test vaccine nonresponders monthly for HBsAg. Patients on hemodialysis who test positive for HBsAg should be cohorted.	Vaccinate against hepatitis B to prevent transmission and revaccinate when serum anti-HBs titer falls below 10mIU/mL.
All pregnant persons	Test for HBsAg during each pregnancy, preferably in the first trimester. Test at the time of admission for delivery if prenatal HBsAg test result is not available or if person was at risk for infection during pregnancy.	If HBsAg-positive, refer for medical management. To prevent perinatal transmission, infants of persons who test positive for HBsAg and persons with unknown HBsAg status should receive vaccination and postexposure immunoprophylaxis in accordance with recommendations and within 12 hours of delivery.
Infants born to persons who test positive for HBsAg	Test for HBsAg and anti-HBs 1–2 mos after completion of at least 3 doses of a licensed hepatitis B vaccine series (i.e., at age 9–18 months, generally at the next well-child visit to assess effectiveness of postexposure immunoprophylaxis). Testing should not be performed	Vaccinate in accordance <a href="#">with recommendations</a> .

Population	Recommendation	
	Testing	Vaccination/Follow-up
	before age 9 months or within 1 month of the most recent vaccine dose.	
Household, needle-sharing, or sex contacts of persons known to have tested positive for HBsAg	Test for HBsAg, as well as anti-HBc or anti-HBs to identify susceptible persons.	First vaccine dose should be given at the same visit as testing. Susceptible persons should complete a hepatitis B vaccine series to prevent transmission from ongoing exposure.
Persons who are the sources of blood or body fluids resulting in an exposure (e.g., needlestick, sexual assault) that might require postexposure prophylaxis	Test source for HBsAg.	Vaccinate health care and public safety workers with reasonably anticipated occupational exposures to blood or infectious body fluids. Provide postexposure prophylaxis to exposed person if needed.
Persons with HIV infection	Test for HBsAg, as well as for anti-HBc or anti-HBs to identify susceptible persons.	Vaccinate susceptible persons against hepatitis B to prevent transmission.

(CDC, 2022).

Serologic tests for chronic hepatitis B infections should include three HBV seromarkers: HBsAg, anti-HBs, and Total anti-HBc, while testing for acute infection should include HBsAg and IgM anti-HBc. The CDC provides the following chart on interpreting serologic testing results:

HBsAg anti-HBc anti-HBs	negative negative negative	Susceptible
HBsAg anti-HBc anti-HBs	negative positive positive	Immune due to natural infection
HBsAg anti-HBc anti-HBs	negative negative positive	Immune due to hepatitis B vaccination
HBsAg anti-HBc IgM anti-HBc anti-HBs	positive positive positive negative	Acutely infected
HBsAg anti-HBc IgM anti-HBc anti-HBs	positive positive negative negative	Chronically infected
HBsAg anti-HBc anti-HBs	negative positive negative	Interpretation unclear; four possibilities: 1. Resolved infection (most common) 2. False-positive anti-HBc, thus susceptible 3. "Low level" chronic infection 4. Resolving acute infection

Adapted from: A Comprehensive Immunization Strategy to Eliminate Transmission of Hepatitis B Virus Infection in the United States: Recommendations of the Advisory Committee on Immunization Practices. Part I: Immunization of Infants, Children, and Adolescents. *MMWR* 2005;54(No. RR-16).

(CDC, 2020a)

**Figure 1:** Interpreting HBV serologic testing results.

For health care providers and viral hepatitis, the CDC makes the following recommendation: “Health care providers should be vaccinated against hepatitis B and tested for hepatitis C after a potential exposure. . . For continued protection, CDC and the Advisory Committee on Immunization Practices (ACIP) recommend that health-care providers and public-safety workers with reasonably anticipated risk for exposures to blood or infectious body fluids receive the complete hepatitis B vaccine series and have their immunity documented through postvaccination testing” (CDC, 2020b).

### *Hepatitis A*

Hepatitis A does not present as a chronic infection, and so the CDC offers no testing recommendations. For acute infections, IgM anti-HAV is the serologic test of choice, but “testing for past acute infection is generally not recommended” (CDC, 2020a).

### **United States Preventive Services Task Force (USPSTF)**

The USPSTF also recommends hepatitis C virus screening in adults aged 18 to 79 years (B recommendation) with anti-HCV antibody testing followed by confirmatory PCR testing (Owens et al., 2020).



The United States Preventive Services Task Force (USPSTF) recommends screening for hepatitis B virus (HBV) infection in adolescents and adults at increased risk for infection. This applies to all asymptomatic, nonpregnant adolescents and adults at increased risk for HBV infection, including those who were vaccinated before being screened for HBV infection. The USPSTF defines some increased-risk groups as “Persons born in the US with parents from regions with higher prevalence are also at increased risk of HBV infection during birth or early childhood, particularly if they do not receive appropriate passive and active immunoprophylaxis (and antiviral therapy for pregnant women with a high viral load)” and also “persons who have injected drugs in the past or currently; men who have sex with men; persons with HIV; and sex partners, needle sharing contacts, and household contacts of persons known to be HBsAg positive” (US Preventive Services Task Force, 2020).

USPSTF recommends the following in relation to screening tests for HBV: “Screening for hepatitis B should be performed with HBsAg tests approved by the US Food and Drug Administration, followed by a confirmatory test for initially reactive results. A positive HBsAg result indicates chronic or acute infection. Serologic panels performed concurrently with or after HBsAg screening allow for diagnosis and to determine further management. (See the Additional Tools and Resources section for serologic test interpretation [Figure 1]).”

### **American Association for the Study of Liver Diseases (AASLD) and the Infectious Disease Society of America (IDSA)**

AASLD-IDSA guidelines recommend one-time HCV testing in the following situations:

- “One-time, routine, opt out HCV testing is recommended for all individuals aged 18 years and older. Rating: I, B
- One-time HCV testing should be performed for all persons less than 18 years old with activities, exposures, or conditions or circumstances associated with an increased risk of HCV infection (see below). Rating: I, B
- Prenatal HCV testing as part of routine prenatal care is recommended with each pregnancy. Rating: I, B
- Periodic repeat HCV testing should be offered to all persons with activities, exposures, or conditions or circumstances associated with an increased risk of HCV exposure (see below). Rating: IIa, C
- Annual HCV testing is recommended for all persons who inject drugs, for HIV-infected men who have unprotected sex with men, and men who have sex with men taking pre-exposure prophylaxis (PrEP). Rating: IIa, C

#### Risk Activities

- Injection-drug use (current or ever, including those who injected once)
- Intranasal illicit drug use
- Use of glass crack pipes
- Male engagement in sex with men
- Engagement in chem sex (defined as the intentional combining of sex with the use of particular nonprescription drugs in order to facilitate or enhance the sexual encounter)

#### Risk exposures

- Persons on long-term hemodialysis (ever)
- Persons with percutaneous/parenteral exposures in an unregulated setting
- Healthcare, emergency medical, and public safety workers after needlestick, sharps, or mucosal exposures to HCV-infected blood
- Children born to HCV-infected [individuals]
- Recipients of a prior transfusion or organ transplant, including persons who:
  - Were notified that they received blood from a donor who later tested positive for HCV
  - Received a transfusion of blood or blood components, or underwent an organ transplant before July 1992
  - Received clotting factor concentrates produced before 1987.
- Persons who were ever incarcerated

#### Other considerations and circumstances

- HIV infection
- Sexually active persons about to start pre-exposure prophylaxis (PreP) for HIV
- Chronic liver disease and/or chronic hepatitis, including unexplained elevated alanine aminotransferase (ALT) levels.
- Solid organ donors (living and deceased) and solid organ transplant recipients” (AASLD-IDSA, 2021)

#### *Recommendations for Initial HCV Testing and Follow-up*

- “HCV-antibody testing with reflex HCV RNA polymerase chain reaction (PCR) is recommended for initial HCV testing. Rating: Class I, Level A
- Among persons with a negative HCV-antibody test who were exposed to HCV within the prior 6 months, HCV-RNA or follow-up HCV-antibody testing 6 months or longer after exposure is recommended. HCV-RNA testing can also be considered for immunocompromised persons. Rating: Class I, Level C
- Among persons at risk of reinfection after previous spontaneous or treatment-related viral clearance, initial HCV-RNA testing is recommended because a positive HCV-antibody test is expected. Rating: Class I, Level C
- Persons found to have a positive HCV-antibody test and negative results for HCV RNA by PCR should be informed that they do not have evidence of current (active) HCV infection but are not protected from reinfection. Rating: Class I, Level A
- Quantitative HCV-RNA testing is recommended prior to the initiation of antiviral therapy to document the baseline level of viremia (ie, baseline viral load). Rating: Class I, Level A
- HCV genotype testing may be considered for those in whom it may alter treatment recommendations. Rating: Class I, Level A” (AASLD-IDSA, 2021; Ghany & Morgan, 2020)

For diagnosing and monitoring acute HCV infection, AASLD-IDSA issued the following recommendation:

- “HCV antibody and HCV RNA testing are recommended when acute HCV infection is suspected due to exposure, clinical presentation, or elevated aminotransferase levels.” (Rating: Class I, Level C)
- “After the initial diagnosis of acute HCV with viremia (defined as quantifiable RNA), HCV treatment should be initiated without awaiting spontaneous resolution.” (Rating: Class I, Level B) (AASLD-IDSA, 2022)

For monitoring patients who are starting hepatitis C treatment, are on treatment, or have completed therapy, AASLD-IDSA issued the following recommendations:

- “The following laboratory tests are recommended within 6 months prior to starting DAA (direct-acting antiviral) therapy:
  - Complete blood count (CBC)
  - International normalized ratio (INR)
  - Hepatic function panel (ie, serum albumin, total and direct bilirubin, alanine aminotransferase [ALT], aspartate aminotransferase [AST], and alkaline phosphatase levels)
  - Estimated glomerular filtration rate (eGFR)
- The following laboratory tests are recommended any time prior to starting DAA therapy:
  - Quantitative HCV RNA (HCV viral load)
  - If a nonpangenotypic DAA will be prescribed, then test for HCV genotype and subtype” (Rating: Class I, Level C)
- “Quantitative HCV viral load testing is recommended 12 or more weeks after completion of therapy to document sustained virologic response (SVR), which is consistent with cure of chronic infection” (Rating: Class I, Level B)

#### *Recommendations for Post-Treatment Follow-Up for Patients in Whom Treatment Failed*

- “Disease progression assessment every 6 to 12 months with a hepatic function panel, complete blood count (CBC), and international normalized ratio (INR) is recommended if patients are not retreated or fail a second or third DAA treatment course. (Rating: Class I, Level C)
- Surveillance for hepatocellular carcinoma with liver ultrasound examination, with or without alpha fetoprotein (AFP), every 6 months is recommended for patients with cirrhosis in accordance with the AASLD guidance on the diagnosis, staging, and management of hepatocellular carcinoma. Rating: Low, Conditional” (AASLD-IDSA, 2021).

#### *Recommendations for Monitoring HCV-Infected [Individuals] During Pregnancy*

- “As part of prenatal care, all pregnant [individuals] should be tested for HCV infection, ideally at the initial visit. (Rating: IIb, C)

- HCV RNA and routine liver function tests are recommended at initiation of prenatal care for HCV-antibody-positive pregnant [individuals] to assess the risk of mother-to-child transmission (MTCT) and severity of liver disease. (Rating: I, B)
- All pregnant [individuals] with HCV infection should receive prenatal and intrapartum care that is appropriate for their individual obstetric risk(s) as there is no currently known intervention to reduce MTCT. (Rating: I, B)
- In HCV-infected pregnant [individuals] with pruritus or jaundice, there should be a high index of suspicion for intrahepatic cholestasis of pregnancy (ICP) with subsequent assessment of alanine aminotransferase (ALT), aspartate aminotransferase (AST), and serum bile acids. (Rating: I, B)
- HCV-infected [individuals] with cirrhosis should be counseled about the increased risk of adverse maternal and perinatal outcomes. Antenatal and perinatal care should be coordinated with a maternal-fetal medicine (ie, high-risk pregnancy) obstetrician (Rating: I, B)” (AASLD-IDSA, 2020).

### *Assessment of Liver Disease Severity*

A section focused on determining the severity of liver diseases associated with an HCV infection is also included as part of the background of these AASLD-IDSA guidelines. The authors state the following:

“The severity of liver disease associated with chronic HCV infection is a key factor in determining the initial and follow-up evaluation of patients. Noninvasive tests using serum biomarkers, elastography, or liver imaging allow for accurate diagnosis of cirrhosis in most individuals (see pretreatment workup in *When and in Whom to Initiate HCV Therapy*). Liver biopsy is rarely required but may be considered if other causes of liver disease are suspected.

Noninvasive methods frequently used to estimate liver disease severity include:

- Liver-directed physical exam (normal in most patients)
- Routine blood tests (eg, ALT, AST, albumin, bilirubin, international normalized ratio [INR], and CBC with platelet count)
- Serum fibrosis marker panels
- Transient elastography
- Liver imaging (eg, ultrasound or CT scan)” (AASLD-IDSA, 2021)

### *Testing of Perinatally Exposed Children and Siblings of Children with HCV Infection*

- All children born to women with acute or chronic hepatitis C should be tested for HCV infection.
- Antibody-based testing is recommended at or after 18 months of age. (I, A)
- Testing with an HCV-RNA assay can be considered in the first year of life, but the optimal timing of such testing is unknown. (IIa, C)
- Testing with an HCV-RNA assay can be considered as early as 2 months of age. (IIa, B)
- Repetitive HCV-RNA testing prior to 18 months of age is not recommended. (III, A)
- Children who are anti-HCV-positive after 18 months of age should be tested with an HCV RNA assay after 3 to confirm chronic hepatitis C infection. (I, A)

- The siblings of children with vertically acquired chronic hepatitis C should be tested for HCV infection, if born from the same mother. (I, C) (Ghany & Morgan, 2020)

Testing recommendations relating to the monitoring and medical management of children include

- “Routine liver biochemistries at initial diagnosis and at least annually thereafter are recommended to assess for HCV disease progression. (I, C)”
- “Disease severity assessment by routine laboratory testing and physical examination, as well as use of evolving noninvasive modalities (i.e., transient elastography, imaging, or serum fibrosis markers) is recommended for all children with chronic hepatitis C. (I, B)” (Ghany & Morgan, 2020)

### **American Association for the Study of Liver Diseases (AASLD)**

#### *Hepatitis B*

The guidance statements surrounding screening for hepatitis B infection is (shown in more detail following) declare that

1. Screening should be performed using both HBsAg and anti-HBs.
2. Screening is recommended in all persons born in countries with a HBsAg seroprevalence of  $\geq 2\%$ , U.S.-born persons not vaccinated as infants whose parents were born in regions with high HBV endemicity ( $\geq 8\%$ ), pregnant women, persons needing immunosuppressive therapy, and the at-risk groups listed in Table 3.
3. Anti-HBs–negative screened persons should be vaccinated.
4. Screening for anti-HBc to determine prior expo-sure is not routinely recommended but is an important test in patients who have HIV infection, who are about to undergo HCV or anti-cancer and other immunosuppressive therapies or renal dialysis, and in donated blood (or, if feasible, organs)(Terrault et al., 2018).

AASLD recommends that the following groups are at high risk for HBV infection and should be screened and immunized if seronegative:

**TABLE 3. Groups at High Risk for HBV Infection Who Should Be Screened**

- Persons born in regions of high or intermediate HBV endemicity (HBsAg prevalence of  $\geq 2\%$ )
  - Africa (all countries)
  - North, Southeast, East Asia (all countries)
  - Australia and South Pacific (all countries except Australia and New Zealand)
  - Middle East (all countries except Cyprus and Israel)
  - Eastern Europe (all countries except Hungary)
  - Western Europe (Malta, Spain, and indigenous populations of Greenland)
  - North America (Alaskan natives and indigenous populations of Northern Canada)
  - Mexico and Central America (Guatemala and Honduras)
  - South America (Ecuador, Guyana, Suriname, Venezuela, and Amazonian areas)
  - Caribbean (Antigua-Barbuda, Dominica, Grenada, Haiti, Jamaica, Saint Kitts and Nevis, Saint Lucia, and Turks and Caicos Islands)
- U.S.-born persons not vaccinated as an infant whose parents were born in regions with high HBV endemicity ( $\geq 8\%$ )<sup>\*</sup>
- Persons who have ever injected drugs<sup>\*</sup>
- Men who have sex with men<sup>\*</sup>
- Persons needing immunosuppressive therapy, including chemotherapy, immunosuppression related to organ transplantation, and immunosuppression for rheumatological or gastroenterologic disorders.
- Individuals with elevated ALT or AST of unknown etiology<sup>\*</sup>
- Donors of blood, plasma, organs, tissues, or semen
- Persons with end-stage renal disease, including predialysis, hemodialysis, peritoneal dialysis, and home dialysis patients<sup>\*</sup>
- All pregnant women
- Infants born to HBsAg-positive mothers<sup>\*</sup>
- Persons with chronic liver disease, e.g., HCV<sup>\*</sup>
- Persons with HIV<sup>\*</sup>
- Household, needle sharing, and sexual contacts of HBsAg positive persons<sup>\*</sup>
- Persons who are not in a long-term, mutually monogamous relationship (e.g.,  $>1$  sex partner during the previous 6 months)<sup>\*</sup>
- Persons seeking evaluation or treatment for a sexually transmitted disease<sup>\*</sup>
- Health care and public safety workers at risk for occupational exposure to blood or blood-contaminated body fluids<sup>\*</sup>
- Residents and staff of facilities for developmentally disabled persons<sup>\*</sup>
- Travelers to countries with intermediate or high prevalence of HBV infection<sup>\*</sup>
- Persons who are the source of blood or body fluid exposures that might require postexposure prophylaxis
- Inmates of correctional facilities<sup>\*</sup>
- Unvaccinated persons with diabetes who are aged 19 through 59 years (discretion of clinician for unvaccinated adults with diabetes who are aged  $\geq 60$  years)<sup>\*</sup>

<sup>\*</sup>Indicates those who should receive hepatitis B vaccine, if seronegative.

Sources: (23,35,36)

(Terrault et al., 2018).

AASLD proposes the use of various screening methods for the diagnose of hepatitis B infection: “HBsAg and antibody to hepatitis B surface antigen (anti-HBs) should be used for screening (Table 4). Alternatively, antibody to hepatitis B core antigen (anti-HBc) can be utilized for screening as long as those who test positive are further tested for both HBsAg and anti-HBs to differentiate current infection from previous HBV exposure. HBV vaccination does not lead to anti-HBc positivity .” The interpretations and follow-up steps of the screening results are summarized in their table:

**TABLE 4. Interpretation of Screening Tests for HBV Infection**

Screening Test Results			Interpretation	Management	Vaccinate?
HBsAg	Anti-HBc	Anti-HBs			
+	+	-	Chronic hepatitis B	Additional testing and management needed	No
-	+	+	Past HBV infection, resolved	No further management unless immunocompromised or undergoing chemotherapy or immunosuppressive therapy	No
-	+	-	Past HBV infection, resolved or false positive	HBV DNA testing if immunocompromised patient	Yes, if not from area of intermediate or high endemicity
-	-	+	Immune	No further testing	No
-	-	-	Uninfected and not immune	No further testing	Yes

*Hepatitis C*

AASLD recommends not repeating hepatitis C viral load testing outside of antiviral therapy, stating that “the results of virologic testing do not change clinical management or outcomes” (AASLD, 2014).

**World Health Organization (WHO)**
*Hepatitis C*

Recommendations on screening for HCV infection (WHO, 2017, 2018):

<b>Testing approach</b>	<b>Recommendations</b>
<b>Focused testing in most affected populations</b>	<p>In all settings (and regardless of whether delivered through facility- or community-based testing), it is recommended that serological testing for HCV antibody (anti-HCV) be offered with linkage to prevention, care and treatment services to the following:</p> <ul style="list-style-type: none"> <li>• Adults and adolescents from populations most affected by HCV infection (i.e. who are either part of a population with high HCV seroprevalence or who have a history of exposure and/or high-risk behaviors for HCV infection);</li> <li>• Adults, adolescents and children with a clinical suspicion of chronic viral hepatitis (i.e. symptoms, signs, laboratory markers) (strong recommendation, low quality of evidence)</li> </ul> <p><i>Note: Periodic re-testing using HCV NAT should be considered for those with ongoing risk of acquisition or reinfection.</i></p>
<b>General population testing</b>	<p>In settings with a <math>\geq 2\%</math> (intermediate) or <math>\geq 5\%</math> (high) HCV antibody seroprevalence in the general population, it is recommended that all adults have access to and be offered HCV serological testing with linkage to prevention, care and treatment services.</p> <p>General population testing approaches should make use of existing community- or facility-based testing opportunities or programs such as HIV or TB clinics, drug treatment services and antenatal clinics (conditional recommendation, low quality of evidence)</p>

<b>Which serological assay to use</b>	<p>To test for serological evidence of past or present infection in adults, adolescents and children (&gt;18 months of age), an HCV serological assay (antibody or antibody/antigen) using either a rapid diagnostic test (RDT) or laboratory-based immunoassay formats that meet minimum safety, quality and performance standards (with regard to both analytical and clinical sensitivity and specificity) is recommended.</p> <ul style="list-style-type: none"> <li>• In settings where there is limited access to laboratory infrastructure and testing, and/or in populations where access to rapid testing would facilitate linkage to care and treatment, RDTs are recommended. (Strong recommendation, low/moderate quality of evidence)</li> </ul>
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In a guideline pertaining to the screening, care, and treatment of people with chronic hepatitis C infection, the WHO has provided the following recommendations on hepatitis C screening:

- “It is recommended that HCV serology testing be offered to individuals who are part of a population with high HCV seroprevalence or who have a history of HCV risk exposure/behaviour. Strong recommendation, moderate quality of evidence
- It is suggested that nucleic acid testing (NAT) for the detection of HCV ribonucleic acid (RNA) be performed directly following a positive HCV serological test to establish the diagnosis of chronic HCV infection, in addition to NAT for HCV RNA as part of the assessment for starting treatment for HCV infection. - Conditional recommendation, very low quality of evidence (WHO, 2016).”

The WHO also includes a table which shows the populations with a high HCV prevalence or who have a history of HCV risk. The following groups are included:

- “Persons who have received medical or dental interventions in health-care settings where infection control practices are substandard
- Persons who have received blood transfusions prior to the time when serological testing of blood donors for HCV was initiated or in countries where serological testing of blood donations for HCV is not routinely performed
- People who inject drugs (PWID)
- Persons who have had tattoos, body piercing or scarification procedures done where infection control practices are substandard
- Children born to mothers infected with HCV
- Persons with HIV infection
- Persons who use/have used intranasal drugs
- Prisoners and previously incarcerated persons (WHO, 2016)”

The WHO also mentions liver function tests several times, stating that “A number of clinical considerations are important for the management of persons with chronic HCV infection”; further, “Pre-treatment evaluation of the risk of adverse events should be based on the patient’s clinical details, concomitant medications, and knowledge of treatment regimen to be administered. The potential for DDIs [drug-drug interactions] should be assessed before treatment, and a regimen that has a low risk of DDI selected. Standard laboratory tests that are



assessed prior to treatment initiation include a full blood count (FBC), international normalized ratio (INR), renal function and liver function tests: ALT, AST, bilirubin, albumin and alkaline phosphatase (WHO, 2016).”

The WHO also mentions that “in persons with HCV infection being treated for TB, it is important to monitor liver function tests” and that “Baseline liver function tests for individuals with chronic liver disease are encouraged prior to initiating treatment for latent TB infection. For individuals with abnormal baseline test results, routine periodic laboratory testing should be carried out during the treatment of latent TB infection (WHO, 2016).”

The recommendations of the WHO for assays and strategies regarding hepatitis C testing are summarized in their table, captured below.

HOW TO TEST FOR CHRONIC HCV INFECTION AND MONITOR TREATMENT RESPONSE	
Topic	Recommendations*
Which serological assays to use	<ul style="list-style-type: none"> <li>To test for serological evidence of past or present infection in adults, adolescents and children (&gt;18 months of age<sup>1</sup>), an HCV serological assay (antibody or antibody/antigen) using either RDT or laboratory-based immunoassay formats<sup>2</sup> that meet minimum safety, quality and performance standards<sup>3</sup> (with regard to both analytical and clinical sensitivity and specificity) is recommended.               <ul style="list-style-type: none"> <li>In settings where there is limited access to laboratory infrastructure and testing, and/or in populations where access to rapid testing would facilitate linkage to care and treatment, RDTs are recommended.</li> </ul> <p><i>Strong recommendation, low/moderate quality of evidence</i></p> </li> </ul>
Serological testing strategies	<p>In adults and children older than 18 months<sup>1</sup>, a single serological assay for initial detection of serological evidence of past or present infection is recommended prior to supplementary nucleic acid testing (NAT) for evidence of viraemic infection.</p> <p><i>Conditional recommendation, low quality of evidence</i></p>
Detection of viraemic infection	<ul style="list-style-type: none"> <li>Directly following a reactive HCV antibody serological test result, the use of quantitative or qualitative NAT for detection of HCV RNA is recommended as the preferred strategy to diagnose viraemic infection.</li> <p><i>Strong recommendation, moderate/low quality of evidence</i></p> <li>An assay to detect HCV core (p22) antigen, which has comparable clinical sensitivity to NAT, is an alternative to NAT to diagnose viraemic infection<sup>4</sup>.</li> <p><i>Conditional recommendation, moderate quality of evidence</i></p> </ul>
Assessment of HCV treatment response	<ul style="list-style-type: none"> <li>Nucleic acid testing for qualitative or quantitative detection of HCV RNA should be used as test of cure at 12 or 24 weeks (i.e. sustained virological response [SVR12 or SVR24]) after completion of antiviral treatment.</li> <p><i>Conditional recommendation, moderate/low quality of evidence</i></p> </ul>

(WHO, 2017)  
Hepatitis B

The below table details the populations who should be tested for chronic hepatitis B infection, according to the WHO.

WHO TO TEST FOR CHRONIC HBV INFECTION	
Testing approach and population	Recommendations*
<b>General population testing</b>	<p>1. In settings with a <math>\geq 2\%</math> or <math>\geq 5\%</math><sup>1</sup> HBsAg seroprevalence in the general population, it is recommended that all adults have routine access to and be offered HBsAg serological testing with linkage to prevention, care and treatment services.</p> <p>General population testing approaches should make use of existing community- or health facility-based testing opportunities or programmes such as at antenatal clinics, HIV or TB clinics.</p> <p><i>Conditional recommendation, low quality of evidence</i></p>
<b>Routine testing in pregnant women</b>	<p>2. In settings with a <math>\geq 2\%</math> or <math>\geq 5\%</math><sup>1</sup> HBsAg seroprevalence in the general population, it is recommended that HBsAg serological testing be routinely offered to all pregnant women in antenatal clinics<sup>2</sup>, with linkage to prevention, care and treatment services. Couples and partners in antenatal care settings should be offered HBV testing services.</p> <p><i>Strong recommendation, low quality of evidence</i></p>
<b>Focused testing in most affected populations</b>	<p>3. In all settings (and regardless of whether delivered through facility- or community-based testing), it is recommended that HBsAg serological testing and linkage to care and treatment services be offered to the following individuals:</p> <ul style="list-style-type: none"> <li>• <b>Adults and adolescents from populations most affected by HBV infection<sup>3</sup></b> (i.e. who are either part of a population with high HBV seroprevalence or who have a history of exposure and/or high-risk behaviours for HBV infection);</li> <li>• <b>Adults, adolescents and children with a clinical suspicion of chronic viral hepatitis<sup>4</sup></b> (i.e. symptoms, signs, laboratory markers);</li> <li>• <b>Sexual partners, children and other family members, and close household contacts</b> of those with HBV infection<sup>5</sup>;</li> <li>• <b>Health-care workers:</b> in all settings, it is recommended that HBsAg serological testing be offered and hepatitis B vaccination given to all health-care workers who have not been vaccinated previously (adapted from existing guidance on hepatitis B vaccination<sup>6</sup>).</li> </ul> <p><i>Strong recommendation, low quality of evidence</i></p>
<b>Blood donors</b> <i>Adapted from existing 2010 WHO guidance (Screening donated blood for transfusion transmissible infections<sup>7</sup>)</i>	<p>4. In all settings, screening of blood donors should be mandatory with linkage to care, counselling and treatment for those who test positive.</p>

(WHO, 2017)

Similarly, the recommendations of the WHO for assays and strategies regarding hepatitis B testing are summarized in their table, captured below.

HOW TO TEST FOR CHRONIC HBV INFECTION AND MONITOR TREATMENT RESPONSE	
Topic	Recommendations*
<b>Which serological assays to use</b>	<ul style="list-style-type: none"> <li>For the diagnosis of chronic HBV infection in adults, adolescents and children (&gt;12 months of age), a serological assay (in either RDT or laboratory-based immunoassay format) that meets minimum quality, safety and performance standards<sup>2</sup> (with regard to both analytical and clinical sensitivity and specificity) is recommended to detect hepatitis B surface antigen (HBsAg).               <ul style="list-style-type: none"> <li>In settings where existing laboratory testing is already available and accessible, laboratory-based immunoassays are recommended as the preferred assay format.</li> <li>In settings where there is limited access to laboratory testing and/or in populations where access to rapid testing would facilitate linkage to care and treatment, use of RDTs is recommended to improve access.</li> </ul> </li> </ul> <p><i>Strong recommendation, low/moderate quality of evidence</i></p>
<b>Serological testing strategies</b>	<ul style="list-style-type: none"> <li>In settings or populations with an HBsAg seroprevalence of <math>\geq 0.4\%</math>, a single serological assay for detection of HBsAg is recommended, prior to further evaluation for HBV DNA and staging of liver disease.</li> <li>In settings or populations with a low HBsAg seroprevalence of <math>&lt; 0.4\%</math>, confirmation of HBsAg positivity on the same immunoassay with a neutralization step or a second different RDT assay for detection of HBsAg may be considered<sup>2</sup>.</li> </ul> <p><i>Conditional recommendation, low quality of evidence</i></p>
<b>Detection of HBV DNA – assessment for treatment</b> <i>Adapted from existing guidance (WHO HBV 2015 guidelines<sup>2</sup>)</i>	<ul style="list-style-type: none"> <li>Directly following a positive HBsAg serological test, the use of quantitative or qualitative nucleic acid testing (NAT) for detection of HBV DNA is recommended as the preferred strategy and to guide who to treat or not treat.</li> </ul> <p><i>Strong recommendation, moderate/low quality of evidence</i></p>
<b>Monitoring for HBV treatment response and disease progression</b> <i>Existing guidance (WHO HBV 2015 guidelines<sup>2</sup>)</i>	<ul style="list-style-type: none"> <li><b>It is recommended that the following be monitored at least annually:</b> <ul style="list-style-type: none"> <li>ALT levels (and AST for APRI), HBsAg<sup>3</sup>, HBeAg<sup>3</sup>, and HBV DNA levels (where HBV DNA testing is available)</li> <li>Non-invasive tests (APRI score or transient elastography) to assess for presence of cirrhosis in those without cirrhosis at baseline.</li> <li>If on treatment, adherence should be monitored regularly and at each visit.</li> </ul> </li> </ul> <p><i>Strong recommendation, moderate quality of evidence</i></p> <p><b>More frequent monitoring is recommended:</b></p> <ul style="list-style-type: none"> <li><b>In persons on treatment or following treatment discontinuation:</b> more frequent on-treatment monitoring (at least every 3 months for the first year) is indicated in: persons with more advanced disease (compensated or decompensated cirrhosis<sup>3</sup>), during the first year of treatment to assess treatment response and adherence; where treatment adherence is a concern; in HIV-coinfected persons; and in persons after discontinuation of treatment. <i>Conditional recommendation, very low quality of evidence</i></li> <li><b>In persons who do not yet meet the criteria for antiviral therapy:</b> i.e. persons who have intermittently abnormal ALT levels or HBV DNA levels that fluctuate between 2000 IU/mL and 20 000 IU/mL (where HBV DNA testing is available) and in HIV-coinfected persons<sup>2</sup>. <i>Conditional recommendation, low quality of evidence</i></li> </ul>

(WHO, 2017)

## American Gastroenterological Association (AGA)

### Hepatitis B

“The AGA recommends screening for HBV (HBsAg and anti-HBc, followed by a sensitive HBV DNA test if positive) in patients at moderate or high risk who will undergo immunosuppressive drug therapy. (*Strong recommendation; Moderate-quality evidence*) The AGA suggests against routinely screening for HBV in patients who will undergo immunosuppressive drug therapy and are at low risk. (*Weak recommendation; Moderate-quality evidence*) Comments: *Patients in populations with a baseline prevalence likely exceeding 2% for chronic HBV should be screened according to Centers for Disease Control and Prevention and US Preventive Services Task Force recommendations*” (Reddy et al., 2015).

### Hepatitis C

The AGA released best practice statements for care of patients with chronic HCV that have achieved a sustained virologic response (SVR).

- “SVR should be confirmed by undetectable HCV RNA at 12 weeks after completion of an all-oral DAA treatment regimen.”
- “Routine confirmation of SVR at 48 weeks post end of treatment is recommended. Testing for HCV RNA at 24 weeks post treatment should be considered on an individual patient basis.”
- “Routine testing for HCV RNA beyond 48 weeks after end of treatment to evaluate for late virologic relapse is not supported by available evidence; periodic testing for HCV RNA is recommended for patients with ongoing risk factors for reinfection” (Jacobson et al., 2017).

The AGA has also released a “pathway” for HCV treatment (an algorithm).

Prior to treatment, the AGA recommends identifying the HCV genotype, as well as taking a hepatic function panel (defined as albumin, total and direct bilirubin, alanine aminotransferase, aspartate aminotransferase, and alkaline phosphatase).

For all three lengths of treatment courses (8, 12, 16 weeks), the AGA recommends assessing viral load and liver function (the same hepatic panel listed above) (Kanwal et al., 2017).

### **European Association for the Study of the Liver (EASL)**

#### *Hepatitis C*

The EASL released guidelines on treatment of hepatitis C. The EASL recommends:

- “Screening strategies for HCV infection should be defined according to the local epidemiology of HCV infection, ideally within the framework of local, regional or national action plans.
- Liver disease severity must be assessed prior to therapy.
- Rapid diagnostic tests using serum, plasma, fingerstick whole blood or crevicular fluid (saliva) as matrices can be used instead of classical EIAs as point-of-care tests to facilitate anti-HCV antibody screening and improve access to care.
- “It is still useful to determine the HCV genotype and subtype where such determination is available and does not limit access to care, to identify patients who may benefit from treatment tailoring. However, “testing for HCV resistance prior to treatment is not recommended” (EASL, 2018, 2020).

#### *Hepatitis B*

EASL states that “The initial evaluation of a subject with chronic HBV infection should include a complete history, a physical examination, assessment of liver disease activity and severity and markers of HBV infection (Fig. 1). In addition, all first-degree relatives and sexual partners of subjects with chronic HBV infection should be advised to be tested for HBV serological markers (HBsAg, anti-HBs, anti-HBc) and to be vaccinated if they are negative for these markers.”

“The assessment of the severity of liver disease is important to identify patients for treatment and HCC surveillance. It is based on a physical examination and biochemical parameters (aspartate aminotransferase [AST] and ALT, gamma-glutamyl transpeptidase [GGT], alkaline phosphatase, bilirubin, and serum albumin and gamma globulins, full blood count and prothrombin time). An abdominal hepatic ultrasound is recommended in all patients. A liver biopsy or a non-invasive test should be performed to determine disease activity in cases where biochemical and HBV markers reveal inconclusive results.”

“HBeAg and anti-HBe detection are essential for the determination of the phase of chronic HBV infection.”

“Measurement of HBV DNA serum level is essential for the diagnosis, establishment of the phase of the infection, the decision to treat and subsequent monitoring of patients.”

“Serum HBsAg quantification can be useful, particularly in HBeAg-negative chronic HBV infection and in patients to be treated with interferon-alfa (IFN $\alpha$ ).”

“HBV genotype is not necessary in the initial evaluation, although it may be useful for selecting patients to be treated with IFN $\alpha$  offering prognostic information for the probability of response to IFN $\alpha$  therapy and the risk of HCC.”

“Co-morbidities, including alcoholic, autoimmune, metabolic liver disease with steatosis or steatohepatitis and other causes of chronic liver disease should be systematically excluded including co-infections with hepatitis D virus (HDV), hepatitis C virus (HCV) and HIV.”

“Testing for antibodies against hepatitis A virus (anti-HAV) should be performed, and patients with negative anti-HAV should be advised to be vaccinated against HAV.”

“Screening for HBsAg in the first trimester of pregnancy is strongly recommended (Evidence level 1, grade of recommendation 1)” (European Association for the Study of the Liver, 2017).

### **Indian Health Services (IHS)**

Indian Health Services published recommendations on Hepatitis C screening. IHS recommends using an anti-HCV antibody test such as a point-of-care test on a fingerstick capillary or venipuncture whole-blood sample or a laboratory-based HCV ELISA test on a serum sample. IHS recommends screening the following patients:

- "Adults 18 years and older, including people with diabetes, at least once for HCV infection, regardless of their risk factors.
- All pregnant persons, regardless of age, during each pregnancy.
- People at higher risk of HCV exposure (IHS, 2021)."

IHS also provides guidance on how to diagnose a chronic HCV infection:

- "For individuals with a positive HCV antibody screening test result, perform the laboratory-based HCV RNA PCR test to confirm the presence of HCV.
- The presence of HCV indicates active infection. These individuals should be referred for direct acting anti-viral (DAA) agents treatment.

- The absence of HCV indicates no active infection.
- For individuals with a negative HCV antibody test result who might have been exposed to HCV within the previous 6 months, perform an HCV RNA PCR or follow-up HCV antibody test at least 6 months after exposure (IHS, 2021)."

Regarding hepatitis B, the HIS suggests that

“People who inject drugs illicitly, including participants in substance abuse treatment programs, should be offered screening and counseling for chronic HBV infection. Testing should include a serologic assay for hepatitis B surface antigen (HBsAg) offered as a part of routine care, and if the result is positive, be accompanied by appropriate counseling and referral for recommended clinical evaluation and care. Previous and current sex partners and household and needle-sharing contacts of HBsAg-positive persons should be identified. Unvaccinated sex partners and household and needle-sharing contacts should be tested for HBsAg and for antibody to the hepatitis B core antigen (anti-HBc) or antibody to the hepatitis B surface antigen (anti-HBsAg)” (Indian Health Service).

## VI. Applicable State and Federal Regulations

### Food and Drug Administration (FDA)

Many labs have developed specific tests that they must validate and perform in house. These laboratory-developed tests (LDTs) are regulated by the Centers for Medicare and Medicaid (CMS) as high-complexity tests under the Clinical Laboratory Improvement Amendments of 1988 (CLIA '88). LDTs are not approved or cleared by the U. S. Food and Drug Administration; however, FDA clearance or approval is not currently required for clinical use.

## VII. Applicable CPT/HCPCS Procedure Codes

Procedure codes appearing in medical policy documents are only included as a general reference. This list may not be all inclusive and is subject to updates. In addition, codes listed are not a guarantee of payment.

Code NumberCPT	Code Description
86704	Hepatitis B core antibody (HBcAb); total
86705	Hepatitis B core antibody (HBcAb); IgM antibody
86706	Hepatitis B surface antibody (HBsAb)
86803	Hepatitis C antibody
86804	Hepatitis C antibody; confirmatory test (eg, immunoblot)
87340	Infectious agent antigen detection by immunoassay technique, (eg, enzyme immunoassay [EIA], enzyme-linked immunosorbent assay [ELISA], fluorescence immunoassay [FIA], immunochemiluminometric assay [IMCA]) qualitative or semiquantitative; hepatitis B surface antigen (HBsAg)
87341	Infectious agent antigen detection by immunoassay technique, (eg, enzyme immunoassay [EIA], enzyme-linked immunosorbent assay [ELISA], fluorescence immunoassay [FIA], immunochemiluminometric assay [IMCA])

Code NumberCPT	Code Description
	qualitative or semiquantitative; hepatitis B surface antigen (HBsAg) neutralization
87517	Infectious agent detection by nucleic acid (DNA or RNA); hepatitis B virus, quantification
87520	Infectious agent detection by nucleic acid (DNA or RNA); hepatitis C, direct probe technique
87521	Infectious agent detection by nucleic acid (DNA or RNA); hepatitis C, amplified probe technique, includes reverse transcription when performed
87522	Infectious agent detection by nucleic acid (DNA or RNA); hepatitis C, quantification, includes reverse transcription when performed
87902	Infectious agent genotype analysis by nucleic acid (DNA or RNA); Hepatitis C virus
G0472	Hepatitis C antibody screening, for individual at high risk and other covered indication(s)
G0499	Hepatitis b screening in non-pregnant, high risk individual includes hepatitis b surface antigen (hbsag), antibodies to hbsag (anti-hbs) and antibodies to hepatitis b core antigen (anti-hbc), and is followed by a neutralizing confirmatory test, when performed, only for an initially reactive hbsag result

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## IX. Revision History

Revision Date	Summary of Changes
12/01/2021	Initial Effective Date

Revision Date	Summary of Changes
04/12/2022	<p>Updated background, guidelines, and evidence-based scientific references. Based on the 2020 CDC guidelines, updated following CC:</p> <p>Added “Screening at least once in a lifetime” to CC1. Added “for all adults (≥18 years old) with recognized conditions or exposures” to CC2. Also combined CC2 with CC describing testing based on recognized exposure, combined CC2a and 2b to “illicit intranasal or injectable drug use”; removed “or current” from hemodialysis.</p> <p>Added following CC:</p> <p>3. Routine periodic HCV testing MEETS COVERAGE CRITERIA for individuals with ongoing risk factors, while risk factors persist:</p> <p>a. Individuals who currently inject drugs and share needles, syringes, or other drug preparation equipment</p> <p>b. individuals who are receiving ongoing hemodialysis</p>
10/02/2022	<p>Updated background, guidelines, and evidence-based scientific references. Guideline updates necessitated the following CC updates:</p> <p>Addition of new CC1: “Screening for Hepatitis C virus (HCV) infection in pregnant individuals is covered in accordance with Avalon Policy AHS-G2035-Prenatal Screening (Nongenetic).</p> <p>Upper age limit removed from CC2, additional edits for clarity. Formerly read “Screening at least once in a lifetime for Hepatitis C infection MEETS COVERAGE CRITERIA for adults between the ages of 18 years and 79 years.” Now reads “Once per lifetime testing for Hepatitis C virus (HCV) infection MEETS COVERAGE CRITERIA for all individuals over 18 years of age.”</p> <p>CC3 edited for clarity and age limit removed based on guideline updates, which allow for those under 18 to get tested if they have one of the known risk exposures.</p> <p>Previously read “Testing for Hepatitis C infection for all adults (&gt;18 years old) with recognized conditions or exposures MEETS COVERAGE CRITERIA in the following situations” Now reads “For any individual with the following recognized conditions or exposures, one-time, post-exposure testing for HCV infection MEETS COVERAGE CRITERIA:”</p> <p>CC31 changed “Children born to HCV-positive women” to “Children born from an HCV-positive individual” for consistency with removing gendered language from Avalon CC when possible</p> <p>Addition of CC4c: “Individuals engaging in high-risk sexual behavior”</p> <p>Revised code disclaimer statement</p>
09/15/2023	Literature review necessitated the following changes in coverage criteria:

Revision Date	Summary of Changes
	<p>Policy name changed from “Hepatitis C” to “Hepatitis Testing” and coverage was broadened to include coverage on Hepatitis B.</p> <p>Guidance on HCV testing in pregnant individuals moved from being a coverage criteria to being a note in the Policy Description.</p> <p><b>**Addition of new CC1, 2, 3, and 4 to address screening and following up testing for Hepatitis B “1) For asymptomatic, non-pregnant individuals, the following annual Hepatitis B virus (HBV) infection screening MEETS COVERAGE CRITERIA:</b></p> <p>a) Hepatitis B surface antigen (HBsAg) when one of the following high-risk situations is met:</p> <ul style="list-style-type: none"> <li>i) For individuals born in geographic regions with a HBsAg prevalence 2% or higher.</li> <li>ii) For U.S.-born individuals not vaccinated as infants whose parents were born in geographic regions with a HBsAg prevalence 8% or higher.</li> <li>iii) For individuals with elevated alanine aminotransferase (ALT)/aspartate transaminase (AST) of unknown etiology.</li> </ul> <p>b) HBsAg and hepatitis B surface antibody (anti-HBs) for infants born from an HBsAg-positive individual.</p> <p>c) HBsAg, anti-HBs, and total hepatitis B core antibody (anti-HBc), when one of the following high-risk situations is met:</p> <ul style="list-style-type: none"> <li>i) For injection-drug users.</li> <li>ii) For men who have sex with men.</li> <li>iii) For individuals with select medical conditions who require cytotoxic or immunosuppressive therapy.</li> <li>iv) For individuals who are on long-term hemodialysis treatment.</li> <li>v) For household contacts, needle-sharing contacts, and sex partners of HBV-infected individuals.</li> <li>vi) For individuals infected with HIV.</li> <li>vii) For healthcare and public safety workers exposed to blood or body fluids.</li> </ul> <p><b>**2) For individuals who test positive for HBV by initial antibody screening, follow up serologic testing (anti-HBc, anti-HBs, IgM antibody to anti-HBc [IgM anti-HBc]) MEETS COVERAGE CRITERIA.</b></p> <p><b>**3) For the confirmation of seroconversion after hepatitis B vaccination, anti-HBs testing MEETS COVERAGE CRITERIA.</b></p>

Revision Date	Summary of Changes
	<p>**4) For individuals who test positive for HBV by initial antibody screening and who will undergo immunosuppressive drug therapy, HBV DNA testing MEETS COVERAGE CRITERIA.</p> <p>All other CC edited for clarity and consistency.</p> <p>Specified antibody screening as the allowed test for initial HCV screening in CC5, CC6, and CC7. Reflex NAT testing allowed with new CC8: “7) For individuals who test positive for HCV by initial antibody screening, follow up nucleic acid testing for HCV to differentiate between active infection and resolved infection MEETS COVERAGE CRITERIA.”</p> <p>Removed former CC6: “For patients with acute HCV infection, monitoring HCV RNA MEETS COVERAGE CRITERIA to determine spontaneous clearance of HCV infection versus persistence of infection. Testing can be performed every 4 to 8 weeks for 6 to 12 months.” because of updates to ASSLD-ISDA guidelines that now recommend HCV treatment begin without awaiting spontaneous resolution. Previous versions of this guideline provided recommendations for RNA monitoring for spontaneous clearance of HCV.</p> <p>Addition of “forty-eight weeks” in CC10d: “d) Twelve, twenty-four, and forty-eight weeks after completion of treatment.”</p> <p>Addition of new Note 1: “Note 1: The CDC defines HBsAg prevalence by geographic region: <a href="https://wwwnc.cdc.gov/travel/yellowbook/2020/travel-related-infectious-diseases/hepatitis-b">https://wwwnc.cdc.gov/travel/yellowbook/2020/travel-related-infectious-diseases/hepatitis-b</a>.”</p> <p>Coverage criteria edited for clarity and consistency.</p> <p>**Added CPT codes 86704, 86705, 86706, 87340, 87341, 87517, G0499</p> <p>Committee approved: 06/12/2023</p> <p>DCH approved: 09/15/2023</p>