





2018-2019 PROGRESS REPORT





The 2018-2019 Viral Hepatitis Progress Report was prepared under the direction of the Office of Infectious Disease and HIV/AIDS Policy (OIDP), Office of the Assistant Secretary for Health (OASH), U.S. Department of Health and Human Services (HHS). The National Viral Hepatitis Action Plan 2017–2020 (Action Plan) was developed collaboratively with input from representatives of agencies and offices from across HHS as well as from the U.S. Departments of Housing and Urban Development, Justice, and Veterans Affairs.



VISION

The United States will be a place where new viral hepatitis infections have been eliminated, where all people with chronic hepatitis B and hepatitis C know their status, and everyone with chronic hepatitis B and hepatitis C has access to high-quality health care and curative treatments, free from stigma and discrimination.

COMMITMENT

To help achieve our vision, agencies and offices from across the U.S. Department of Health and Human Services and partners from the U.S. Departments of Housing and Urban Development, Justice, and Veterans Affairs have joined together to improve viral hepatitis prevention and the care and treatment provided to people with hepatitis B and hepatitis C. To be successful in our efforts, we must continue to improve the efficiency, effectiveness, and impact of our work. We must remain flexible to adapt to changing needs and funding levels, and make the best use of scientific, clinical, and programmatic advances. The National Viral Hepatitis Action Plan 2017-2020 (Action Plan) provides a roadmap for this important work, and the federal government is committed to achieving the Action Plan's goals.

Although this is a federal progress report, we acknowledge the tremendous support and commitment of a broad mix of nonfederal stakeholders from various sectors, both public and private, whose work contributes substantially to the nation's progress. Many of the actions reported reflect the work of nonfederal stakeholders, including actions that are supported through grants, cooperative agreements, partnerships, and other collaborative efforts.

The Action Plan itself is a national plan. It emphasizes that all sectors of society have roles to play if we are to achieve our vision and national goals, prevent disease and death, and reduce costs to the health care system. As progress to address viral hepatitis faces new threats, most notably from the opioid crisis, and in 2020 from the COVID-19 pandemic, we must find new ways to work together with a broad variety of stakeholders to sustain our achievements and continue to advance toward the nation's viral hepatitis prevention and care goals. All sectors of society – both federal and nonfederal – are needed to achieve the Action Plan's goals and to realize a future where viral hepatitis in this nation has been eliminated.





CONTENTS

INTRODUCTION	1
Overview of the National Viral Hepatitis Action Plan 2017-2020	1
In This Report	2
PROGRESS ON THE INDICATOR MEASURES	3
Overall Progress on Action Plan Indicator Measures: Based on 2018 Data	4
Graphs and Tables of Each Indicator and Progress Toward 2020 Targets	5
Data Sources	10
FEDERAL ACTIVITIES TO ADVANCE THE UNITED STATES TOWARD NATIONAL HEPATITIS GOALS	11
Departments, Agencies, and Offices that Implemented the Action Plan in FY2018 – FY2019	11
Timeline	15
APPENDIX 1: FEDERAL PROGRESS ON ACTION PLAN	
Goal 1: Prevent New Viral Hepatitis Infections	16
Goal 2: Reduce Deaths and Improve the Health of People Living with Viral Hepatitis	18
Goal 3: Reduce Viral Hepatitis Health Disparities	23
Goal 4: Coordinate, Monitor, and Report on Implementation of Viral Hepatitis Activities	25
APPENDIX 2: 2018/2019 VIRAL HEPATITIS-RELATED PUBLICATIONS, ARTICLES, AND REPORTS BY FEDERAL PARTNERS	
Goal 1: Prevent New Viral Hepatitis Infections	28
Goal 2: Reduce Deaths and Improve the Health of People Living with Viral Hepatitis	33
Goal 3: Reduce Viral Hepatitis Health Disparities	42
Goal 4: Coordinate, Monitor, and Report on Implementation of Viral Hepatitis Activities	43
APPENDIX 3: ABBREVIATIONS	45

INTRODUCTION

This report provides an overview of progress toward achieving the goals of the National Viral Hepatitis Action Plan based on 2018 surveillance data (the most recent data currently available) and key federal actions that were taken during fiscal years (FY) 2018 – 2019.

OVERVIEW OF THE NATIONAL VIRAL HEPATITIS ACTION PLAN 2017-2020

The National Viral Hepatitis Action Plan 2017–2020 (Action Plan) is the third iteration of a strategic roadmap to address viral hepatitis in the United States. Building on progress under the previous iterations, the Action Plan sets four goals and recommends more than 20 strategies to achieve the goals. These strategies, if implemented by the full range of stakeholders, are expected to improve the prevention, diagnosis, and treatment of viral hepatitis in the United States. Federal agencies engaged in implementing these strategies include the U.S. Departments of Health and Human Services (HHS), Housing and Urban Development (HUD), Justice (DOJ), and Veterans Affairs (VA). Nonfederal stakeholders include a wide range of state and local governments, nonprofit and advocacy organizations, academic institutions, health plans, healthcare providers, and professional organizations, as well as private sector groups and companies.

The Action Plan has guided the nation's response to the viral hepatitis epidemic through its goals:

- Goal 1: Prevent new viral hepatitis infections
- Goal 2: Reduce deaths and improve the health of people living with viral hepatitis
- Goal 3: Reduce viral hepatitis health disparities
- Goal 4: Coordinate, monitor, and report on implementation of viral hepatitis activities

In order to help stakeholders with limited resources focus their efforts for the greatest impact, the Action Plan identifies the following disproportionately impacted populations, referred to as priority populations 1, which have higher rates and/or risk for transmission of viral hepatitis:

- Baby boomers (people born during 1945–1965)
- People who inject drugs
- American Indians and Alaska Natives (AI/AN)
- Asian Americans and Pacific Islanders (AAPI)
- African Americans
- People in correctional facilities
- Veterans, particularly those who served during the Vietnam War era
- Homeless individuals
- Men who have sex with men (MSM)

¹ The Action Plan refers to the priority populations utilizing this terminology, but it is recognized that language and preferred terminology change over time.



- Pregnant women
- People living with HIV/AIDS

A key feature of the Action Plan is the 17 indicators used to measure progress toward the national goals. These are reported below.

In support of efforts across the federal government to implement the Action Plan, the Office of Infectious Disease and HIV/AIDS Policy (OIDP), within the Office of the Assistant Secretary for Health (OASH) in HHS, convenes the Viral Hepatitis Implementation Group (VHIG). The VHIG coordinates and monitors implementation of the Action Plan. Its members include representatives from across HHS agencies and other federal departments engaged in implementing the Action Plan. VHIG members meet regularly to share information about resources and initiatives, advance and collaborate on implementation of the Action Plan's strategies, and address new opportunities and challenges. The members represent their respective agencies and offices on matters related to viral hepatitis.

IN THIS REPORT

This report includes:

- Table of overall progress on indicators based on 2018 data.
- Graphs and tables of each indicator and progress toward 2020 targets
- Description of data sources
- A description of each federal partner's role
- Timeline of significant policy, program, and scientific actions by federal partners during FY2018 FY2019.

Appendices:

- Appendix 1: A more detailed list of reported federal actions undertaken in FY2018 FY2019
- Appendix 2: Publications, articles, and reports by federal partners in FY2018 FY2019 that contribute to the growing body of evidence in the field of viral hepatitis
- Appendix 3: Abbreviations used in this report.

PROGRESS ON THE INDICATOR MEASURES

Indicators are important tools that help measure progress toward meeting the goals established in the Action Plan. The indicators were selected because they represent the best way to measure national progress on viral hepatitis prevention and care based on the available data and in alignment with other national plans. The baseline year for the indicators is 2014, the most recent year national surveillance data was available at the time the Action Plan was published. This report uses 2018 surveillance data, which are the most recent data available, to measure progress. Centers for Disease Control and Prevention (CDC) generally reports viral hepatitis surveillance data two years after the calendar year in which they occurred. The lag is due to the time needed to collect data from all jurisdictions, ensure completeness and accuracy, and conduct analyses.

Based on 2018 data, across the 17 indicators:

- Five are on track to meet 2020 targets;
- Three are moving in the right direction but require additional effort to meet the 2020 targets;
- Five are trending in the wrong direction; and
- Four do not have updated data available.

Below is a full-page chart that presents the 17 indicators and, based on 2018 data, summarizes progress toward the 2020 targets. Following that chart is a series of graphs – one for each indicator. The graphs illustrate the annual targets set forth in the Action Plan and the current trends based on CDC surveillance data. They include linear trend projections based on available data; these projections may change as new national data are published. The color of the lines in each graph correspond with progress in meeting the 2020 target, as indicated by the key. The data sources for the indicators are described following the series of graphs.

OVERALL PROGRESS ON ACTION PLAN INDICATOR MEASURES: BASED ON 2018 DATA

	Indicator and Measure	Baseline (2014)	Progre as of 20		2020 Goal	Data Source [†]
GC	DAL 1					
L.	Decrease the number of new HBV infections by at least 60% # estimated and (reported) acute hepatitis B cases in the U.S.	18,090 (2,791)*	21,600	x	7,236 (1,116)*	NNDSS
2.	Increase the rate of hepatitis B vaccine "birth dose" coverage to 85% % children who received the first dose of hepatitis B vaccine within three days of birth	71.8%	NO NEW DATA	0	85.0%	NIS-Chile
3.	Increase the rate of hepatitis B vaccination among health care personnel to 90% % health care personnel 19 years of age and older with direct patient care responsibilities reporting they have had at least three doses of hepatitis B vaccine	67.7%	NO NEW DATA	x	90.0%	NHIS
4.	Decrease the number of new HCV infections by at least 60% # estimated and (reported) acute hepatitis C cases in the U.S.	30,500 (2,194)*	50,300	x	10,889 (783)*	NNDSS
GC	DAL 2					
5.	Increase the percentage of persons aware of their HBV infection to 66% % respondents who indicate they were aware they had hepatitis B prior to laboratory testing	33.0%	NO NEW DATA	\bigcirc	66.0%	NHANES
5.	Reduce the number of HBV-related deaths by 20% # deaths in the U.S. for which hepatitis B is listed as the underlying or a contributing cause of death	1,843	1,649	✓	1,474	NVSS
7.	Increase the percentage of persons aware of their HCV infection to 66% % respondents who indicate they were aware they had hepatitis C prior to laboratory testing	54.0% (2013–2016)	NO NEW DATA	\bigcirc	66.0%	NHANES
8.	Reduce the number of HCV-related deaths by 25% # deaths in the U.S. for which hepatitis C is listed as the underlying or a contributing cause of death	19,659	15,713	✓	14,744	NVSS
GC	DAL 3					
9.	Decrease the number of new HBV infections among individuals 30–49 years of age by at least 60% # reported acute hepatitis B cases for adults 30–49 years of age living in the U.S.	1,706	1,920	x	682	NNDSS
10	Reduce the number of HBV-related deaths among AAPI by at least 20% # deaths among AAPI living in the U.S. for which hepatitis B is listed as the underlying or a contributing cause of death	478	439	\rightarrow	382	NVSS
11	Reduce the number of HBV-related deaths among African Americans by at least 20% # deaths among African Americans living in the U.S. for which hepatitis B is listed as the underlying or a contributing cause of death	330	304	\rightarrow	264	NVSS
12	Reduce the number of HBV-related deaths among individuals 45 years of age and older by at least 20% # deaths among persons ages 45 and older in the U.S. for which hepatitis B is listed as the underlying or a contributing cause of death	1,682	1,495	✓	1,346	NVSS
13	Decrease the number of new HCV infections among individuals 20–39 years of age by at least 60% # acute hepatitis C cases reported for adults 20–39 years of age in the U.S.	1,561	2,380	x	624	NNDSS
14	Decrease the number of new HCV infections among AI/AN by at least 60% # reported acute hepatitis C cases for AI/AN living in the U.S.	29	83	x	12	NNDSS
15	Reduce the number of HCV-related deaths among individuals 55–74 years of age by at least 25% # deaths among persons ages 55–74 in the U.S. for which hepatitis C is listed as the underlying or a contributing cause of death	13,389	11,726	\rightarrow	10,042	NVSS
16	Reduce the number of HCV-related deaths among AI/AN by at least 25% # deaths among AI/AN in the U.S. for which hepatitis C is listed as the underlying or a contributing cause of death	317	264	✓	238	NVSS
17.	Reduce the number of HCV-related deaths among African Americans by at least 25% # deaths among African Americans living in the U.S. for which hepatitis C is listed as the underlying or a contributing cause of death	3,540	2,978	✓	2,655	NVSS
•	on track to achieve 2020 target trending in the right direction target target	🔵 data	not availab	ole		

*In cells that contain two numbers, the initial number is estimated cases, the number in parentheses is reported cases.

⁺ NHANES = <u>National Health and Nutrition Examination Survey</u>; NHIS = <u>National Health Interview Survey</u>; NIS-Child = <u>National Immunization Survey</u>; <u>Children</u>; NNDSS = <u>National Notifiable Diseases Surveillance System</u>; NVSS= <u>National Vital Statistics System</u>



GRAPHS AND TABLES OF EACH INDICATOR AND PROGRESS TOWARD 2020 TARGETS

This section provides graphs of annual targets that are set forth in the Action Plan and the current trends based on available surveillance data. The linear trend projections are based on available data and may change when new national data are published. The color of the lines in each graph correspond with progress in meeting the 2020 target, as indicated by the key below. The methodology for measuring indicators from the NNDSS and NVSS is available in CDC's <u>2018 Viral Hepatitis Surveillance Report</u>. The methodology for measuring indicators from NHANES, NHIS, and NIS-Child is available from each of those data sources.

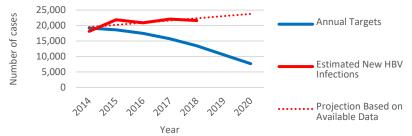
on track to achieve 2020 target

trending in the right direction

not on track to achieve 2020 target

data not available



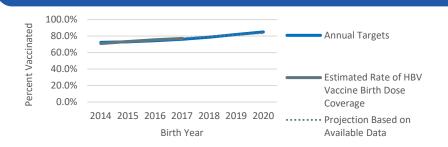


Source: National Notifiable Diseases Surveillance System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS*	19,200	18,624	17,472	15,744	13,440	10,560	7,680
EST. NEW INFECTIONS	18,142	21,905	20,917	22,100	21,600		

* Annual target numbers do not match those reported in the Action Plan because CDC published updated numbers after the Action Plan was published; the updated numbers and targets are presented here.

2. Increase the rate of hepatitis B vaccine "birth dose" coverage to 85% % children who received the first dose of hepatitis B vaccine within three days of birth



Source: National Immunization Surveys - Children

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS*	72.4%	73.0%	74.3%	76.2%	78.7%	81.9%	85.0%
EST. RATE OF COVERAGE	70.9%	73.3%	75.6%	77.1%†	Data not available		

*Annual targets and estimated coverage for this indicator differs from data published in the Action Plan and 2017 Progress Report. CDC transitioned from reporting NIS-Child data by survey year to birth year and this data reflects the change.

⁺Estimates for children born in 2017 are considered preliminary and will be finalized after the data for survey year 2020 are available.



3. Increase the rate of hepatitis B vaccination among health care personnel to 90%

% health care personnel 19 years of age and older with direct patient care responsibilities reporting they have had at least three doses of hepatitis B vaccine



rear

Source: National Health Interview Survey

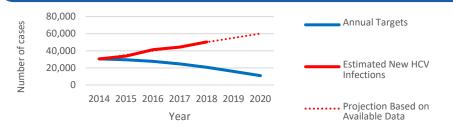
	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	67.7%	68.8%	71.1%	74.4%	78.9%	84.4%	90.0%
EST. RATE OF VACCINATION	67.7%	64.7%	61.4%	69.8%	Data not available		

5. Increase the percentage of persons aware of their HBV infection by 66%

% respondents who indicate they were aware they had hepatitis B prior to laboratory testing

No new data is available for this indicator. Progress is reported on this indicator utilizing 4-year estimates from NHANES. The next 4-year estimate will be from 2017-2020 and is expected to be available in 2021.

4. Decrease the number of new HCV infections by at least 60% # estimated acute hepatitis C cases in the U.S.

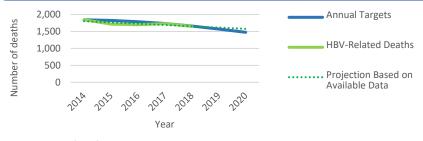


Source: National Notifiable Diseases Surveillance System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	30,500	29,519	27,558	24,617	20,694	15,791	10,889
EST. NEW INFECTIONS	30,497	33,860	41,241	44,300	50,300		

6. Reduce the number of HBV-related deaths by 20%

deaths in the U.S. for which hepatitis B is listed as the underlying or a contributing cause of death



Source: National Health and Nutrition Examination Survey

Source: National Vital Statistics System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	1,843	1,825	1,788	1,732	1,659	1,567	1,474
HBV-RELATED DEATHS	1,843	1,715	1,698	1,727	1,649		

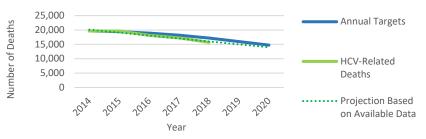


7. Increase the percent of persons aware of their HCV infection to 66% % respondents who indicate they were aware they had hepatitis C prior to laboratory testing

No new data is available for this indicator. Progress is reported on this indicator utilizing 4-year estimates from NHANES. The next 4-year estimate will be from 2017-2020 and is expected to be available in 2021.

8. Reduce the number of HCV-related deaths by 25%

deaths in the U.S. for which hepatitis C is listed as the underlying or a contributing cause of death



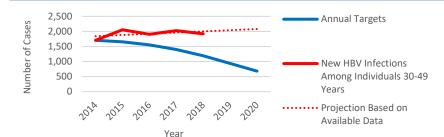
Source: National Health and Nutrition Examination Survey

Source: National Vital Statistics System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	19,659	19,413	18,922	18,185	17,202	15,973	14,744
HCV-RELATED DEATHS	19,659	19,629	18,153	17,253	15,713		

9. Decrease the number of new HBV infections among individuals 30-49 years of age by at least 60%

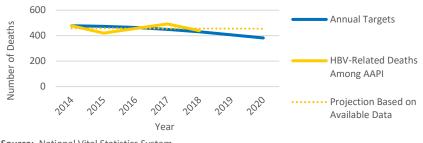
reported acute hepatitis B cases for adults 30-49 years of age living in the U.S.



Source: National Notifiable Diseases Surveillance System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	1,706	1,655	1,552	1,399	1,194	938	682
NEW HBV INFECTIONS AMONG INDIVIDUALS 30-49 YEARS	1,706	2,055	1,906	2,024	1,920		

10. Reduce the number of HBV-related deaths among AAPI by at least 20% # deaths among AAPI living in the U.S. for which hepatitis B is listed as the underlying or a contributing cause of death



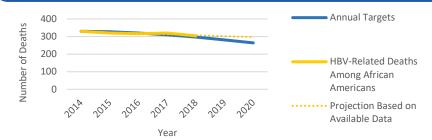
Source: National Vital Statistics System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	478	473	464	449	430	406	382
HBV-RELATED DEATHS AMONG AAPI	478	420	457	492	439		



11. Reduce the number of HBV-related deaths among African Americans by at least 20%

deaths among African Americans living in the U.S. for which hepatitis B listed as the underlying or a contributing cause of death

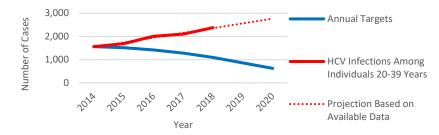


Source: National Vital Statistics System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	330	327	320	310	297	281	264
HBV-RELATED DEATHS AMONG AFRICAN AMERICANS	330	320	316	320	304		

13. Decrease the number of new HCV infections among individuals 20-39 years of age by at least 60%

acute hepatitis C cases reported for adults 20-39 years of age in the U.S.

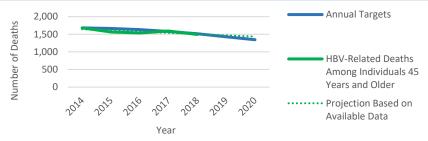


Source: National Notifiable Diseases Surveillance System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	1,561	1,514	1,421	1,280	1,093	859	624
NEW HCV INFECTIONS AMONG INDIVIDUALS 20–39 YEARS	1,561	1,692	2,003	2,105	2,380		

12. Reduce the number of HBV-related deaths among individuals 45 years of age and older by at least 20%

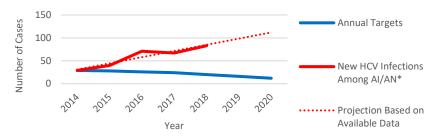
deaths among persons aged 45 and older in the U.S. for which hepatitis B is listed as the underlying or a contributing cause of death



Source: National Vital Statistics System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	1,682	1,665	1,632	1,581	1,514	1,430	1,346
HBV-RELATED DEATHS AMONG INDIVIDUALS 45 YEARS AND OLDER	1,682	1,563	1,540	1,591	1,495		

14. Decrease the number of new HCV infections among AI/AN by at least 60% # reported acute hepatitis C cases for AI/AN living in the U.S.



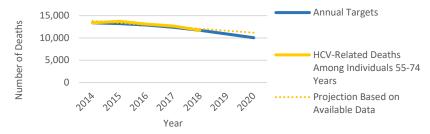
Source: National Notifiable Diseases Surveillance System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	29	28	26	24	20	16	12
NEW HCV INFECTIONS AMONG AI/AN	29	40	71	67	83		



15. Reduce the number of HCV-related deaths among individuals 55-74 years of age by at least 25%

deaths among persons aged 55-74 in the U.S. for which hepatitis C is listed as the underlying or a contributing cause of death

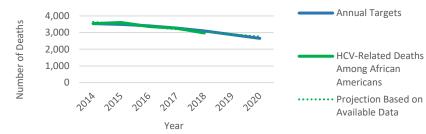


Source: National Vital Statistics System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	13,389	13,222	12,887	12,385	11,715	10,879	10,042
HCV-RELATED DEATHS AMONG INDIVIDUALS 55-74 YEARS	13,389	13,725	13,118	12,672	11,726		

17. Reduce the number of HCV-related deaths among African Americans by at least 25%

deaths among African Americans living in the U.S. for which hepatitis C is listed as the underlying or a contributing cause of death

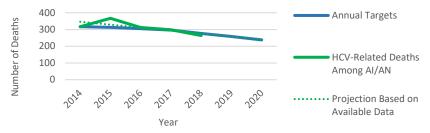


Source: National Vital Statistics System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	3,540	3,496	3,407	3,275	3,098	2,876	2,655
HCV-RELATED DEATHS AMONG AFRICAN AMERICANS	3,540	3,606	3,365	3,262	2,978		

16. Reduce the number of HCV-related deaths among AI/AN by at least 25%

deaths among AI/AN in the U.S. for which hepatitis C is listed as the underlying or a contributing cause of death



Source: National Vital Statistics System

	2014	2015	2016	2017	2018	2019	2020
ANNUAL TARGETS	317	313	305	297	277	258	238
HCV-RELATED DEATHS AMONG AI/AN	317	367	312	299	264		

DATA SOURCES

The National Health and Nutrition Examination Survey (NHANES) is a CDC program designed to assess the health and nutritional status of adults and children in the United States. A survey and physical examination of a nationally representative sample of 5,000 persons each year collects demographic, socioeconomic, dietary, and health-related data as well as medical, dental, and physiological measurements and results of laboratory tests administered by medical personnel.

The National Health Interview Survey (NHIS) is an annual, cross-sectional in-person household survey collected by CDC's National Center for Health Statistics and conducted by interviewers trained by the U.S. Census Bureau. Data are used to monitor self-reported trends in illness and disability among the U.S. civilian noninstitutionalized population. NHIS provides adult vaccination coverage estimates.

The National Immunization Surveys (NIS) are a group of telephone surveys sponsored and conducted by CDC's National Center for Immunization and Respiratory Diseases. <u>NIS-Child</u> targets children in the United States who are or will be 19–35 months old within a few weeks of the survey. Annually collected data are used to monitor vaccination coverage among 2-year-old children, including the hepatitis B birth dose, at the national, state, and selected local levels, and in some U.S. territories.

The National Notifiable Diseases Surveillance System (NNDSS) is a CDC program managed by its Division of Health Informatics and Surveillance that collects, analyzes, and publishes health data for approximately 120 diseases. These data, which CDC collects annually, help public health officials monitor, control, and prevent disease in the United States.

The <u>National Vital Statistics System</u> (NVSS) is the mechanism by which CDC's National Center of Health Statistics (NCHS) annually collects and disseminates the nation's official vital statistics. These data are provided through contracts between NCHS and vital registration systems operated in the various jurisdictions legally responsible for the registration of vital events – births, deaths, marriages, divorces, and fetal deaths.

FEDERAL ACTIVITIES TO ADVANCE THE UNITED STATES TOWARD NATIONAL HEPATITIS GOALS

The Action Plan was developed collaboratively by partners from federal agencies with input from nonfederal stakeholders from a variety of sectors. These federal partners have implemented a wide range of viral hepatitis activities, from prevention to care to researching new viral hepatitis therapies. The activities are described in detail in Appendix 2.

DEPARTMENTS, AGENCIES, AND OFFICES THAT IMPLEMENTED THE ACTION PLAN IN FY2018 – FY2019

	Department of Health and Human Services (HHS)
Agency for Healthcare Research and Quality (AHRQ)	AHRQ's mission is to produce evidence to make health care safer, higher quality, more accessible, equitable, and affordable, and to work within the U.S. Department of Health and Human Services and with other partners to make sure that the evidence is understood and used.
Centers for Disease Control and Prevention (CDC)	
National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP)	NCCDPHP's mission is to help people and communities prevent chronic diseases and promote health and wellness for all.
National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP)	NCHHSTP maximizes public health and safety nationally and internationally through the elimination, prevention, and control of disease, disability, and death caused by HIV/AIDS, non-HIV retroviruses, viral hepatitis, other sexually transmitted diseases, and tuberculosis.
Division of Viral Hepatitis (DVH)	DVH's mission is to end the viral hepatitis epidemics through leadership in science and public health practices.
National Center for Immunization and Respiratory Diseases (NCIRD)	NCIRD's mission is the prevention of disease, disability, and death through immunization and by control of respiratory and related diseases.
Centers for Medicare & Medicaid Services (CMS)	CMS provides health coverage to more than 100 million people through Medicare, Medicaid, the Children's Health Insurance Program, and the private health insurance market including Health Insurance Exchanges. CMS seeks to strengthen and modernize the nation's health care system, to provide access to high quality care and improved health at lower costs.
Food and Drug Administration (FDA), Center for Drug Evaluation and Research (CDER)	CDER's mission is to protect and promote public health by helping to ensure that human drugs are safe and effective for their intended use, that they meet established quality standards, and that they are available to patients.



	Department of Health and Human Services (HHS)
Health Resources and Services Administration (HRSA), HIV/AIDS Bureau (HAB)	HRSA HAB's mission is to provide leadership and resources to assure access to and retention in high quality, integrated care and treatment services for vulnerable people with HIV/AIDS and their families.
Indian Health Service (IHS)	IHS is responsible for providing federal health services to American Indians and Alaska Natives. The mission of IHS is to raise the physical, mental, social, and spiritual health of American Indians and Alaska Natives to the highest level.
National Institutes of Health (NIH)	
Eunice Kennedy Shriver National Institute of Child Health and Human Development (NICHD)	NICHD's mission is to lead research and training to understand human development, improve reproductive health, enhance the lives of children and adolescents, and optimize abilities for all.
National Cancer Institute (NCI)	NCI leads, conducts, and supports cancer research across the nation to advance scientific knowledge and help all people live longer, healthier lives.
National Institute of Allergy and Infectious Diseases (NIAID)	NIAID's mission is to conduct and support basic and applied research to better understand, treat, and ultimately prevent infectious, immunologic, and allergic diseases.
National Institute of Diabetes and Digestive and Kidney Diseases (NIDDK)	The mission of NIDDK is to conduct and support medical research and research training and to disseminate science-based information on diabetes and other endocrine and metabolic diseases; digestive diseases, nutritional disorders, and obesity; and kidney, urologic, and hematologic diseases, to improve people's health and quality of life.
National Institute on Drug Abuse (NIDA)	The mission of NIDA is to advance science on the causes and consequences of drug use and addiction and to apply that knowledge to improve individual and public health.
Office of Intergovernmental and External Affairs (OIEA)	OIEA strengthens relationships between state and local partners and external stakeholders with the Office of the Secretary. OIEA also serves as liaison for governmental and non-governmental partners in communicating with Departmental offices and the Bureaus.
Office of the Assistant Secretary for Health (OASH)	
Office of Disease Prevention and Health Promotion (ODPHP)	ODPHP provides leadership for disease prevention and health promotion efforts for all Americans. To promote the health of the country, ODPHP sets national health goals and supports programs, services, and educational activities. ODPHP leads Healthy People 2020/2030, Dietary Guidelines for Americans, Physical Activity Guidelines for Americans, National Clinical Care Commission, National Youth Sports Strategy, President's Council on Sports, Fitness and Nutrition, and healthfinder.gov.



	Department of Health and Human Services (HHS)
Office of Infectious Disease and HIV/AIDS Policy (OIDP)	OIDP provides strategic leadership and management to the Department, while encouraging collaboration, coordination, and innovation among federal agencies and stakeholders, to reduce the burden of infectious diseases. OIDP includes the previously separate Office of HIV/AIDS and Infectious Disease Policy (OHAIDP) and National Vaccine Program Office (NVPO), which were reorganized and combined as OIDP in June 2019.
Office of Population Affairs (OPA)	OPA promotes health across the reproductive lifespan through innovative, evidence-based adolescent health and family planning programs, services, strategic partnerships, evaluation, and research. OPA administers the <u>Title</u> <u>X family planning program, Teen Pregnancy Prevention program,</u> <u>Pregnancy Assistance Fund program, and embryo adoption program. OPA</u> advises the Secretary and the Assistant Secretary for Health on a wide range of topics, including adolescent health, family planning, sterilization, and other population issues.
Office of the Surgeon General (OSG)	As the nation's doctor, the Surgeon General provides Americans with the best scientific information available on how to improve their health and reduce their risk of illness and injury. The Surgeon General brings this information to the public by issuing Surgeon General's Advisories, Calls to Action, and Reports on critical issues and communicating directly with the public via a number of communication channels. As Vice Admiral of the U.S. Public Health Service Commissioned Corps, the Surgeon General oversees the operations of the <u>U.S. Public Health Service Commissioned Corps (USPHS)</u> , an elite group of over 6,000 uniformed officers whose mission is to protect, promote, and advance the health of our nation.
Office on Women's Health (OWH)	OWH provides national leadership and coordination to improve the health of women and girls through policy, education, and innovative programs.
Regional Health Administrators (RHA)	As the senior federal public health official and scientist in the region, each Regional Health Administrator performs essential functions for HHS in three major areas: prevention, preparedness, and agency-wide coordination. These functions support the work of OASH and the Department. There are ten OASH regions nationwide and ten RHAs.
Office of Minority Health (OMH)	OMH is dedicated to improving the health of racial and ethnic minority populations through the development of health policies and programs to help eliminate health disparities.
Office of the National Coordinator for Health Information Technology (ONC)	ONC is the principal federal entity charged with coordination of nationwide efforts to implement and use the most advanced health information technology and the electronic exchange of health information.
Substance Abuse and Mental Health Services Administration (SAMHSA)	SAMHSA leads public health efforts to advance the behavioral health of the nation and to improve the lives of individuals living with mental and substance use disorders, and their families.



	Department of Housing and Urban Development
Agency for Healthcare Research and Quality (AHRQ)	AHRQ's mission is to produce evidence to make health care safer, higher quality, more accessible, equitable, and affordable, and to work within the U.S. Department of Health and Human Services and with other partners to make sure that the evidence is understood and used.
	Department of Justice
Federal Bureau of Prisons (BOP)	The mission of the BOP is to protect society by confining offenders in the controlled environments of prisons and community-based facilities that are safe, humane, cost-efficient, and appropriately secure, and that provide work and other self-improvement opportunities to assist offenders in becoming law-abiding citizens.
Civil Rights Division (CRT)	CRT works to uphold the civil and constitutional rights of all Americans, particularly some of the most vulnerable members of our society. The Division enforces federal statutes prohibiting discrimination on the basis of race, color, sex, disability, religion, familial status and national origin.
	Department of Veteran Affairs
Veterans Health Administration's (VHA) Office of Patient Care Services (PCS)	PCS is dedicated to ensuring the full continuum of health care, comprised of health promotion, disease prevention, diagnostics, therapeutic and rehabilitative care, recovery and palliative care. PCS provides care through policy and program development that promotes dignity and respect, and is achieved by utilizing innovative approaches and technologies through interdisciplinary collaboration both within and outside of VHA.



TIMELINE: SELECTED HIGHLIGHTS (BASED ON FISCAL YEARS)



APPENDIX 1: FEDERAL PROGRESS ON ACTION PLAN

This appendix documents contributions made by federal partners during FY2018 and FY2019 on strategies detailed in the <u>National Viral Hepatitis Action Plan 2017–2020</u> (Action Plan). It does not provide a complete summary of all the actions federal agencies have taken that are related to Action Plan goals. For example, some agencies initiated new programs that were not anticipated at the time the Action Plan was updated and therefore do not correlate with a specific Action Plan strategy.

The following information is organized by Action Plan goal and strategy. The information presented here provides only a brief snapshot of the effort that went into these actions.

GOAL 1: PREVENT NEW VIRAL HEPATITIS INFECTIONS

Agency	Activity
	Strategy 1.1: Increase community awareness of viral hepatitis and decrease stigma and discrimination
CDC	The Cherokee Nation Comprehensive Cancer Control Program collaborated with the Cherokee Nation HCV Elimination Program within Cherokee Nation's Health Services (CNHS) to increase knowledge and awareness of liver cancer prevention among the Cherokee Nation community by conducting presentations to 26 community coalition organizations.
DOJ	The Disability Rights Section of the Civil Rights Division continues to receive and review referrals of potential hepatitis-based discrimination through direct calls and online at <u>http://www.ada.gov</u>
SAMHSA	To reduce the impact of substance use, HIV, and viral hepatitis in high-risk communities, SAMHSA funds the HIV Capacity Building Initiative and the HIV Prevention Navigator grant programs to:
	 provide HIV and viral hepatitis testing services in non-traditional settings; develop strategies that combine education and awareness programs; and produce social marketing campaigns with substance abuse and HIV prevention programming for the population of focus.
	Strategy 1.2: Build capacity and support innovation by the health care workforce to prevent viral hepatitis
HRSA	HRSA HIV/AIDS Bureau (HAB) developed an online Hepatitis C Prescriber Toolkit which provides state-specific guidance, current prescribing restrictions, and resources for additional assistance to prescribers of Hepatitis C medications.
	Strategy 1.3: Address critical data gaps and improve viral hepatitis surveillance
CDC	Surveillance provides data for action to control the spread of disease. CDC and health departments use surveillance data to detect viral hepatitis outbreaks; quantify, characterize, and monitor trends in new infections, burden of disease, and transmission risk factors; and identify opportunities to link individuals to viral hepatitis preventive and treatment services—all of which are vital to develop and evaluate prevention and control strategies. Health departments must have systems in place to detect, classify, and notify CDC of viral hepatitis cases; however, due to varying infrastructure, not all health departments notify CDC of all cases of acute or newly reported chronic viral hepatitis. Through the <u>Strengthening Surveillance in Jurisdictions with High Incidence of Hepatitis C Virus and Hepatitis B Virus Infections</u> cooperative agreement (CDC-RFA-PS17-1703), CDC funded 14 states experiencing high rates of acute cases of hepatitis B and/or hepatitis C infections to improve active surveillance, data completeness, and case notification to CDC. In 2018, awardees achieved 48.5% and 30% increases in risk factor completeness for hepatitis B and hepatitis C case notifications submitted to CDC, respectively.
	Strategy 1.4: Achieve universal hepatitis A and hepatitis B vaccination for children and vulnerable adults
VA	In April 2019, VA issued system-wide a memo on hepatitis A and hepatitis B immunization in homeless Veterans. The memo outlined the CDC recommendations, provided clinical tools and resources, and advised facility viral hepatitis lead clinicians, homeless program providers, health promotion and disease prevention program managers and facility health behavior coordinators to coordinate efforts to improve vaccination rates.



National Viral Hepatitis Action Plan, 2017–2020

Agency	Activity
	Strategy 1.5: Eliminate mother-to-child transmission of hepatitis B and hepatitis C
AHRQ	The U.S. Preventive Services Task Force (USPSTF) released Screening for Hepatitis B Virus Infection in Pregnant Women: US Preventive Services Task Force Reaffirmation Recommendation Statement in July 2019 <u>https://www.uspreventiveservicestaskforce.org/Page/Document/UpdateSummaryFinal/hepatitis-b-virus-infection-in-pregnant-women-screening?ds=1&s=hepatits</u>
FDA	FDA's Center for Drug Evaluation and Research (CDER) approved supplemental application for VIREAD (tenofovir disoproxil fumarate) to update product labeling to include safety and pregnancy-related outcome information from three published controlled trials in pregnant women with chronic hepatitis B virus infection who were administered VIREAD during their third trimester.
	Strategy 1.6: Ensure that people who inject drugs have access to viral hepatitis prevention services
CDC	In FY 2019, CDC began a new three-year cooperative agreement National Harm Reduction Technical Assistance and Syringe Services Program (SSP) Monitoring and Evaluation Funding Opportunity. This program aims to strengthen the capacity and improve the performance of SSPs throughout the United States by supporting enhanced technical assistance to ensure the provision of high-quality, comprehensive harm reduction services; implementing a national SSP monitoring and evaluation program; and supporting the development and implementation of best practices for patient navigation from SSPs to community-based health and social services. Activities under this program also seek to increase the capacity to understand injection drug use and risk in the United States with an injection drug use surveillance demonstration project.
CDC	CDC funded nine jurisdictions to test and link people to care in high-impact settings, such as SSPs, substance use disorder treatment facilities, emergency departments, and correction facilities.
IHS	 While there are many resources available to the public on harm reduction, they are scattered or not tribal-specific. To ensure that the tribes are not only aware of current and promising harm reduction practices and strategies for opioid response, both regionally and nationally, the Northwest Portland Area Indian Health Board Indian Country Opioid Response Monthly Newsletter and Community of Learning webinar serieswere developed. The goal of these two tools is to use them as a way to cultivate a community of practice and to disseminate the strategies and promising practices currently being implemented to address injection drug use and substance use disorder across Indian Country. More at https://www.indiancountryecho.org/substance-use-disorder/community-resources. As of September 2019, 350 are signed up for the Opioid Newsletter. In 2019, the project developed a number of campaigns for communities. This included electronic and print material for several new resources including "A Trickster Tale – Outsmarting Through Education and Action",
	"Words Matter When Providers Talk About Addiction", "Words Matter When Providers Talk About Addiction -For Patients", and "Supporting Someone with Opioid Addiction", among others. See <u>https://www.indiancountryecho.org/substance-use-disorder/community-resources</u> .
	Strategy 1.8: Conduct research leading to new or improved viral hepatitis vaccines, diagnostic tests, and treatments, and the optimal use of existing tools to prevent, detect, and treat viral hepatitis
NIH	NIAID has conducted a double-blind, randomized, Phase II trial on the safety, immunogenicity and efficacy of an innovative, experimental hepatitis C vaccine designed to stimulate T cell reactivity to the non-structural proteins of the hepatitis C virus (https://www.clinicaltrials.gov/ct2/show/NCT01436357). A total of 548 adults (18-45 years old) at high risk of HCV infection because of active injection drug use received the "prime-boost" regimen of hepatitis C vaccine or similar-appearing placebo injections. They were followed carefully for side effects, immune responses, and evidence of HCV infection. Chronic HCV infection developed in 7 percent of both groups but the vaccine was safe, led to vigorous T cell responses to HCV, and was associated with lower levels of HCV RNA in infected persons. Thus, while this innovative vaccine did not provide protection against HCV, it was safe and yielded excellent T cell immune responses suggesting that combination with B cell vaccines or means of improving innate immunity might ultimately provide an effective vaccine against this important cause of liver disease.
NIH	The National Heart, Lung, and Blood Institute (NHLBI) has collaborated with FDA on the Transfusion- Transmissible Infections Monitoring System (TTIMS) to monitor the residual risk of HBV and HCV as well as HIV among blood donations and the safety of the blood supply in the U.S. No major trends were observed over 4 years covering the MSM policy change from indefinite to a 12-month deferral, but ongoing monitoring is indicated.

GOAL 2: REDUCE DEATHS AND IMPROVE THE HEALTH OF PEOPLE LIVING WITH VIRAL HEPATITIS

Agency	Activity
	Strategy 2.1: Build the capacity of the health care workforce to diagnose viral hepatitis and provide care and treatment to persons living with chronic viral hepatitis
CDC	Training of Healthcare Providers
	As part of its efforts to build the capacity of the health care workforce to diagnose and treat viral hepatitis, CDC supports the development of up-to-date, comprehensive, web-based, hepatitis materials, resources, and trainings for health professionals. The University of Washington National Hepatitis Training Center, through CDC's <u>Viral Hepatitis Networking, Capacity Building, and Training</u> cooperative agreement (CDC-RFA- PS16-1608), has developed Hepatitis C Online. <u>Hepatitis C Online</u> is a free, self-study, interactive course on hepatitis C virus infection for medical providers. The comprehensive training addresses the diagnosis, monitoring and management of hepatitis C and includes dedicated sections for HCV medications, clinical calculators (APRI, MELD, Glasgow Coma Scale, etc.), and master bibliography. Free CME credit and free CNE credit are offered to clinicians who complete the course. Between September 2018 and October 2019, Hepatitis C Online had 618,486 total users who initiated at least one session of the course.
CDC	The Cherokee Nation's HCV Elimination Program was responsible for conducting didactic sessions for CNHS health care providers on HCC epidemiology, diagnosis, and surveillance through the Project Extension for Community Healthcare Outcomes (ECHO) platform (the Echo Model) and through conducting health care provider education workshops focused on liver cancer at 8 CNHS facilities.
HRSA	HRSA staff continues to monitor potential HIV/HCV outbreaks and provide technical assistance on a case by case basis to enhance coordination and the availability of federal, state, and community resources to provide linkage to HIV/hepatitis care and prevention services. Through the RWHAP AIDS Education and Training Centers, training on HIV and HCV testing, outreach, and case monitoring is provided.
IHS	To improve the capacity of treatment of HCV at the primary care level, IHS and tribal partners made extensive use of the ECHO (Extension for Community Health Outcomes) model of telehealth and teleconsultation University of California, San Francisco NCCC (National Clinical Consultation Center). The Northwest Portland Area Indian Health Board (NPAIHB) has provided recommendations via the Indian Country ECHO (<u>www.indiancountryecho.org</u>) for over 700 patients across the United States with the highest case presentations coming from Washington, Oregon, Montana, South Dakota, North Dakota and Minnesota. The Indian Country ECHO website launched July 11, 2019. In September of 2019, the Indian Country ECHO website received: Users = 182, Sessions = 312, Page views = 961, Pages/Session = 3.08, Average session
	duration = 4:43, Bounce Rate = 39.74%. IHS, tribal, and urban Indian health clinics from all 12 IHS Areas participated in the NPAIHB ECHO. Fifty-six
	unique clinics used the NCCC teleconsultation services.



Agency	Activity
ОМН	Hepatitis B Demonstration Summary Document
	OMH in consultation with the OIDP and CDC's Division of Viral Hepatitis, funded development of model comprehensive hepatitis B programs that included strategic partnerships between: community-based organizations servicing communities at risk; departments of health, perinatal hepatitis B programs; safety net providers, research centers, and healthcare facilities that have capacity to deliver widespread vaccination; and scale-up testing, care and link/provide treatment services.
	These hepatitis B programs will advance progress toward the national hepatitis B elimination goals and strategic actions recommended by National Academies of Sciences, Engineering, and Medicine (NASEM): end transmission of hepatitis B virus (HBV) (perinatal, children and adults); and reduce morbidity and mortality attributable to ongoing HBV infection.
	The program is directly aligned with the goals of the National Viral Hepatitis Action Plan. Grantees were required to deliver widespread vaccination, scale-up testing, and provide care and treatment services.
	OMH will support model programs to implement all of the following strategic actions: prevent new HBV cases, reduce deaths, and improve the health of people living with viral hepatitis; provide the birth dose coverage (vaccination); provide maternal HBsAg testing followed by HBV DNA testing as appropriate; treat HBV-exposed newborns; provide community-based testing and linkage to care; provide people who are HBV-positive linkage to care, treatment, and contact tracing; vaccinate people who are HBV-negative; increase availability of testing kits and provide education to health care providers and patients; utilize care coordinators for linkage to care; and implement standing orders or utilization of Electronic Health Records (EHR) for testing and linkage to care.
ОМН	OMH contracted with MayaTech in FY2018 to implement the Viral Hepatitis Program-Primary Care Physicians Capacity Building Initiative . The major objectives of this initiative were to support training for physicians, nurses, pharmacists, and other health workers, and enhance capacity building in primary care settings. Of particular interest was a focus on screening guidelines, hepatitis vaccinations guidelines, and guidelines for management and treatment of individuals infected with HBV and/or HCV. Additional objectives were to establish standard operating practices (e.g., implementation of provider prompts, data-driven chart reviews to identify patients tested/diagnosed but not currently in care, and screening and vaccination coverage assessments) and create hepatitis comprehensive care models that facilitate collaboration between primary care providers and hepatitis specialists. Three sites accepted the invitation for and participated in trainings: AIDS Care Group, Philadelphia, PA; Housing Works, New York, NY; and, Mattapan Community Health Center, Boston, MA.
VA	In 2018, the Hepatic Innovation Team (HIT) Collaboratives were transitioned to focus on addressing advanced liver disease (ALD). These teams picked up on the highly successful work accomplished by the Hepatitis Innovation Team Collaborative which focused on testing and treatment of HCV in VA. The HIT Collaboratives focus on education and communication to improve and support provider practice, Veterans' health and engagement in care. They employ population-based approaches with a focus on quality improvement initiatives to address gaps in care and anticipate system-wide needs. The HIT Collaboratives consist of multi-disciplinary teams across each Veteran Integrated Service Network (VISN).
	Strategy 2.2: Identify persons infected with viral hepatitis early in the course of their disease
BOP	The BOP has adopted an "opt-out" policy for hepatitis C screening. In this model, when inmates are processed during intake, the policy is to obtain bloodwork to screen for hepatitis C as a routine part of the process unless the patient declines, or "opts out". This is in contrast to prior policy where the inmate was offered hepatitis C screening as an option.
AHRQ	The USPSTF is currently updating its HBV recommendation statement. The USPSTF posted a draft research plan, <i>Hepatitis B Virus Infection in Nonpregnant Adolescents and Adults: Screening</i> , for public comment. The final research plan is found at <u>https://www.uspreventiveservicestaskforce.org/Page/Document/final-</u> <u>research-plan/hepatitis-b-virus-infection-screening-nonpregnant</u> .



2018-2019 PROGRESS REPORT

National Viral Hepatitis Action Plan, 2017–2020

Agency	Activity
CDC	Prevention - Testing for hepatitis C and hepatitis B, when linked to care and treatment, is cost-effective and improves health outcomes. However, only about half of the estimated 2.4 million people living with hepatitis C are aware of their infection and most have not received recommended care and treatment. Similarly, only about one-third of the estimated 862,000 people living with hepatitis B in the United States are aware of their infection. In FY2017, CDC awarded \$5.7M over a four-year project period (2017 –2020) to state and local health departments in 46 U.S. states, three cities, and the District of Columbia through the Improving Hepatitis B and C Care Cascades: Focus on Increased Testing and Diagnosis cooperative agreement (CDC-RFA-PS17-1702). These resources support activities to increase the number of persons living with HBV and/or HCV infection who are tested for these infections, made aware of their infection status, and promoted linkage to care and treatment services, if needed. In FY2018, 100% of funded jurisdictions completed a situational analysis, a foundational step to assist jurisdictions to achieve the greatest impact with this investment.
SAMHSA	SAMHSA has two Minority AIDS Initiative (MAI) programs with 5% of each grant must be used to address vira hepatitis:
	1. 37 HIV Capacity Building Initiative – FY18 (N=37)
	2. 6 HIV Prevention Navigator – FY19 (N=6)
	These have increased viral hepatitis testing with:
	 Total viral hepatitis test kits purchased with MAI funds (as of 1/23/2020): 110,177
	 Total tested for viral hepatitis with MAI funds (as of 1/23/2020): 34,558
VA	VA birth cohort screening rates were 84.8% in 2018 and 84.9% in 2019.
	Strategy 2.3: Improve access to and quality of care and treatment for persons infected with viral hepatitis
CMS	CMS approved a Washington State Plan Amendment proposal that permits the state to negotiate supplemental rebate agreements involving value-based purchasing arrangements with drug manufacturers that links payment for prescription drugs to the value delivered. Washington's proposal was specifically designed to allow the state to negotiate under a subscription model with manufacturers of prescription drugs that treat patients with the hepatitis C. Under this subscription model, the state would pay a fixed annual amount to a pharmaceutical manufacturer to purchase an unrestricted supply of hepatitis C drugs. This innovative proposal demands value from pharmaceutical companies and takes steps to eradicate hepatitis C in Washington state.
FDA	FDA's CDER approved supplemental application for VIREAD (tenofovir disoproxil fumarate) to expand the treatment of chronic hepatitis B indication to include pediatric patients 2 years and older weighing at least 10 kg. This fulfilled post-marketing requirements under the Pediatric Research Equity Act (PREA).
FDA	FDA's CDER approved supplemental application for VEMLIDY (tenofovir alafenamide) to update product labeling with data in adult patients with creatinine clearance below 15 mL per minute (end-stage renal disease) who are receiving chronic hemodialysis.
FDA	FDA's CDER approved supplemental application for Mavyret (glecaprevir and pibrentasvir) to expand the indication to adolescents 12 years and older or weighing at least 45 kilograms (kg). This approval provides a shorter duration treatment option in adolescents for six major genotypes of HCV.
VA	As of October 1, 2018, there were approximately 9,000 Veterans with cirrhosis awaiting hepatitis C treatmen in VA and approximately 6,800 remaining to be treated by September 30, 2019.
VA	As of October 1, 2018, there nearly 33,000 Veterans with hepatitis C waiting on treatment and approximately nearly 25,000 remaining to be treated by September 30, 2019.
VA	As of September 30, 2019, 71.2% of Veterans in care with chronic hepatitis B were on treatment.
CMS	CMS approved a Louisiana State Plan Amendment proposal for Supplemental Rebate Agreements using a modified subscription model for hepatitis C therapies in Medicaid. Louisiana's proposal permits the state to negotiate supplemental drug rebates to assist the state in controlling expenditures for hepatitis C drugs while providing unlimited access to the therapies. This modified subscription model initially focuses on antiviral agents for hepatitis C and promotes eliminating the hepatitis C virus statewide. This supplemental rebate agreement will be used by the state for a modified subscription model, allowing Louisiana to cap gross expenditures at a fixed amount for hepatitis C drugs, while providing the state with unlimited access to clinically necessary doses of these therapies for Medicaid beneficiaries.



National Viral Hepatitis Action Plan, 2017–2020

Agency	Activity
FDA	FDA's CDER approved supplemental applications for Sovaldi (sofosbuvir) and Harvoni (ledipasvir and sofosbuvir) to treat chronic HCV infection in children 3 years of age and older, weighing at least 17 kg. Harvoni and Sovaldi were previously approved to treat HCV in adults and children aged 12 to 17 years. These direct-acting antiviral treatments address an unmet need in younger children.
FDA	FDA's Center for Drug Evaluation and Research approved supplemental application for Mavyret (glecaprevir and pibrentasvir) to support an 8-week dosing regimen for the treatment of genotypes 1, 2, 3, 4, 5, and 6, chronic HCV infection in treatment-naïve subjects with compensated cirrhosis. Mavyret is the first treatment of eight weeks duration approved for HCV genotypes 1-6 in adult patients without cirrhosis and with compensated cirrhosis who have not been previously treated. This approval provides a shorter treatment duration for many patients.
	Strategy 2.4: Improve viral hepatitis treatment among persons with HIV
ВОР	While the goal of the BOP is to treat all hepatitis C-infected patients, the order of treatment is prioritized based on a number of factors. One of the criteria for prioritized treatment for hepatitis C is HIV coinfection.
FDA	FDA's CDER approved supplemental application for Mavyret (glecaprevir and pibrentasvir) to include safety and efficacy data in adult patients with HCV/HIV-1 coinfection.
HRSA	HRSA-funded RWHAP AIDS Education and Training Centers (AETC) continued to provide clinical training and clinical consultation on the screening and treatment of HBV and HCV for coinfected people living with HIV. As part of this effort, the AETC National Coordinating Resource Center developed and published a free, online e-Learning platform that offers healthcare providers and health profession educators training on HIV/HCV coinfection, including prevention, screening, diagnosis, and treatment recommendations. The curriculum also examines barriers and other factors that may impede optimal treatment outcomes for coinfected people of color.
HRSA	HRSA's Curing Hepatitis C among People of Color Living with HIV, funded through the 2017 Secretary's Minority AIDS Initiative Fund, awarded two academic centers for a three-year project intending to improve care for people with HIV through the development of comprehensive jurisdiction-level HCV screening, care, and treatment systems and the enhancement of state and local health department surveillance systems to increase capacity to monitor acute and chronic coinfections of HIV and HCV. Additionally, this project focused on improving coordination with SAMHSA-funded Substance Use Disorder (SUD) treatment providers to deliver behavioral health and SUD treatment support to achieve treatment completion, prevent HCV infection, and re- infection.
VA	As of October 1, 2018, there were nearly 3,000 Veterans with HIV waiting on HCV treatment, and approximately 600 remaining to be treated by September 30, 2019.
	Strategy 2.5: Ensure that people who inject drugs have access to viral hepatitis care and evidence-based treatment services
FDA	FDA's CDER initiated revisions to the Prescribing Information for HCV drugs to include a subsection on Medication-Assisted Treatment (MAT) for Opioid Use Disorder, to highlight drug interaction information.
SAMHSA	SAMHSA's Substance Abuse Treatment Block Grant (SABG) program's objective is to help plan, implement, and evaluate activities that prevent and treat substance abuse. The SABG program targets the following populations and service areas: Pregnant women and women with dependent children, intravenous drug users, early intervention services for HIV/AIDS and viral hepatitis, and primary prevention services. Injection drug use (IDU) risk reduction is best approached in a step-wise fashion; for example, abstinence from illicit drug use is the best way to address a person's health and diminish the chance of becoming infected with HIV or viral hepatitis. For those with OUD the best way of stopping IDU is to provide MAT and psychosocial services in a SUD program. Syringe service programs (SSPs) represent an opportunity to try to engage people in treatment and short of that educating on reducing HIV risk through not sharing syringes and using only sterile syringes if they are going to inject. SAMHSA resources are used to support SSPs.
	Strategy 2.6: Expand access to and delivery of hepatitis prevention, care, and treatment services in correctional settings



2018-2019 PROGRESS REPORT

National Viral Hepatitis Action Plan, 2017–2020

Agency	Activity
BOP	The Federal Bureau of Prisons offers HCV treatment to all sentenced inmates with sufficient time remaining to treat and no major obstacles to treatment such as continued high-risk behavior (decided on a case-by-case basis). Treatment numbers have substantially increased over time: almost triple from FY2016 (298 patients) to FY2017 (874 patients), >1.5 times from FY2017 to FY2018 (1,421 patients), and more than double from FY2018 to FY2019 (>3,000 inmates). These significant leaps in treatment numbers are due to multiple factors, including: 1) protocol where \$25-30 million is set aside per fiscal year to reimburse individual institutions for HCV treatment, thus removing financial burden as one of the biggest hurdles to treatment; 2) continued use of the Regional Hepatitis Clinical Pharmacist Consultant Program to approve non-formulary requests and serve as subject matter experts for the field; and 3) continued focus on hepatitis C care by BOP medical leadership leading to increased exposure and comfort level of institution providers.
	Strategy 2.8: Advance research to enhance identification, care, treatment, and cure for persons infected with viral hepatitis
NIH	The NIDDK's Hepatitis B Research Network has completed two multicenter studies—one in adults and another in children—of the combination of interferon alfa (an immune cytokine) and entecavir (an oral antiviral agent) in immune tolerant chronic hepatitis B, a common, but resistant form of this infection (https://www.ncbi.nlm.nih.gov/pubmed/30549279; https://www.ncbi.nlm.nih.gov/pubmed/30318613). Therapy was given for one year. While the treatment was well-tolerated, response rates were low. No adult had a sustained response, but 5 percent of children had a dramatic response, becoming negative for hepatitis B antigen and developing antibody, suggesting a "functional cure." These results indicate that treatment will likely require the combination of three or more agents against HBV, directed at different viral "targets," but that full recovery from chronic hepatitis B may be possible even in the most difficult-to-treat instances.
NIH	An international trial of prevention of mother-to-child transmission of hepatitis B funded by <i>Eunice Kennedy</i> <i>Shriver</i> National Institute of Child Health and Human Development (NICHD) has been completed (https://www.nejm.org/doi/full/10.1056/NEJMoa1708131). A total of 331 pregnant women with high levels of serum HBV were randomly assigned to receive tenofovir (an oral antiviral agent) or placebo during the last trimester of pregnancy. All babies born to the mothers were given the usual regimen to prevent infection: hepatitis B vaccine and hepatitis B immune globulin. In follow up, none of the 147 newborns of mothers treated with tenofovir but 3 of the 147 infants born to mothers given placebo developed hepatitis B during the first year of life. The treatment was safe and did not cause significant side effects or worsening of the mother's HBV infection. These results and those from similar trials indicate that use of antiviral therapy in mothers at high risk of transmitting HBV to their newborns can decrease or eliminate the risk of transmission. Guidelines supporting this approach have now been published by academic societies in the United States and elsewhere.
NIH	The NCI has established a multicenter U.S. Liver Cancer Consortium, which is charged with developing a large clinical network to conduct advanced translational research on the early detection, diagnosis, clinical management, prevention and treatment of liver cancer in patients with chronic liver disease who are at high risk for this highly fatal malignancy (<u>https://grants.nih.gov/grants/guide/rfa-files/RFA-CA-17-025.html</u>). The Consortium will bring together clinical and basic research expertise and apply state-of-the-art investigational techniques aimed at identifying biomarkers for early detection and diagnosis of liver cancer, as well as insights into personalized medical approach to its treatment.

GOAL 3: REDUCE VIRAL HEPATITIS HEALTH DISPARITIES

 Agency
 Activity

 Strategy 3.1: Decrease health disparities by partnering with and educating priority populations and their communities about viral hepatitis and the benefits of available prevention, care, and treatment



National Viral Hepatitis Action Plan, 2017–2020

Agency	Activity
CDC	Communications Campaign and Outreach Activities for Hepatitis B Awareness
	CDC is committed to leveraging the successes and best practices of its partners to help improve hepatitis B screening in the highest-risk and most underserved Asian American and Pacific Islander communities in the United States. In 2013, CDC launched Know Hepatitis B, a national communications campaign promoting hepatitis B testing among Asian Americans. This multilingual campaign is delivered through a variety of multi-media channels. To further expand the reach of the Know Hepatitis B campaign, CDC's <u>Viral Hepatitis</u> <u>Networking, Capacity Building, and Training</u> cooperative agreement (CDC-RFA- PS16-1608) funds a partner to provide capacity building, training, and technical assistance to over 30 hepatitis B coalition partners in 28 cities and 20 states, including the District of Columbia. Activities include coalition building, webinars, mini grants, peer mentoring, a national hepatitis B storytelling campaign, and in-person annual summits. Since its inception, the Know Hepatitis B campaign has achieved at least 474 million impressions worth \$3.8 million of paid media value.
CDC	CDC Response to Widespread Hepatitis A Outbreaks
	More than half of states across the country have reported outbreaks of hepatitis A. Since first identified in 2016, over 27,000 cases with approximately 60% hospitalizations and at least 275 deaths have been reported (as of November 2019). In these outbreaks, the virus is being spread person-to-person primarily among people who use drugs and people who are experiencing homelessness. Vaccinating people at risk can stop the spread. Since March 2017, CDC has been supporting state and local health departments by providing national situational awareness and ongoing remote and on-site technical assistance with outbreak response and prevention. CDC has deployed experts representing a mix of expertise from CDC's viral hepatitis, vaccine, and preparedness programs, including epidemiologists, laboratorians, public health advisors, and disease intervention specialists, to eight states to support their outbreak responses. CDC's laboratory has processed more than 5,000 hepatitis A virus specimens since the outbreaks began and continues to support vaccine supply and vaccine policy development. CDC shares its best practices through ongoing engagement and communication with impacted states and health departments nationwide. Furthermore, CDC has launched an <u>outbreak-specific website</u> to provide all stakeholders and the public with up-to-date information about the outbreaks, educational resources, and links to useful guidance documents.
HUD	HUD's Office of Special Needs Assistance Programs outreached to and provided support for homeless assistance providers and Continuums of Care located in communities impacted by hepatitis A outbreaks. One key component of these engagements is facilitating partnerships between homeless assistance providers, public health departments and health care providers to improve access to and quality of care and treatment for persons experiencing homelessness who are infected with hepatitis A.
IHS	The Northwest Portland Area Indian Health Board has created a National HCV Social Marketing Campaign with the focus of relaying the message, "Hepatitis C is everybody's responsibility." The HCV Print & Video Campaign can be found at http://www.npaihb.org/hcv/#Community-Resources . Through FY2019, 10,000 items (posters, rack cards, pamphlets) have been printed and mailed out to IHS, tribal and urban Indian clinics. The video has received 944 video views on YouTube, and reached 5,515 on Facebook. Through FY2019, the project has sent 18,444 and received 1,976 messages from 432 text message subscribers. The project sent four marketing emails and had a reach of 1,754 through constant contact in the month of September. (Text HCV to 97779)
SAMHSA	The Assistant Secretary for Mental Health and Substance Abuse, Dr. Elinore McCance-Katz, released a Dear Colleague letter urging partnership with providers to increase testing and referrals for HIV and viral hepatitis within the priority populations of those with mental illness and substance use disorder. (https://www.samhsa.gov/sites/default/files/hivhepadvisory_letter-signed-508.pdf).
VA	In FY18-19, the HIV, Hepatitis, and Related Conditions Program (HHRC) increased awareness around viral hepatitis (hepatitis A, B, and C) in several ways. A primary source for awareness and information sharing is the website, <u>www.hepatitis.va.gov</u> . HHRC created and revised patient and provider education materials for the website throughout the fiscal year. This fiscal year HHRC had a major revision of the website which made it easier for users to find information on hepatitis A, hepatitis B, hepatitis C, and liver disease such as cirrhosis. For hepatitis testing day and hepatitis awareness month, HHRC conducted targeted outreach activities. This included emails and trainings for HHRC providers, features on the website and in facilities, and blog posts and social media posts. In addition to outreach and promotion of marking over 100,000 Veterans cured of hepatitis C, HHRC's work this year also focused on increasing hepatitis A and hepatitis B vaccination.



Agency	Activity
IHS	As part of IHS efforts to improve access to viral hepatitis treatment, two key policies were enacted in 2019. First, HCV DAAs were added to the IHS National Core Formulary. Second, based on cost-effectiveness modeling, IHS issued a Special General Memorandum to widen HCV screening to all persons 18 years and older. In addition to changes made at the IHS, our tribal partners have engaged with state Medicaid programs in Oregon, Arizona, South Dakota and Montana to help reduce restrictions on access to HCV medications.
	Strategy 3.3: Monitor viral hepatitis-associated health disparities in transmission, disease, and deaths
AHRQ	AHRQ published Characteristics of Inpatient Stays Involving Hepatitis C, 2005-2014. The statistical brief used Healthcare Cost and Utilization Project (HCUP) data to describe trends in the number and rate of hepatitis C- related inpatient hospital stays among adults over 18 years of age, both with and without the co-morbidities of HBV infection, HIV, and alcoholic liver disease.

GOAL 4: COORDINATE, MONITOR, AND REPORT ON IMPLEMENTATION OF VIRAL HEPATITIS ACTIVITIES

Agency	Activity
	Strategy 4.1: Increase coordination of viral hepatitis programs across the federal government and among federal agencies; state, territorial, tribal, and local governments; as well as non-governmental stakeholders from all sectors of society
CDC	In July 2019, CDC, in partnership with the Association of State and Territorial Health Officials (ASTHO), hosted the National Viral Hepatitis Program Planning Meeting to provide an opportunity for a national dialogue on viral hepatitis program planning to move forward on the path to elimination. As the first formal national elimination planning conference in a decade, the two-day meeting featured presentations and discussions on innovative drug pricing models for hepatitis C treatment, improving prevention and surveillance, decreasing barriers to treatment, and engaging people affected by viral hepatitis in elimination planning. Over 300 attendees from 48 states, 5 local jurisdictions, multiple federal agencies, and partner organizations attended. Additionally, states that
	received funding from ASTHO to implement hepatitis elimination projects were invited to stay for an additional day to learn about elimination strategies used by other jurisdictions and to discuss their own state-specific issues with partners.
FDA	A multidisciplinary working group in CDER is developing a guidance document for the development of drugs to treat chronic hepatitis D.
FDA	CDER's Division of Antiviral Products is engaged in discussions with various stakeholders through the HBV Forum (Forum for collaborative Research), and professional societies such as the American Association for the Study of Liver Diseases (AASLD) to enhance development of novel therapies for treatment of chronic hepatitis B.
FDA	CDER's Division of Antiviral Products in collaborative partnership with Hepatitis C Therapeutic Registry and Research Network (HCV-TARGET) is engaged in using real-world evidence to assess the safety and effectiveness of direct acting antivirals approved for the treatment of chronic hepatitis C.
FDA	CDER issued a draft guidance document for the development of drugs to treat chronic hepatitis B.
OIDP	Updating the Viral Hepatitis National Strategic Plan 2021 – 2025
	OIDP led the process to update the National Viral Hepatitis Action Plan 2017 – 2020, for 2021-2025. The new, updated Viral Hepatitis National Strategic Plan 2021-2025 (Hepatitis Plan) will be grounded in the latest science to guide stakeholders at all levels and sectors in key strategies to achieve updated national viral hepatitis goals. The Hepatitis Plan has been developed in alignment with developing the next HIV/AIDS National Strategic Plan as well as the first-ever Sexually Transmitted Infections (STI) National Strategic Plan.
	OIDP convened and collaborated with leadership from many federal departments and agencies to compile the best available evidence and recommendations for the Hepatitis Plan. This federal steering committee was informed by subcommittees (Prevention and Care, Disparities and Coordination, and Indicators) staffed by subject matter experts from throughout the federal government. The development of the plan was also informed by input from a wide variety of stakeholders and the public. Eighteen listening sessions were held across the nation (September 2018 - March 2019), and a <u>Request for Information</u> in the Federal Register solicited written public comments.
	This public input, federal leadership and subject matter expertise will help ensure that the Hepatitis Plan responds to pressing challenges in viral hepatitis, focuses on the most effective and scalable actions, responds to the needs of disproportionately affected communities and populations, and Is based on the latest scientific

evidence regarding viral hepatitis prevention, care, and treatment.



Agency	Activity
OIDP	Integrating infectious disease with opioid and drug misuse prevention
	In response to the syndemic of opioid and viral hepatitis epidemics, OIDP partnered to support expansion and improvement of integrated prevention programs through the following activities:
	National Academies of Science, Engineering, and Medicine (NASEM) Workshop and Evaluation Project
	 Integrating Infectious Disease Considerations with Response to the Opioid Epidemic: A Workshop
	 OIDP and the Office on Women's Health (OWH) cosponsored a one and a half day NASEM workshop on the infectious disease consequences of the opioid epidemic. The workshop aimed to explore the intersections and opportunities to improve our collective response to both problems. Presentations highlighted the scope and strategies to reduce the infectious disease consequences, especially those that emphasize empathy, respectful treatment, and patient satisfaction. A recording of the workshop sessions is <u>available to view</u>.
	 Examination of the Integration of Opioid and Infectious Disease Prevention Efforts in Select Programs To inform future projects and promote integrated programs, OIDP sponsored NASEM to convene an ad hoc committee to conduct a review and assess the extent to which select opioid and infectious disease prevention programs are integrating the services they provide. The committee highlighted programs that are achieving integration, barriers to integration, and related strategies to address these barriers. Syringe Services (SSPs) Webinar Series: to highlight the growing support from HHS, the Regional Health Administrators and OIDP developed and hosted a series of webinars.
	 Syringe Services Programs: A Critical Public Health Intervention (July 30, 2019) Featuring Assistant Secretary for Health ADM Brett Giroir and leadership from CDC, HRSA, and SAMHSA highlighting their activities and support for SSPs.
	• Syringe Services Programs: State and Local Perspectives on the Role of Policy, Funding, and Partnerships (September 16, 2019)
	 Featuring examples of effective SSP implementation by state public health leadership from Kentucky, ASTHO/North Carolina, and New Mexico.
	• Hidden Casualties: Consequences of the Opioid Epidemic on the Spread of Infectious Diseases (October 23, 2017)
	 Featured VADM Jerome Adams, U.S. Surgeon General, CDC, and other federal leaders. Hidden Casualties: National Partners' Response to the Opioid Epidemic & Infectious Diseases (February 22, 2018 & March 8, 2018)
	Featured federal leaders and nonfederal stakeholders highlighting national partners' efforts to respond including Association of State and Territorial Health Officials, National Association of County and City Health Officials, National Governors Association, Infectious Diseases Society of America, National Alliance of State and Territorial AIDS Directors, National Association of Community Health Centers, and National Viral Hepatitis Roundtable.
OIDP	Increasing Hepatitis C Treatment in State Medicaid Programs
	The Hepatitis C Medicaid Affinity Group supports state-generated solutions for eliminating hepatitis C by increasing the number and percentage of Medicaid beneficiaries diagnosed with hepatitis C who are successfully treated and cured. The project is convened by OIDP and supported by CDC, CMS, HRSA, OMH, and SAMHSA. It brings together state teams including Medicaid, public health, and correctional agencies. In 2018, its second year, most states focused on the intersection of Medicaid and corrections populations. Fourteen states participated in FY18-FY19; and evaluation of the project demonstrated its success in advancing state hepatitis C activities and improving coordination between Medicaid and the public health agencies.



Agency	Activity
	Examples of state activities include: calculating the HCV care cascade; enhancing provider knowledge of HCV testing and treatment; assessing and revising prior authorization processes for HCV medication; improving treatment for people who inject drugs; and linking people in correctional settings to screening and treatment.
OIDP	Mapping Viral Hepatitis Elimination in Action
	To share resources and encourage and support jurisdictions to launch or expand viral hepatitis elimination efforts, OIDP developed a resource map of viral hepatitis elimination projects. This micro-website helps jurisdictions and stakeholders learn from each other's elimination efforts. OIDP and OMH also cosponsored a webinar featuring elimination project examples, practical steps and key messages to help stakeholders develop and implement hepatitis elimination efforts. These projects implement the National Viral Hepatitis
	Action Plan's call for more coordinated and
	collaborative elimination efforts including a wide range of stakeholders such as states, local jurisdictions, health systems, and non-governmental organizations.
OIDP	Expanding Safe Organ Transplantation
	A Federal Workshop on Transplantation from Hepatitis C Infected Donors was hosted by OIDP and CDC to examine and develop a proactive coordinated approach to issues related to the increasing availability and potential use of organs from donors living with hepatitis C, especially as a result of the opioid crisis and increased overdose deaths. The workshop was held in part to address anticipated 2020 updates to the Public Health Service Guideline to Reduce the Risk of HIV/HBV/HCV Transmission via Organ Transplantation. Participants included CDC, CMS, HRSA, NIH, OASH, and transplant researchers and experts.
	Strategy 4.2: Strengthen timely availability and use of data
ВОР	The BOP developed and continues to work on a patient clinical "dashboard" system that allows for tracking select clinical data. There is a dashboard that currently tracks the number of identified hepatitis C-positive patients, and the numbers for patients who have been previously treated, currently on treatment and who may be eligible for treatment.
OIDP	Release and promotion of the Partner Planning Guide to enhance stakeholder engagement in the National Viral Hepatitis Action Plan
	OIDP developed and released the <u>Partner Planning Guide</u> , a companion document to the <u>National Viral Hepatitis</u> <u>Action Plan 2017-2020</u> to support partners in their efforts to build and strengthen viral hepatitis activities. Stakeholders in all levels and sectors have a role to play in viral hepatitis elimination but many are not sure where to start. The Guide is a tool designed for use by individuals, groups, and organizations that are conducting viral hepatitis strategic planning efforts, assessing existing activities, and planning new ones that align with the Action Plan that contribute toward reaching our national viral hepatitis goals.
	Strategy 4.3: Encourage development of improved mechanisms to monitor and report on progress toward achieving national viral hepatitis goals
VA	VA deployed several new/updated data tools in 2018 and created an internal website page as a central location to access them. These include clinical and data tools for HCV infection, HBV infection, and ALD which makes epidemiologic data for these patient populations more broadly accessible at the national, VISN, facility, and provider level. These tools also provide detailed patient-level data for local population health management and to improve clinical care in these areas.
	Strategy 4.4: Regularly report on progress toward achieving the goals of the National Viral Hepatitis Action Plan
FDA	CDER's Division of Antiviral Products utilizes List Serve email notification to enhance communication and for timely dissemination of information regarding new drug approvals and labeling revisions for previously approved drug products.

APPENDIX 2: 2018/2019 VIRAL HEPATITIS-RELATED PUBLICATIONS, ARTICLES, AND REPORTS BY FEDERAL PARTNERS

Federal partners make important contributions to addressing gaps in our understanding of the prevention, care, and treatment of viral hepatitis through peer-reviewed journal articles and other technical documents. These publications help advance efforts to develop and implement evidence-based programs, clinical services, and policies. There were a total of 282 articles or reports published in FY2018 and FY2019 with authors from CDC (51%), VA (44%), IHS (2%), FDA (2%), AHRQ (1%), and OIDP (0.4%). The publications are organized by Action Plan goal area and type of hepatitis. The majority of publications fall within goal one (33%) and goal two (60%). Nine percent of publications focused on hepatitis A, 26% on hepatitis B, 66% on hepatitis C, and 5% on hepatitis E.

The publications contribute to the scientific literature and inform policy recommendations. For example:

- Hofmeister, et al.'s "Estimating prevalence of hepatitis C virus infection in the United States, 2013 – 2016" provides an estimate of the burden of hepatitis C in the United States without a robust surveillance system.
- Jordain, et al.'s "Tenofovir versus placebo to prevent perinatal transmission of hepatitis B" provides evidence for the role of tenofivir for hepatitis B therapy during pregnancy.
 Backus, et al.'s "Impact of sustained virologic response with direct-acting antiviral treatment on mortality in patients with advanced liver disease" provides evidence for decreased risk of all-cause mortality and incident hepatocellular carcinoma with DAA treatment.

GOAL 1: PREVENT NEW VIRAL HEPATITIS INFECTIONS

Hepatitis A

Broderick M, Kamili S, Nelson NP, Le T, Faix D, Romero-Steiner S. Serosurveillance of first-year military personnel for hepatitis A and B. *Am J Public Health*. 2018;108(Suppl 3):S204–206. doi:10.2105/AJPH.2018.304713

Doshani M, Weng M, Moore KL, Romero JR, Nelson NP. Recommendations of the Advisory Committee on Immunization Practices for use of hepatitis A vaccine for persons experiencing homelessness. *MMWR Morb Mortal Wkly Rep.* 2019;68(6):153–156. doi:10.15585/mmwr.mm6806a6

Epstein RL, Sabharwal V, Wachman EM, et al. Perinatal transmission of hepatitis C virus: defining the cascade of care. *J Pediatr.* Published online August 28, 2018;203:34–40.E1. <u>doi:10.1016/j.jpeds.2018.07.006</u>

Foster MA, Hofmeister MG, Kupronis BA, et al. Increase in hepatitis A virus infections—United States, 2013–2018. *MMWR Morb Mortal Wkly Rep.* 2019;68(18):413–415. doi:10.15585/mmwr.mm6818a2

Hofmeister MG, Foster MA, Teshale EH. Epidemiology and transmission of hepatitis A virus and hepatitis E virus infections in the United States. *Cold Spring Harb Perspect Med*. Published online April 30, 2018;9:a033431. doi:10.1101/cshperspect.a033431

Hofmeister MG, McCready JA, Link-Gelles R, et al. Notes from the field: increase in hepatitis A virus infections—Marshall Islands, 2016–2017. *MMWR Morb Mortal Wkly Rep.* 2018;67(17):504–505. doi:10.15585/mmwr.mm6717a5

Link-Gelles R, Hofmeister MG, Nelson NP. Use of hepatitis A vaccine for post-exposure prophylaxis in individuals over 40 years of age: a systematic review of published studies and recommendations for vaccine use. *Vaccine*. Published online April 16, 2018;36(20):2745–2750. doi:10.1016/j.vaccine.2018.04.015



Moorman AC, Xing J, Nelson NP, et al. Need for increasing hepatitis A virus vaccination among patients infected with hepatitis B virus and hepatitis C virus [editorial]. *Gastroenterology*. 2018;154:2015–2017. doi:10.1053/j.gastro.2018.04.031

Mosites E, Gounder P, Snowball M, et al. Hepatitis A vaccine immune response 22 years after vaccination. *J Med Virol.* Published online April 16, 2018;90(8):1418–1422. doi:10.1002/jmv.25197

Nelson NP, Link-Gelles R, Hofmeister MG, et al. Update: recommendations of the Advisory Committee on Immunization Practices for use of hepatitis A vaccine for postexposure prophylaxis and for preexposure prophylaxis for international travel. *MMWR Morb Mortal Wkly Rep.* 2018;67(43):1216–1220. doi:10.15585/mmwr.mm6743a5

Nelson NP, Yankey D, Singleton JA, Elam-Evans LD. Hepatitis A vaccination coverage among adolescents (13–17 years) in the United States, 2008–2016. *Vaccine*. 2018;36(12):1650–1659. doi:10.1016/j.vaccine.2018.01.090

Shing JZ, Ly KN, Xing J, Teshale EH, Jiles RB. Prevalence of hepatitis B virus infection among US adults aged 20–59 years with a history of injection drug use: National Health and Nutrition Examination Survey, 2001–2016. *Clin Infect Dis.* Published online July 27, 2019;70(12):2619–2627. doi:10.1093/cid/ciz669

Tejada-Strop A, Tohme RA, Andre-Alboth J, et al. Seroprevalence of hepatitis A and hepatitis E viruses among pregnant women in Haiti. *Am J Trop Med Hyg.* Published online May 20, 2019;101(1):230–232. <u>doi:10.4269/ajtmh.19-0020</u>

Tejada-Strop A, Zafrullah M, Kamili S, Stramer SL, Purdy MA. Distribution of hepatitis A antibodies in US blood donors. *Transfusion.* Published online October 4, 2018;58(12):2761–2765. doi:10.1111/trf.14916

Viray MA, Hofmeister MG, Johnston DI, et al. Public health investigation and response to a hepatitis A outbreak from imported scallops consumed raw—Hawaii, 2016. *Epidemiol Infect*. Published online October 17, 2018;147(e28):1–8. doi:10.1017/S0950268818002844

Yue X, Black CL, O'Halloran A, Lu PJ, Williams WW, Nelson NP. Hepatitis A and hepatitis B vaccination coverage among adults with chronic liver disease. *Vaccine*. Published online February 7, 2018;36(9):1183–1189. doi:10.1016/j.vaccine.2018.01.033

Hepatitis B

Beste LA, Ioannou GN, Chang MF, et al. Prevalence of hepatitis B virus exposure in the Veterans Health Administration and association with military-related risk factors. *J Clin Gastroenterol Hepatol.* Published online August 5, 2019;18(4):954–962.E6. doi:10.1016/j.cgh.2019.07.056

Broderick M, Kamili S, Nelson NP, Le T, Faix D, Romero-Steiner S. Serosurveillance of first-year military personnel for hepatitis A and B. *Am J Public Health*. 2018;108(Suppl 3):S204–206. doi:10.2105/AJPH.2018.304713

Chahal HS, Peters MG, Harris AM, McCabe D, Volberding P, Kahn JG. Cost-effectiveness of hepatitis B virus infection screening and treatment or vaccination in 6 high-risk populations in the United States. *Open Forum Infect Dis.* 2019;6(1):ofy353. doi:10.1093/ofid/ofy353

Chak E, Taefi A, Li CS, et al. Electronic medical alerts increase screening for chronic hepatitis B: a randomized, double-blind, controlled trial. *Cancer Epidemiol Biomarkers Prev*. Published online August 8, 2018;27(11):1352–1357. doi:10.1158/1055-9965.EPI-18-0448

Choi YH, Perez-Cuevas MB, Kodani M, et al. Feasibility of hepatitis B vaccination by microneedle patch: cellular and humoral immunity studies in rhesus macaques. *J Infect Dis.* Published online August 13, 2019;220(12):1926–1934. doi:10.1093/infdis/jiz399

Collier MG, Doshani M, Asher A. Using population based hospitalization data to monitor increases in conditions causing morbidity among persons who inject drugs. *J Community Health*. Published online January 5, 2018;43(3):598–603. doi:10.1007/s10900-017-0458-9

Cressey TR, Harrison L, Achalapong J, et al. Tenofovir exposure during pregnancy and postpartum in women receiving tenofovir disoproxil fumarate for the prevention of mother-to-child transmission of hepatitis B virus. *Antimicrob Agents Chemother*. Published online October 1, 2018;62(12):e01686-18. doi:10.1128/AAC.01686-18

Fusco DN, Ganova-Raeva L, Khudyakov Y, et al. Reactivation of a vaccine escape hepatitis B virus mutant in a Cambodian patient during anti-hepatitis C virus therapy. *Front Med (Lausanne)*. 2018;5:97. doi:10.3389/fmed.2018.00097

Gidudu JF, Shaum A, Habersaat K, et al. An approach for preparing and responding to adverse events following immunization reported after hepatitis B vaccine birth dose administration. *Vaccine*. Published online ahead of print July 20, 2019. doi:10.1016/j.vaccine.2019.07.041



Haber P, Moro PL, Ng C, et al. Safety of currently licensed hepatitis B surface antigen vaccines in the United States, Vaccine Adverse Event Reporting System (VAERS), 2005–2015. *Vaccine*. Published online December 11, 2017;36(4):559–564. doi:10.1016/j.vaccine.2017.11.079

Hall EW, Rosenberg ES, Trigg M, Nelson N, Schillie S. Cost analysis of single-dose hepatitis B revaccination among infants born to hepatitis B surface antigen-positive mothers and not responding to the initial vaccine series. *Public Health Rep.* Published online April 17, 2018;133(3):338–346. doi:10.1177/003354918768224

Harris AM, Isenhour C, Schillie S, Vellozzi C. Hepatitis B virus testing and care among pregnant women using commercial claims data, United States, 2011–2014. *Infect Dis Obstet Gynecol*. 2018;2018:4107329. doi:10.1155/2018/4107329

Hendrickson B, Kamili S, Timmons T, Iwen PC, Pedati C, Safranek T. Notes from the field: false-negative hepatitis B surface antigen test results in a hemodialysis patient—Nebraska, 2017. *MMWR Morb Mortal Wkly Rep.* 2018;67(10):311–312. doi:10.15585/mmwr.mm6710a6

Jones JM, Gurbaxani BM, Asher A, et al. Quantifying the risk of undetected HIV, hepatitis B virus, or hepatitis C virus infection in Public Health Service increased risk donors. *Am J Transplant*. Published online April 13, 2019;19(9):2583–2593. doi:10.1111/ajt.15393

Jourdain G, Ngo-Giang-Huong N, Harrison L, et al. Tenofovir versus placebo to prevent perinatal transmission of hepatitis B [note]. *Obstet Gynecol Surv.* 2018;73:443–445. https://www.nejm.org/doi/full/10.1056/NEJMoa1708131

Jourdain G, Ngo-Giang-Huong N, Harrison L, et al. Tenofovir versus placebo to prevent perinatal transmission of hepatitis B. *N Engl J Med.* 2018;378:911–923. <u>doi:10.1056/NEJMoa1708131</u>

Kilmer GA, Barker LK, Ly KN, Jiles RB. Hepatitis B vaccination and screening among foreign-born women of reproductive age in the United States: 2013–2015. *Clin Infect Dis.* Published online June 1, 2018;68(2):256–265. doi:10.1093/cid/ciy479

King H, Xing J, Dean HD, Holtzman D. Trends in prevalence of protective levels of hepatitis B surface antibody among adults aged 18–49 years with risk factors for hepatitis B virus infection—United States, 2003–2014. *Clin Infect Dis.* Published online June 20, 2019;70(9):1907–1915. doi:10.1093/cid/ciz537

Koneru A, Schillie S, Roberts H, et al. Estimating annual births to hepatitis B surface antigen-positive women in the United States by using data on maternal country of birth. *Public Health Rep.* Published online April 3, 2019;134(3):255–263. doi:10.1177/0033354919836958

Lu PJ, O'Halloran AC, Williams WW, Nelson NP. Hepatitis B vaccination coverage among adults aged ≥18 years traveling to a country of high or intermediate endemicity, United States, 2015. *Vaccine*. Published online March 28, 2018;36(18):2471–2479. doi:10.1016/j.vaccine.2018.03.030

Mittal S, Kramer JR, Omino R, et al. Role of age and race in the risk of hepatocellular carcinoma in veterans with hepatitis B virus infection. *Clin Gastroenterol Hepatol.* 2018;16(2):252–259. doi:10.1016/j.cgh.2017.08.042

Mixson-Hayden T, Purdy MA, Ganova-Raeva L, McGovern D, Forbi JC, Kamili S. Evaluation of performance characteristics of hepatitis B e antigen serologic assays. *Clin Diagn Virol.* 2018;109:22–28. doi:10.1016/j.jcv.2018.10.005

Moorman AC, Xing J, Nelson NP, et al. Need for increasing hepatitis A virus vaccination among patients infected with hepatitis B virus and hepatitis C virus [editorial]. *Gastroenterology*. 2018;154:2015–2017. doi:10.1053/j.gastro.2018.04.031

Paul RC, Rahman M, Wiesen E, et al. Hepatitis B surface antigen seroprevalence among prevaccine and vaccine era children in Bangladesh. *Am J Trop Med Hyg.* Published online July 16, 2018;99(3):764–771. doi:10.4269/ajtmh.17-0721

Pauly MD, Kamili S, Hayden TM. Impact of nucleic acid extraction platforms on hepatitis virus genome detection. *J Virol Methods*. 2019;273:113715. doi:10.1016/j.jviromet.2019.113715

Perez Cuevas MB, Kodani M, Choi Y, et al. Hepatitis B vaccination using a dissolvable microneedle patch is immunogenic in mice and rhesus macaques. *Bioeng Transl Med.* 2018;3:186–196. doi:10.1002/btm2.10098

Ramachandran S, Thai H, Forbi JC, et al. A large HCV transmission network enabled a fast-growing HIV outbreak in rural Indiana, 2015. *EBioMedicine*. Published online November 15, 2018;37:374–381. doi:10.1016/j.ebiom.2018.10.007

Reardon JM, O'Connor SM, Njau JD, Lam EK, Staton CA, Cookson ST. Cost-effectiveness of birth-dose hepatitis B vaccination among refugee populations in the African region: a series of case studies. *Confl Health*. 2019;13:5. <u>doi:10.1186/s13031-019-0188-y</u>

Roberts H, Boktor SW, Waller K, et al. Underreporting of hepatitis B and C virus infections—Pennsylvania, 2001–2015. *PLoS One.* 2019;14(6):e0217455. doi:10.1371/journal.pone.0217455



Salvadori N, Fan B, Teeyasoontranon W, et al. Maternal and infant bone mineral density 1 year after delivery in a randomized, controlled trial of maternal tenofovir disoproxil fumarate to prevent mother-to-child transmission of hepatitis B virus. *Clin Infect Dis.* Published online March 29, 2019;69(1):144–146. doi:10.1093/cid/ciy982

Schillie S, Harris A, Link-Gelles R, Romero J, Ward J, Nelson N. Recommendations of the Advisory Committee on Immunization Practices for use of a hepatitis B vaccine with a novel adjuvant. *MMWR Morb Mortal Wkly Rep.* 2018;67(15):455–458. doi:10.15585/mmwr.mm6715a5

Schillie S, Vellozzi C, Reingold A, et al. Prevention of hepatitis B virus infection in the United States: recommendations of the Advisory Committee on Immunization Practices. *MMWR Recomm Rep.* 2018;67(No. RR-1):1–31. doi:10.15585/mmwr.rr6701a1

Shing JZ, Ly KN, Xing J, Teshale EH, Jiles RB. Prevalence of hepatitis B virus infection among US adults aged 20–59 years with a history of injection drug use: National Health and Nutrition Examination Survey, 2001–2016. *Clin Infect Dis*. Published online July 27, 2019;70(12):2619–2627. <u>doi:10.1093/cid/ciz669</u>

US Preventive Services Task Force. Screening for hepatitis B virus infection in pregnant women: US Preventive Services Task Force reaffirmation recommendation statement. *JAMA*. 2019;322(4):349–354. doi:10.1001/jama.2019.9365

Yue X, Black CL, O'Halloran A, Lu PJ, Williams WW, Nelson NP. Hepatitis A and hepatitis B vaccination coverage among adults with chronic liver disease. *Vaccine*. Published online February 7, 2018;36(9):1183–1189. doi:10.1016/j.vaccine.2018.01.033

Hepatitis C

Abara WE, Moorman AC, Zhong Y, et al. The predictive value of International Classification of Disease codes for chronic hepatitis C virus infection surveillance: the utility and limitations of electronic health records. *Popul Health Manag.* 2018;21(2):110–115. doi:10.1089/pop.2017.0004

Abara WE, Spradling P, Zhong Y, et al. Hepatocellular carcinoma surveillance in a cohort of chronic hepatitis C virus-infected patients with cirrhosis. *J Gastrointest Cancer*. Published online May 23, 2019;51:464–468. doi:10.1007/s12029-019-00255-4

Alroy-Preis S, Daly ER, Adamski C, et al. Large outbreak of hepatitis C virus associated with drug diversion by a healthcare technician. *Clin Infect Dis*. Published online May 14, 2018;67(6):845–853. doi:10.1093/cid/ciy193

Barocas JA, Tasillo A, Eftekhari G, Wang J, Vellozzi C, Hariri S, Isenhour C, Randall L, Ward JW, Mermin J, Salomon JA, Linas BP. Population Level Outcomes and Cost-Effectiveness of Expanding the Recommendation for Age-Based Hepatitis C Testing in the United States, Clinical Infectious Diseases. 2018 Feb. 6. Accessed 2/9/2018. <u>doi: 10.1093/cid/ciy098</u>

Campo DS, Khudyakov Y. Intelligent Network DisRuption Analysis (INDRA): a targeted strategy for efficient interruption of hepatitis C transmissions. *Infect Genet Evol*. Published online May 31, 2018;63:204–215. doi:10.1016/j.meegid.2018.05.028

Campo DS, Zhang J, Ramachandran S, Khudyakov Y. Transmissibility of intra-host hepatitis C virus variants. *BMC Genomics*. 2017;18(Suppl 10):881. doi:10.1186/s12864-017-4267-4

Collier MG, Doshani M, Asher A. Using population based hospitalization data to monitor increases in conditions causing morbidity among persons who inject drugs. *J Community Health*. Published online January 5, 2018;43(3):598–603. doi:10.1007/s10900-017-0458-9

Epstein RL, Sabharwal V, Wachman EM, et al. Perinatal transmission of hepatitis C virus: defining the cascade of care. *J Pediatr.* Published online August 28, 2018;203:34–40.E1. <u>doi:10.1016/j.jpeds.2018.07.006</u>

Fill MA, Sizemore LA, Rickles M, et al. Epidemiology and risk factors for hepatitis C virus infection in a high-prevalence population. *Epidemiol Infect*. Published online February 12, 2018;146(4):508–514. doi:10.1017/S0950268818000080

Fusco DN, Ganova-Raeva L, Khudyakov Y, et al. Reactivation of a vaccine escape hepatitis B virus mutant in a Cambodian patient during anti-hepatitis C virus therapy. *Front Med (Lausanne).* 2018;5:97. <u>doi:10.3389/fmed.2018.00097</u>

Ganova-Raeva L, Dimitrova Z, Alexiev I, et al. HCV transmission in high-risk communities in Bulgaria. *PLoS One*. 2019;14(3):e0212350. doi:10.1371/journal.pone.0212350

Glebova O, Knyazev S, Melnyk A, et al. Inference of genetic relatedness between viral quasispecies from sequencing data. *BMC Genomics.* 2017;18(Suppl 10):918. doi:10.1186/s12864-017-4274-5

Hofmeister MG, Rosenthal EM, Barker LK, et al. Estimating prevalence of hepatitis C virus infection in the United States, 2013–2016. *Hepatology*. Published online November 6, 2018;69(3):1020–1031. <u>doi:10.1002/hep.30297</u>

Irvin R, Ward K, Agee T, et al. Comparison of hepatitis C virus testing recommendations in high-income countries. *World J Hepatol.* 2018;10(10):743–751. doi:10.4254/wjh.v10.i10.743



Jones JM, Gurbaxani BM, Asher A, et al. Quantifying the risk of undetected HIV, hepatitis B virus, or hepatitis C virus infection in Public Health Service increased risk donors. *Am J Transplant*. Published online April 13, 2019;19(9):2583–2593. doi:10.1111/ajt.15393

Ko JY, Haight SC, Schillie SF, Bohm MK, Dietz PM. National trends in hepatitis C infection by opioid use disorder status among pregnant women at delivery hospitalization—United States, 2000–2015. *MMWR Morb Mortal Wkly Rep.* 2019;68(39):833–838. doi:10.15585/mmwr.mm6839a1

Kodani M, Martin M, de Castro VL, Drobeniuc J, Kamili S. An automated immunoblot method for detection of IgG antibodies to hepatitis C virus: a potential supplemental antibody confirmatory assay. *J Clin Microbiol*. Published online January 16, 2019;57(3):e01567. doi:10.1128/JCM.01567-18

Lara J, Teka M, Khudyakov Y. Identification of recent cases of hepatitis C virus infection using physical-chemical properties of hypervariable region 1 and a radial basis function neural network classifier. *BMC Genomics*. 2017;18(Suppl 10):880. doi:10.1186/s12864-017-4269-2

Lu M, Li J, Rupp LB, et al. Changing trends in complications of chronic hepatitis C. *Liver Int*. 2018;38(2):239–247. doi:10.1111/liv.13501

Lu M, Wu KH, Li J, et al. Adjuvant ribavirin and longer direct-acting antiviral treatment duration improve sustained virological response among hepatitis C patients at risk of treatment failure. *J Viral Hepat*. Published online June 13, 2019;26(10):1210–1217. doi:10.1111/jvh.13162

Ly KN, Kim AA, Drobenuic J, et al. The prevalence of hepatitis C virus antibody in HIV-negative persons in Kenya, 2007. *Am J Trop Med Hyg.* Published online April 23, 2018;98(6):1876. doi:10.4269/ajtmh.17-0830

Mane A, Sacks J, Sharma S, et al. Evaluation of five rapid diagnostic tests for detection of antibodies to hepatitis C virus (HCV): a step towards scale-up of HCV screening efforts in India. *PLoS One*. 2019;14(1):e0210556. doi:10.1371/journal.pone.0210556

Moorman AC, Xing J, Nelson NP, et al. Need for increasing hepatitis A virus vaccination among patients infected with hepatitis B virus and hepatitis C virus [editorial]. *Gastroenterology*. 2018;154:2015–2017. <u>doi:10.1053/j.gastro.2018.04.031</u>

Njuguna HN, Stinson D, Montgomery P, et al. Hepatitis C virus potentially transmitted by opioid drug diversion from a nurse— Washington, August 2017–March 2018. *MMWR Morb Mortal Wkly Rep*. 2019;68(16):374–376. doi:10.15585/mmwr.mm6816a3

Nolen LD, Gustin C, Seeman S, et al. Risk-based prenatal hepatitis C testing practices and results, Alaska 2013–2016. *Can J Gastroenterol Hepatol.* 2019;2019:8654741. doi:10.1155/2019/8654741

Nwaohiri A, Schillie S, Bulterys M, Kourtis AP. Hepatitis C virus infection in children: how do we prevent it and how do we treat it? *Expert Rev Anti Infect Ther.* Published online August 21, 2018;16(9):689–694. doi:10.1080/14787210.2018.1509707

Osinubi A, Harris AM, Vellozzi C, Lom J, Miller L, Millman AJ. Evaluation of the performance of algorithms that use serial hepatitis C RNA tests to predict treatment initiation and sustained virological response among patients infected with hepatitis C virus. *Am J Epidemiol.* Published online December 7, 2018;188(3):555–561. doi:10.1093/aje/kwy270

Pauly MD, Kamili S, Hayden TM. Impact of nucleic acid extraction platforms on hepatitis virus genome detection. *J Virol Methods*. 2019;273:113715. doi:10.1016/j.jviromet.2019.113715

Poe A, Duong NT, Bedi K, Kodani M. Stability of hepatitis C virus RNA and anti-HCV antibody in air-dried and freeze-dried human plasma samples. *J Virol Methods*. Published online December 15, 2017;253:53–55. doi:10.1016/j.jviromet.2017.12.004

Ramachandran S, Thai H, Forbi JC, et al. A large HCV transmission network enabled a fast-growing HIV outbreak in rural Indiana, 2015. *EBioMedicine*. Published online November 15, 2018;37:374–381. <u>doi:10.1016/j.ebiom.2018.10.007</u>

Reilley B, Leston J, Doshani M, et al. Assessing disparities in the rates of HCV diagnoses within American Indian or Alaska Native populations served by the US Indian Health Service, 2005–2015. *J Community Health*. 2018;43(6):1115–1118. doi:10.1007/s10900-018-0528-7

Roberts H, Boktor SW, Waller K, et al. Underreporting of hepatitis B and C virus infections—Pennsylvania, 2001–2015. *PLoS One*. 2019;14(6):e0217455. doi:10.1371/journal.pone.0217455

Rosenberg ES, Rosenthal EM, Hall EW, et al. Prevalence of hepatitis C virus infection in US states and the District of Columbia, 2013 to 2016. *JAMA Netw Open*. 2018;1(8):e186371. doi:10.1001/jamanetworkopen.2018.6371

Schillie SF, Canary L, Koneru A, et al. Hepatitis C virus in women of childbearing age, pregnant women, and children. *Am J Prev Med.* 2018;55(5):633–641. doi:10.1016/j.amepre.2018.05.029



Sims S, Longmire AG, Campo DS, et al. Automated quality control for a molecular surveillance system. *BMC Bioinformatics*. 2018;19(Suppl 11):358. doi:10.1186/s12859-018-2329-5

Skums P, Zelikovsky A, Singh R, et al. QUENTIN: reconstruction of disease transmissions from viral quasispecies genomic data. *Bioinformatics*. 2018;34(1):163–170. doi:10.1093/bioinformatics/btx402

Sood A, Suryaprasad A, Trickey A, et al. The burden of hepatitis C virus infection in Punjab, India: a population-based serosurvey. *PLoS One.* 2018;13(7):e0200461. doi:10.1371/journal.pone.0200461

Tasillo A, Eftekhari Yazdi G, Nolen S, et al. Short-term effects and long-term cost-effectiveness of universal hepatitis C testing in prenatal care. *Obstet Gynecol.* Published online February 1, 2019;133(2):289–300. doi:10.1097/AOG.0000000000003062

Tsyvina V, Campo DS, Sims S, Zelikovsky A, Khudyakov Y, Skums P. Fast estimation of genetic relatedness between members of heterogeneous populations of closely related genomic variants. *BMC Bioinformatics*. 2018;19(Suppl 11):360. doi:10.1186/s12859-018-2333-9

Vohra D, Huang M, O'Neil S. The costs and benefits of expanding hepatitis C screening in the Indian Health Service. Published September 2018. Accessed August 219, 2020. <u>https://aspe.hhs.gov/system/files/pdf/260026/HepC.pdf</u>

Yin S, Barker L, White JZ, Jiles RB. Sofosbuvir-based regimens for chronic hepatitis C in a well-insured U.S. population: patient characteristics, treatment adherence, effectiveness, and health care costs, 2013–2015. *J Manag Care Spec Pharm.* 2019;25(2):195–210. doi:10.18553/jmcp.2019.25.2.195

Hepatitis E

Baylis SA, Hanschmann K-MO, Matsubayashi K, et al. Development of a World Health Organization international reference panel for different genotypes of hepatitis E virus for nucleic acid amplification testing. *J Clin Virol*. Published online May 14, 2019;119:60–67. doi:10.1016/j.jcv.2019.05.006

Castro VOL, Tejada-Strop A, Weis SMS, et al. Evidence of hepatitis E virus infections among persons who use crack cocaine from the Midwest region of Brazil. *J Med Virol*. Published online August 22, 2018;91(1):151–154. doi:10.1002/jmv.25288

Choi Y, Zhang X, Skinner B. Analysis of IgG anti-HEV antibody protective levels during hepatitis E virus reinfection in experimentally infected rhesus macaques. *J Infect Dis.* Published online October 16, 2018;219(6):916–924. doi:10.1093/infdis/jiy603

Choi YH, Zhang X, Tran C, Skinner B. Expression profiles of host immune response-related genes against HEV genotype 3 and genotype 1 infections in rhesus macaques. *J Viral Hepat*. Published online March 13, 2018;25(8):986–995. doi:10.1111/jvh.12890

Hofmeister MG, Foster MA, Teshale EH. Epidemiology and transmission of hepatitis A virus and hepatitis E virus infections in the United States. *Cold Spring Harb Perspect Med*. Published online April 30, 2018;9:a033431. doi:10.1101/cshperspect.a033431

Miernyk KM, Bruden D, Parkinson AJ, et al. Human seroprevalence to 11 zoonotic pathogens in the U.S. Arctic, Alaska. *Vector Borne Zoonotic Dis.* Published online August 2, 2019;19(8):563–575. <u>doi:10.1089/vbz.2018.2390</u>

Pauly MD, Kamili S, Hayden TM. Impact of nucleic acid extraction platforms on hepatitis virus genome detection. *J Virol Methods*. 2019;273:113715. doi:10.1016/j.jviromet.2019.113715

Tejada-Strop A, Tohme RA, Andre-Alboth J, et al. Seroprevalence of hepatitis A and hepatitis E viruses among pregnant women in Haiti. *Am J Trop Med Hyg.* Published online May 20, 2019;101(1):230–232. doi:10.4269/ajtmh.19-0020

Zafrullah M, Zhang X, Tran C, et al. Disparities in detection of antibodies against hepatitis E virus in US blood donor samples using commercial assays. *Transfusion*. Published online March 9, 2018;58(5):1254–1263. doi:10.1111/trf.14553

GOAL 2: REDUCE DEATHS AND IMPROVE THE HEALTH OF PEOPLE LIVING WITH VIRAL HEPATITIS

Hepatitis A

Barrett CE, Pape BJ, Benedict KM, et al. Impact of public health interventions on drinking water–associated outbreaks of hepatitis A—United States, 1971–2017. *MMWR Morb Mortal Wkly Rep.* 2019;68(35):766–770. doi:10.15585/mmwr.mm6835a4



Bixler D, Corby-Lee G, Proescholdbell S, et al. Access to syringe services programs—Kentucky, North Carolina, and West Virginia, 2013–2017. *MMWR Morb Mortal Wkly Rep.* 2018;67(18):529–532. doi:10.15585/mmwr.mm6718a5

Cahill JA, Rizvi S, Saeian K. Assessment of adherence to baseline quality measures for cirrhosis and the impact of performance feedback in a regional VA medical center. *Am J Med Qual.* 2018;33(3):262–268. doi:10.1177/1062860617736805

Chen W, Luan J, Wei G, et al. In vivo hepatocellular expression of interleukin-22 using penetratin-based hybrid nanoparticles as potential anti-hepatitis therapeutics. *Biomaterials*. 2018;187:66–80. doi:10.1016/j.biomaterials.2018.09.046

Singal AG, Lim JK, Kanwal F. AGA clinical practice update on interaction between oral direct-acting antivirals for chronic hepatitis C infection and hepatocellular carcinoma: expert review. *Gastroenterology*. 2019;156(8):2149–2157. doi:10.1053/j.gastro.2019.02.046

Hepatitis B

Abara WE, Collier MG, Moorman A, et al. Characteristics of deceased solid organ donors and screening results for hepatitis B, C, and human immunodeficiency viruses—United States, 2010–2017. *Am J Transplant*. 2019;19(3):939–947. doi:10.1111/ajt.15284

Abara WE, Qaseem A, Schillie S, McMahon BJ, Harris AM. Hepatitis B vaccination, screening, and linkage to care: best practice advice from the American College of Physicians and the Centers for Disease Control and Prevention. *Ann Intern Med.* 2017;167(11):794–804. doi:10.7326/M17-1106

Althoff KN, Gebo KA, Moore RD, et al. Contributions of traditional and HIV-related risk factors on non-AIDS-defining cancer, myocardial infarction, and end-stage liver and renal diseases in adults with HIV in the USA and Canada: a collaboration of cohort studies. *Lancet HIV*. 2019;6(2):E93–E104. doi:10.1016/S2352-3018(18)30295-9

Anderson RT, Lim SG, Mishra P, et al. Challenges, considerations, and principles to guide trials of combination therapies for chronic hepatitis B virus. *Gastroenterology*. 2019;156(3):529–533.E4. <u>doi:10.1053/j.gastro.2018.11.062</u>

Bixler D, Annambholta P, Abara WE, et al. Hepatitis B and C virus infections transmitted through organ transplantation investigated by CDC, United States, 2014–2017. *Am J Transplant*. Published online March 12, 2019;19(9):2570–2582. doi:10.1111/ajt.15352

Bixler D, Zhong Y, Ly KN, et al. Mortality among patients with chronic hepatitis B infection: the Chronic Hepatitis Cohort Study (CHeCS). *Clin Infect Dis.* Published online July 28, 2018;68(6):956–963. <u>doi:10.1093/cid/ciy598</u>

Bridges CB, Watson TL, Nelson NP, et al. Challenges with hepatitis B vaccination of high risk adults—a pilot program. *Vaccine*. Published online July 11, 2019;37(35):5111–5120. doi:10.1016/j.vaccine.2019.05.089

Bullard AJ, Cunningham FE, Volpp BD, et al. Preventing hepatitis B reactivation during anti-CD20 antibody treatment in the Veterans Health Administration. *Hepatol Commun.* 2018;2(9):1136–1146. doi:10.1002/hep4.1238

Cahill JA, Rizvi S, Saeian K. Assessment of adherence to baseline quality measures for cirrhosis and the impact of performance feedback in a regional VA medical center. *Am J Med Qual.* 2018;33(3):262–268. doi:10.1177/1062860617736805

Fox RK, Taddei TH, Kaplan DE. Aspirin use and risk of hepatocellular carcinoma in hepatitis B. *JAMA Intern Med.* 2019;179(5):640–641. doi:10.1001/jamainternmed.2018.8314

Ganesan M, Poluektova LY, Kharbanda KK, Osna NA. Human immunodeficiency virus and hepatotropic viruses co-morbidities as the inducers of liver injury progression. *World J Gastroenterol.* 2019;25(4):398–410. doi:10.3748/wjg.v25.i4.398

Harris AM, Link-Gelles R, Kim K, et al. Community-based services to improve testing and linkage to care among non–U.S.-born persons with chronic hepatitis B virus infection—three U.S. programs, October 2014–September 2017. *MMWR Morb Mortal Wkly Rep.* 2018;67(19):541–546. doi:10.15585/mmwr.mm6719a2

Harris AM, Millman AJ, Lora M, Osinubi A, Lom J, Miller LS. Hepatitis B testing, care linkage, and vaccination coverage within a registry of hepatitis C infected patients. *Vaccine*. 2019;37(16):2188–2193. <u>doi:10.1016/j.vaccine.2019.03.012</u>

Hou W, Trieu C, Du Y, Wang C, Syn W-K. Profile of drug resistance mutations in nucleos(t)ide analogue-experienced chronic hepatitis B patients in Tianjin, China. *Int J Antimicrob Agents*. 2018;52(5):735–736. doi:10.1016/j.ijantimicag.2018.07.024

Hutin Y, Nasrullah M, Easterbrook P, et al. Access to treatment for hepatitis B virus infection—worldwide, 2016. *Am J Transplant*. 2018;18(10):2595–2598. doi:10.1111/ajt.15093

Ji Y, Dang X, Nguyen LNT, et al. Topological DNA damage, telomere attrition and T cell senescence during chronic viral infections. *Immun Ageing*. 2019;16:12. <u>doi:10.1186/s12979-019-0153-z</u>



Kim HS, El-Serag HB. The epidemiology of hepatocellular carcinoma in the USA. *Curr Gastroenterol Rep.* 2019;21(4):17. doi:10.1007/s11894-019-0681-x

Kim HS, Yang JD, El-Serag HB, Kanwal F. Awareness of chronic viral hepatitis in the United States: an update from the National Health and Nutrition Examination Survey. *J Viral Hepat.* 2019;26(5):596–602. <u>doi:10.1111/jvh.13060</u>

Kulik L, El-Serag HB. Epidemiology and management of hepatocellular carcinoma. *Gastroenterology*. 2019;156(2):477–491.E1. doi:10.1053/j.gastro.2018.08.065

Li J, Zhang T, Gordon SC, et al. Impact of sustained virologic response on risk of type 2 diabetes among hepatitis C patients in the United States. *J Viral Hepat*. Published online February 25, 2018;25(8):952–958. doi:10.1111/jvh.12887

Mitruka K, Pezzi C, Baack B, et al. Evaluation of hepatitis B virus screening, vaccination, and linkage to care among newly arrived refugees in four states, 2009–2011. *J Immigr Minor Health*. Published online February 7, 2018;21(1):39–46. doi:10.1007/s10903-018-0705-x

Mittal S, Kramer JR, Omino R, et al. Role of age and race in the risk of hepatocellular carcinoma in veterans with hepatitis B virus infection. *Clin Gastroenterol Hepatol.* 2018;16(2):252–259. doi:10.1016/j.cgh.2017.08.042

Momin B, Millman AJ, Nielsen DB, Revels M, Steele CB. Promising practices for the prevention of liver cancer: a review of the literature and cancer plan activities in the National Comprehensive Cancer Control Program. *Cancer Causes Control*. Published online December 1, 2018;29(12):1265–1275. doi:10.1007/s10552-018-1094-0

Moon AM, Singal AG, Tapper EB. Contemporary epidemiology of chronic liver disease and cirrhosis. *Clin Gastroenterol Hepatol.* Published online ahead of print August 8, 2019. <u>doi:10.1016/j.cgh.2019.07.060</u>

Moorman AC, Xing J, Rupp LB, et al. Hepatitis B virus infection and hepatitis C virus treatment in a large cohort of hepatitis C-infected patients in the United States. *Gastroenterology*. 2018;154(3):754–758. doi:10.1053/j.gastro.2017.12.002

Mücke MM, Backus LI, Mücke VT, et al. Hepatitis B virus reactivation during direct-acting antiviral therapy for hepatitis C: a systematic review and meta-analysis. *Lancet Gastroenterol Hepatol*. Published online January 19, 2018;3(3):172–180. doi:10.1016/S2468-1253(18)30002-5

O'Connor SM, Mixson-Hayden T, Ganova-Raeva L, et al. Integrated HIV surveillance finds recent adult hepatitis B virus (HBV) transmission and intermediate HBV prevalence among military in uncharacterized Caribbean country. *PLoS One.* 2019;14(10):e0222835. doi:10.1371/journal.pone.0222835

Patterson S, Schmajuk G, Evans M, et al. Gaps in ambulatory patient safety for immunosuppressive specialty medications. *Jt Comm J Qual Patient Saf.* 2019;45(5):348–357. doi:10.1016/j.jcjq.2018.12.003

Sanaka S, Kasarala GR, Tillmann HL. A downside to hepatitis C virus cure? Vigilance is needed regarding hepatitis B virus reactivation, organ rejection, or hepatocellular carcinoma progression. *J Infect Dis.* 2018;217(6):857–860. doi:10.1093/infdis/jix659

Shah R, Ho EY, Kramer JR, et al. Hepatitis B virus screening and reactivation in a national VA cohort of patients with inflammatory bowel disease treated with tumor necrosis factor antagonists. *Dig Dis Sci*. 2018;63(6):1551–1557. doi:10.1007/s10620-018-5042-3

Talaat N, Tillmann HL. Injury pattern recognition to discriminate competing causes of liver injury. *Liver Int.* 2019;39(5):821–825. doi:10.1111/liv.14056

Taplitz RA, Kennedy EB, Bow EJ, et al. Antimicrobial prophylaxis for adult patients with cancer-related immunosuppression: ASCO and IDSA clinical practice guideline update. *J Clin Oncol.* 2018;36(30):3043–3054. <u>doi:10.1200/JCO.18.00374</u>

Wei MT, Le AK, Chang MS, et al. Antiviral therapy and the development of osteopenia/osteoporosis among Asians with chronic hepatitis B. J Med Virol. 2019;91(7):1288–1294. doi:10.1002/jmv.25433

Zou B, Yeo YH, Jeong D, et al. A nationwide study of inpatient admissions, mortality, and costs for patients with cirrhosis from 2005 to 2015 in the USA. *Dig Dis Sci.* Published online October 9, 2019;65(5):1520–1528. <u>doi:10.1007/s10620-019-05869-z</u>

Hepatitis C

Abara WE, Collier MG, Moorman A, et al. Characteristics of deceased solid organ donors and screening results for hepatitis B, C, and human immunodeficiency viruses—United States, 2010–2017. *Am J Transplant*. 2019;19(3):939–947. doi:10.1111/ajt.15284



Abara WE, Qaseem A, Schillie S, McMahon BJ, Harris AM. Hepatitis B vaccination, screening, and linkage to care: best practice advice from the American College of Physicians and the Centers for Disease Control and Prevention. *Ann Intern Med.* 2017;167(11):794–804. doi:10.7326/M17-1106

Ahmed A, Schriever C, Britt N, et al. Comparing drug interaction frequencies of various hepatitis C treatment regimens among monoinfected patients. *Ann Hepatol.* 2019;18(4):601–606. doi:10.1016/j.aohep.2019.01.005

Althoff KN, Gebo KA, Moore RD, et al. Contributions of traditional and HIV-related risk factors on non-AIDS-defining cancer, myocardial infarction, and end-stage liver and renal diseases in adults with HIV in the USA and Canada: a collaboration of cohort studies. *Lancet HIV*. 2019;6(2):E93–E104. doi:10.1016/S2352-3018(18)30295-9

Assoumou SA, Tasillo A, Vellozzi C, et al. Cost-effectiveness and budgetary impact of hepatitis C virus testing, treatment, and linkage to care in US prisons. *Clin Infect Dis.* Published online May 16, 2019;70(7):1388–1396. doi:10.1093/cid/ciz383

Assoumou SA, Wang J, Tasillo A, Yazdi GE, Tsui JI, Strick L, Linas BP. Hepatitis C Testing and Patient Characteristics in Washington State's Prisons Between 2012 and 2016. American journal of preventive medicine. 2019 Jan 1;56(1):8-16. doi: 10.1016/j.amepre.2018.08.016

Back D, Belperio P, Bondin M, et al. Efficacy and safety of glecaprevir/pibrentasvir in patients with chronic HCV infection and psychiatric disorders: an integrated analysis. *J Viral Hepat.* 2019;26(8):951–960. doi:10.1111/jvh.13110

Backus LI, Belperio PS, Shahoumian TA, Mole LA. Impact of sustained virologic response with direct-acting antiviral treatment on mortality in patients with advanced liver disease. *Hepatology*. 2019;69(2):487–497. doi:10.1002/hep.29408

Backus LI, Shahoumian TA, Belperio PS, et al. Impact of *IFNL4*- Δ G genotype on sustained virologic response in hepatitis C genotype 1 patients treated with direct-acting antivirals. *Diagn Microbiol Infect Dis.* 2018;92(1):34–36. doi:10.1016/j.diagmicrobio.2018.04.004

Bakr O, Gelberg L, Seragaki S, et al. Treating hepatitis C in homeless veterans at the Greater Los Angeles Veterans' Affairs Medical Center. *Hepatology*. 2019;70(3):1071–1073. <u>doi:10.1002/hep.30643</u>

Barbosa C, Fraser H, Hoerger TJ, et al. Cost-effectiveness of scaling up HCV prevention and treatment in the United States for people who inject drugs. *Addiction*. Published online July 15, 2019;114(12):2267–2278. <u>https://doi.org/10.1111/add.14731</u>

Barnett PG, Joyce VR, Lo J, et al. Effect of interferon-free regimens on disparities in hepatitis C treatment of US veterans. *Value Health*. 2018;21(8):921–930. doi:10.1016/j.jval.2017.12.025

Barocas JA, Tasillo A, Eftekhari Yazdi G, et al. Population-level outcomes and cost-effectiveness of expanding the recommendation for age-based hepatitis C testing in the United States. *Clin Infect Dis.* Published online February 6, 2018;67(4):549–556. doi:10.1093/cid/ciy098

Belperio PS, Chartier M, Gonzalez RI, et al. Hepatitis C care in the Department of Veterans Affairs: building a foundation for success. *Infect Dis Clin North Am.* 2018;32(2):281–292. doi:10.1016/j.idc.2018.02.011

Belperio PS, Chartier M, Ross DB. After curing hepatitis C virus infection. Ann Intern Med. 2018;168(9):682–683. doi:10.7326/L18-0014

Belperio PS, Shahoumian TA, Loomis TP, Backus LI. Real-world effectiveness of sofosbuvir/velpatasvir/voxilaprevir in 573 directacting antiviral experienced hepatitis C patients. *J Viral Hepat.* 2019;26(8):980–990. doi:10.1111/jvh.13115

Belperio PS, Shahoumian TA, Loomis TP, Mole LA, Backus LI. Real-world effectiveness of daclatasvir plus sofosbuvir and velpatasvir/sofosbuvir in hepatitis C genotype 2 and 3. *J Hepatol.* 2019;70(1):15–23. doi:10.1016/j.jhep.2018.09.018

Bendich I, Takemoto S, Patterson JT, Monto A, Barber TC, Kuo AC. Preoperative treatment of hepatitis C is associated with lower prosthetic joint infection rates in US veterans. *J Arthroplasty*. 2019;34(Suppl 7):S319–S326.E1. doi:10.1016/j.arth.2019.02.052

Benhammou JN, Dong TS, May FP, et al. Race affects SVR12 in a large and ethnically diverse hepatitis C-infected patient population following treatment with direct-acting antivirals: analysis of a single-center Department of Veterans Affairs cohort. *Pharmacol Res Perspect.* 2018;6(2):e00379. doi:10.1002/prp2.379

Bethea ED, Samur S, Kanwal F, et al. Cost effectiveness of transplanting HCV-infected livers into uninfected recipients with preemptive antiviral therapy. *Clin Gastroenterol Hepatol.* 2019;17(4):739–747.E8. <u>doi:10.1016/j.cgh.2018.08.042</u>

Bixler D, Annambholta P, Abara WE, et al. Hepatitis B and C virus infections transmitted through organ transplantation investigated by CDC, United States, 2014–2017. *Am J Transplant*. Published online March 12, 2019;19(9):2570–2582. doi:10.1111/ajt.15352



Bixler D, Corby-Lee G, Proescholdbell S, et al. Access to syringe services programs—Kentucky, North Carolina, and West Virginia, 2013–2017. *MMWR Morb Mortal Wkly Rep.* 2018;67(18):529–532. doi:10.15585/mmwr.mm6718a5

Bixler D, Zhong Y, Ly KN, et al. Mortality among patients with chronic hepatitis B infection: the Chronic Hepatitis Cohort Study (CHeCS). *Clin Infect Dis*. Published online July 28, 2018;68(6):956–963. <u>doi:10.1093/cid/ciy598</u>

Boyd SD, Harrington P, Komatsu TE, et al. HCV genotype 4, 5 and 6: distribution of viral subtypes and sustained virologic response rates in clinical trials of approved direct-acting antiviral regimens. *J Viral Hepat.* 2018;25(8):969–975. doi:10.1111/jvh.12896

Brady JE, Vellozzi C, Hariri S, et al. Hepatitis C care cascade among persons born 1945–1965: 3 medical centers. *Am J Manag Care*. 2018;24(9):421–427. Accessed August 19, 2020. https://www.ncbi.nlm.nih.gov/pubmed/30222920

Butt AA, Yan P, Lo Re V III, Shaikh OS, Ross DB. Trends in treatment uptake and provider specialty for hepatitis C virus (HCV) infection in the Veterans Affairs healthcare system: results from the Electronically Retrieved Cohort of HCV-Infected Veterans (ERCHIVES). *Clin Infect Dis.* 2019;68(5):857–859. <u>doi:10.1093/cid/ciy697</u>

Cao D, Zhao J, Nguyan LN, et al. Disruption of telomere integrity and DNA repair machineries by KML001 induces T cell senescence, apoptosis, and cellular dysfunctions. *Front Immunol.* 2019;10:1152. doi:10.3389/fimmu.2019.01152

Cash J, Skinner A, Cash S, Jones A, Waters B, Skinner RB Jr. Blistering disease during the treatment of chronic hepatitis C with ledipasvir/sofosbuvir. *Fed Pract.* 2019;36(Suppl 2):S11–S13. Accessed August 19, 2020. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6453604/

Chhatwal J, Chen Q, Ayer T, et al. Hepatitis C virus re-treatment in the era of direct-acting antivirals: projections in the USA. *Aliment Pharmacol Ther.* 2018;47(7):1023–1031. doi:10.1111/apt.14527

Chhatwal J, Chen Q, Bethea ED, Hur C, Spaulding AC, Kanwal F. The impact of direct-acting anti-virals on the hepatitis C care cascade: identifying progress and gaps towards hepatitis C elimination in the United States. *Aliment Pharmacol Ther*. 2019;50(1):66–74. doi:10.1111/apt.15291

Chhatwal J, Chen Q, Wang X, et al. Assessment of the feasibility and cost of hepatitis C elimination in Pakistan. JAMA Netw Open. 2019;2(5):e193613. doi:10.1001/jamanetworkopen.2019.3613

Chhatwal J, Samur S, Bethea ED, et al. Transplanting hepatitis C virus-positive livers into hepatitis C virus-negative patients with preemptive antiviral treatment: a modeling study. *Hepatology*. 2018;67(6):2085–2095. doi:10.1002/hep.29723

Cipriano LE, Liu S, Shahzada KS, Holodniy M, Goldhaber-Fiebert JD. Economically efficient hepatitis C virus treatment prioritization improves health outcomes. *Med Decis Making*. 2018;38(7):849–865. doi:10.1177/0272989X18792284

Dan C, Kaplowitz L. Update on hepatitis C screening and management: actions for emergency departments. *Curr Emerg Hosp Med Rep.* 2019;7(3):53–58. doi:10.1007/s40138-019-00183-4

Eckman MH, Ward JW, Sherman KE. Cost effectiveness of universal screening for hepatitis C virus infection in the era of directacting, pangenotypic treatment regimens. J Clin Gastroenterol Hepatol. 2019;17(5):930–939.E9. doi:10.1016/j.cgh.2018.08.080

El-Serag HB, Christie IC, Puenpatom A, Castillo D, Kanwal F, Kramer JR. The effects of sustained virological response to directacting anti-viral therapy on the risk of extrahepatic manifestations of hepatitis C infection. *Aliment Pharmacol Ther*. 2019;49(11):1442–1447. doi:10.1111/apt.15240

Ganesan M, Dagur RS, Makarov E, Poluektova LI, Kidambi S, Osna NA. Matrix stiffness regulate apoptotic cell death in HIV-HCV co-infected hepatocytes: importance for liver fibrosis progression. *Biochem Biophys Res Commun.* 2018;500(3):717–722. doi:10.1016/j.bbrc.2018.04.142

Ganesan M, Poluektova LY, Enweluzo C, Kharbanda KK, Osna NA. Hepatitis C virus-infected apoptotic hepatocytes program macrophages and hepatic stellate cells for liver inflammation and fibrosis development: role of ethanol as a second hit. *Biomolecules.* 2018;8(4):113. doi:10.3390/biom8040113

Ganesan M, Poluektova LY, Kharbanda KK, Osna NA. Human immunodeficiency virus and hepatotropic viruses co-morbidities as the inducers of liver injury progression. *World J Gastroenterol.* 2019;25(4):398–410. doi:10.3748/wjg.v25.i4.398

Geiger R, Steinert J, McElwee G, et al. A regional analysis of hepatitis C virus collaborative care with pharmacists in Indian Health Service facilities. *J Prim Care Community Health*. 2018;9. <u>doi:10.1177/2150132718807520</u>

Golden-Mason L, McMahan RH, Kriss MS, et al. Early and late changes in natural killer cells in response to ledipasvir/sofosbuvir treatment. *Hepatol Commun*. 2018;2(4):364–375. doi:10.1002/hep4.1166



Hagan LM, Kasradze A, Salyer SJ, et al. Hepatitis C prevalence and risk factors in Georgia, 2015: setting a baseline for elimination. *BMC Public Health*. 2019;19(Suppl 3):480. doi:10.1186/s12889-019-6784-3

Harrington PR, Komatsu TE, Deming DJ, Donaldson EF, O'Rear JJ, Naeger LK. Impact of hepatitis C virus polymorphisms on direct-acting antiviral treatment efficacy: regulatory analyses and perspectives. *Hepatology*. 2018;67(6):2430–2448. doi:10.1002hep.2963

Harrington PR, Komatsu TE, Sun H, Naeger LK. Hepatitis C virus RNA levels following virologic failure with direct-acting antivirals: implications for lower sensitivity diagnostic assays. *Clin Infect Dis.* Published online May 10,2019;70(2):237–330. doi:10.1093/cid/ciz385

Harris AM, Chokoshvili O, Biddle J, et al. An evaluation of the hepatitis C testing, care and treatment program in the country of Georgia's corrections system, December 2013–April 2015. *BMC Public Health*. 2019;19(Suppl 3):466. doi:10.1186/s12889-019-6783-4

Harris AM, Millman AJ, Lora M, Osinubi A, Lom J, Miller LS. Hepatitis B testing, care linkage, and vaccination coverage within a registry of hepatitis C infected patients. *Vaccine*. 2019;37(16):2188–2193. doi:10.1016/j.vaccine.2019.03.012

loannou GN. Reply to: "Individual surveillance using model-based hepatocellular carcinoma risk estimates in chronic hepatitis C patients after antiviral treatment" [letter to the editor]. *J Hepatol*. 2019;70(1):211–212. doi:10.1016/j.jhep.2018.09.026

loannou GN, Beste LA, Green PK, et al. Increased risk for hepatocellular carcinoma persists up to 10 years after HCV eradication in patients with baseline cirrhosis or high FIB-4 scores. *Gastroenterology*. Published online July 26, 2019;157(5):1264–1278.E4. doi:10.1053/j.gastro.2019.07.033

loannou GN, Feld JJ. What are the benefits of a sustained virologic response to direct-acting antiviral therapy for hepatitis C virus infection? *Gastroenterology*. 2019;156(2):446–460.E2. doi:10.1053/j.gastro.2018.10.033

loannou GN, Green P, Lowy E, Mun EJ, Berry K. Differences in hepatocellular carcinoma risk, predictors and trends over time according to etiology of cirrhosis. *PLoS One*. 2018;13(9):e0204412. doi:10.1371/journal.pone.0204412

loannou GN, Green PK, Berry K, Graf SA. Eradication of hepatitis C virus is associated with reduction in hematologic malignancies: major differences between interferon and direct-acting antivirals. *Hepatol Commun.* 2019;3(8):1124–1136. doi:10.1002/hep4.1389

Ioannou GN, Green PK, Beste LA, Mun EJ, Kerr KF, Berry K. Development of models estimating the risk of hepatocellular carcinoma after antiviral treatment for hepatitis C. *J Hepatol.* 2018;69(5):1088–1098. doi:10.1016/j.jhep.2018.07.024

Ji F, Wei B, Yeo YH, et al. Systematic review with meta-analysis: effectiveness and tolerability of interferon-free direct-acting antiviral regimens for chronic hepatitis C genotype 1 in routine clinical practice in Asia. *Aliment Pharmacol Ther.* 2018;47(5):550–562. doi:10.1111/apt.14507

Ji F, Yeo YH, Wei MT, et al. Sustained virologic response to direct-acting antiviral therapy in patients with chronic hepatitis C and hepatocellular carcinoma: a systematic review and meta-analysis. *J Hepatol.* 2019;71(3):473–485. doi:10.1016/j.jhep.2019.04.017

Ji Y, Dang X, Nguyen LNT, et al. Topological DNA damage, telomere attrition and T cell senescence during chronic viral infections. *Immun Ageing*. 2019;16:12. <u>doi:10.1186/s12979-019-0153-z</u>

Jones KM, Shelton ME, Soldano AC, Campbell J. Resolution of atypical lichen myxedematosus following successful treatment of chronic hepatitis C virus infection with sofosbuvir-velpatasvir combination therapy. *JAMA Dermatol.* 2018;154(9):1094–1096. doi:10.1001/jamadermatol.2018.2379

Joshi P, Atherton A. Sustained virologic response with 6 weeks or less of direct-acting antiviral therapy for chronic hepatitis C: experience at a Veterans Affairs healthcare system. *J Gastroenterol Hepatol*. Published online May 6, 2019;34(12):2173–2178. doi:10.1111/jgh.14703

Kanwal F, Kramer JR, Asch SM, Cao Y, Li L, El-Serag HB. Long-term risk of hepatocellular carcinoma in HCV patients treated with direct acting antiviral agents. *Hepatology*. Published online June 20, 2019;71(1):44–55. <u>doi:10.1002/hep.30823</u>

Kanwal F, Pyne JM, Tavakoli-Tabasi S, et al. A randomized trial of off-site collaborative care for depression in chronic hepatitis C virus. *Health Serv Res.* 2018;53(4):2547–2566. doi:10.1111/1475-6773.12758

Kasarala G, Choi S, Lopez K, Britt RB, Boatright C, Tillmann HL. Curing hepatitis C virus (HCV) after organ transplantation: increased risk of rejection following HCV elimination. *Transpl Infect Dis.* 2018;20(1):e12796. doi:10.1111/tid.12796



Kim HS, El-Serag HB. The epidemiology of hepatocellular carcinoma in the USA. *Curr Gastroenterol Rep.* 2019;21(4):17. doi:10.1007/s11894-019-0681-x

Kim HS, Yang JD, El-Serag HB, Kanwal F. Awareness of chronic viral hepatitis in the United States: an update from the National Health and Nutrition Examination Survey. *J Viral Hepat.* 2019;26(5):596–602. <u>doi:10.1111/jvh.13060</u>

Konerman MA, Beste LA, Van T, et al. Machine learning models to predict disease progression among veterans with hepatitis C virus. *PLoS One*. 2019;14(1):e0208141. doi:10.1371/journal.pone.0208141

Kramer JR, Puenpatom A, Erickson KF, et al. Real-world effectiveness of elbasvir/grazoprevir in HCV-infected patients in the US Veterans Affairs healthcare system. *J Viral Hepat.* 2018;25(11):1270–1279. doi:10.1111/jvh.12937

Kulik L, El-Serag HB. Epidemiology and management of hepatocellular carcinoma. *Gastroenterology*. 2019;156(2):477–491.E1. doi:10.1053/j.gastro.2018.08.065

Kushner T, Cohen J, Tien PC, Terrault NA. Evaluating women's preferences for hepatitis C treatment during pregnancy. *Hepatol Commun*. 2018;2(11):1306–1310. doi:10.1002/hep4.1264

Lara J, Teka MA, Sims S, Xia GL, Ramachandran S, Khudyakov Y. HCV adaptation to HIV coinfection. *Infect Genet Evol*. Published online July 31, 2018;65:216–225. doi:10.1016/j.meegid.2018.07.039

Li J, Gordon SC, Rupp LB, et al. Sustained virological response does not improve long-term glycaemic control in patients with type 2 diabetes and chronic hepatitis C. *Liver Int*. Published online December 20, 2018;39(6):1027–1032. doi:10.1111/liv.14031

Li J, Gordon SC, Rupp LB, et al. Sustained virological response to hepatitis C treatment decreases the incidence of complications associated with type 2 diabetes. *Aliment Pharmacol Ther.* Published online January 16, 2019;49(5):599–608. doi:10.1111/apt.15102

Li J, Zhang T, Gordon SC, et al. Impact of sustained virologic response on risk of type 2 diabetes among hepatitis C patients in the United States. *J Viral Hepat*. Published online February 25, 2018;25(8):952–958. doi:10.1111/jvh.12887

Lim AG, Qureshi H, Mahmood H, et al. Curbing the hepatitis C virus epidemic in Pakistan: the impact of scaling up treatment and prevention for achieving elimination. *Int J Epidemiol*. Published online January 3, 2018;47(2):550–560. doi:10.1093/ije/dyx270

Mahale P, Engels EA, Li R, et al. The effect of sustained virological response on the risk of extrahepatic manifestations of hepatitis C virus infection. *Gut.* 2018;67(3):553–561. <u>doi:10.1136/gutjnl-2017-313983</u>

Mahmud N, Kaplan DE, Taddei TH, Goldberg DS. Incidence and mortality of acute-on-chronic liver failure using two definitions in patients with compensated cirrhosis. *Hepatology*. 2019;69(5):2150–2163. <u>doi:10.1002/hep.30494</u>

Maier MM, Zhou XH, Chapko M, Leipertz SL, Wang X, Beste LA. Hepatitis C cure is associated with decreased healthcare costs in cirrhotics in retrospective Veterans Affairs cohort. *Dig Dis Sci*. 2018;63(6):1454–1462. <u>doi:10.1007/s10620-018-4956-0</u>

Mapakshi S, Kramer JR, Richardson P, El-Serag HB, Kanwal F. Positive predictive value of International Classification of Diseases, 10th revision, codes for cirrhosis and its related complications. *Clin Gastroenterol Hepatol.* 2018;16(10):1677–1678. doi:10.1016/j.cgh.2018.01.042

Méndez-Lagares G, Lu D, Chen C, et al. Memory T cell proliferation before hepatitis C virus therapy predicts antiviral immune responses and treatment success. *J Immunol.* 2018;200(3):1124–1132. doi:10.4049/jimmunol.1701364

Miller LS, Millman AJ, Lom J, et al. Defining the hepatitis C cure cascade in an urban health system using the electronic health record. *J Viral Hepat*. Published online September 10, 2019;27(1):13–19. doi:10.1111/jvh.13199

Millman AJ, Luo Q, Nelson NP, Vellozzi C, Weiser J. Missed opportunities for prevention and treatment of hepatitis C among persons with HIV/HCV coinfection. *AIDS Care*. Published online September 23, 2019;32(7):921–929. doi:10.1080/09540121.2019.1668533

Momin B, Millman AJ, Nielsen DB, Revels M, Steele CB. Promising practices for the prevention of liver cancer: a review of the literature and cancer plan activities in the National Comprehensive Cancer Control Program. *Cancer Causes Control*. Published online December 1, 2018;29(12):1265–1275. doi:10.1007/s10552-018-1094-0

Moon AM, Singal AG, Tapper EB. Contemporary epidemiology of chronic liver disease and cirrhosis. *Clin Gastroenterol Hepatol.* Published online ahead of print August 8, 2019. <u>doi:10.1016/j.cgh.2019.07.060</u>

Moorman AC, Rupp LB, Gordon SC, et al. Long-term liver disease, treatment, and mortality outcomes among 17,000 persons diagnosed with chronic hepatitis C virus infection: current chronic hepatitis cohort study status and review of findings. *Infect Dis Clin North Am.* 2018;32(2):253–268. doi:10.1016/j.idc.2018.02.002



Moorman AC, Xing J, Rupp LB, et al. Late diagnosis of hepatitis C virus infection, 2014–2016: continuing missed intervention opportunities. *Am J Manag Care*. 2019;25(8):369–374. Accessed August 19, 2020. https://www.ajmc.com/journals/issue/2019/2019-vol25-n8/late-diagnosis-of-hepatitis-c-virus-infection-20142016-continuing-missed-intervention-opportunities

Moorman AC, Xing J, Rupp LB, et al. Hepatitis B virus infection and hepatitis C virus treatment in a large cohort of hepatitis C-infected patients in the United States. *Gastroenterology*. 2018;154(3):754–758. doi:10.1053/j.gastro.2017.12.002

Mücke MM, Backus LI, Mücke VT, et al. Hepatitis B virus reactivation during direct-acting antiviral therapy for hepatitis C: a systematic review and meta-analysis. *Lancet Gastroenterol Hepatol*. Published online January 19, 2018;3(3):172–180. doi:10.1016/S2468-1253(18)30002-5

Mun EJ, Green P, Berry K, Ioannou GN. No difference between direct-acting antivirals for hepatitis C in hepatocellular carcinoma risk. *Eur J Gastroenterol Hepatol.* 2019;31(1):47–52. doi:10.1097/MEG.00000000001242

Nguyen LN, Zhao J, Cao D, et al. Inhibition of TRF2 accelerates telomere attrition and DNA damage in naïve CD4 T cells during HCV infection. *Cell Death Dis.* 2018;9(9):900. doi:10.1038/s41419-018-0897-y

Njei B, Esserman D, Krishnan S, et al. Regional and rural-urban differences in the use of direct-acting antiviral agents for hepatitis C virus: the veteran birth cohort. *Med Care*. 2019;57(4):279–285. doi:10.1097/MLR.00000000001071

Nwaohiri A, Schillie S, Bulterys M, Kourtis AP. Towards elimination of hepatitis C virus infection in children [note]. *Lancet Child* Adolesc Health. 2018;2(4):235–237. doi:10.1016/S2352-4642(18)30069-5

O'Connor SM, Mixson-Hayden T, Ganova-Raeva L, et al. Integrated HIV surveillance finds recent adult hepatitis B virus (HBV) transmission and intermediate HBV prevalence among military in uncharacterized Caribbean country. *PLoS One*. 2019;14(10):e0222835. doi:10.1371/journal.pone.0222835

Ottman AA, Townsend ML, Hashem MG, Britt RB. Impact of substance use disorder on the rate of sustained virological response in veterans with chronic hepatitis C treated with direct-acting antivirals. *Ann Pharmacother*. 2019;53(6):581–587. doi:10.1177/1060028018824988

Ottman AA, Townsend ML, Hashem MG, DiMondi VP, Britt RB. Incidence of drug interactions identified by clinical pharmacists in veterans initiating treatment for chronic hepatitis C infection. *Ann Pharmacother*. 2018;52(8):763–768. doi:10.1177/1060028018766507

Owens MD, Ioannou GN, Tsui JL, Edelman EJ, Greene PA, Williams EC. Receipt of alcohol-related care among patients with HCV and unhealthy alcohol use. *Drug Alcohol Depend*. 2018;188:79–85. <u>doi:10.1016/j.drugalcdep.2018.03.047</u>

Patterson S, Schmajuk G, Evans M, et al. Gaps in ambulatory patient safety for immunosuppressive specialty medications. *Jt Comm J Qual Patient Saf.* 2019;45(5):348–357. doi:10.1016/j.jcjq.2018.12.003

Reid M, Ma Y, Scherzer R, et al. Contribution of liver fibrosis and microbial translocation to immune activation in persons infected with HIV and/or hepatitis C virus. *J Infect Dis.* 2018;217(8):1289–1297. doi:10.1093/infdis/jix688

Rentsch CT, Cartwright EJ, Gandhi NR, et al. Provider verification of electronic health record receipt and nonreceipt of directacting antivirals for the treatment of hepatitis C virus infection. *Ann Epidemiol.* 2018;28(11):808–811. doi:10.1016/j.annepidem.2018.08.007

Rich NE, Yang JD, Perumalswami PV, et al. Provider attitudes and practice patterns for direct-acting antiviral therapy for patients with hepatocellular carcinoma. *Clin Gastroenterol Hepatol*. Published online July 26, 2019;18(4):974–983. doi:10.1016/j.cgh.2019.07.042

Rife K, Lyman A, LeClerc-Kamieniecki S, et al. Significant HbA1c lowering in patients achieving a hepatitis C virus cure. *Fed Pract.* 2019;36(Suppl 2):S26–S32.

Roberson JL, Lagasca AM, Kan VL. Comparison of the hepatitis C continua of care between hepatitis C virus/HIV coinfected and hepatitis C virus mono-infected patients in two treatment eras during 2008–2015. *AIDS Res Hum Retroviruses*. 2018;34(2):148–155. <u>doi:10.1089/AID.2017.0092</u>

Rogal SS, Beste LA, Youk A, et al. Characteristics of opioid prescriptions to veterans with cirrhosis. *Clin Gastroenterol Hepatol.* 2019;17(6):1165–1174.E3. doi:10.1016/j.cgh.2018.10.021

Rogal SS, Yakovchenko V, Waltz TJ, et al. Longitudinal assessment of the association between implementation strategy use and the uptake of hepatitis C treatment: year 2. *Implement Sci.* 2019;14(1):36. <u>doi:10.1186/s13012-019-0881-7</u>



Romano J, Sims OT, Richman J, et al. Resolution of ascites and hepatic encephalopathy and absence of variceal bleeding in decompensated hepatitis C virus cirrhosis patients. *JGH Open*. 2018;2(6):317–321. doi:10.1002/jgh3.12091

Samur S, Kues B, Ayer T, et al. Cost effectiveness of pre- vs post-liver transplant hepatitis C treatment with direct-acting antivirals. *Clin Gastroenterol Hepatol.* 2018;16(1):115–122.E10. <u>doi:10.1016/j.cgh.2017.06.024</u>

Sanaka S, Kasarala GR, Tillmann HL. A downside to hepatitis C virus cure? Vigilance is needed regarding hepatitis B virus reactivation, organ rejection, or hepatocellular carcinoma progression. *J Infect Dis.* 2018;217(6):857–860. doi:10.1093/infdis/jix659

Shaikh OS, Rogal S, Malik A, Sharma V, Cacciarelli T. Liver transplant from increased-risk donors in the era of direct-acting antivirals for hepatitis C. *Exp Clin Transplant*. Published online ahead of print July 19, 2019. doi:10.6002/ect.2019.0065

Singal AG, Lim JK, Kanwal F. AGA clinical practice update on interaction between oral direct-acting antivirals for chronic hepatitis C infection and hepatocellular carcinoma: expert review. *Gastroenterology*. 2019;156(8):2149–2157. doi:10.1053/j.gastro.2019.02.046

Singal AG, Rich NE, Mehta N, et al. Direct-acting antiviral therapy not associated with recurrence of hepatocellular carcinoma in a multicenter North American cohort study. *Gastroenterology*. 2019;156(6):1683–1692.E1. <u>doi:10.1053/j.gastro.2019.01.027</u>

Skolnik AA, Noska A, Yakovchenko V, et al. Experiences with interferon-free hepatitis C therapies: addressing barriers to adherence and optimizing treatment outcomes. *BMC Health Serv Res.* 2019;19(1):91. doi:10.1186/s12913-019-3904-9

Stephens D, Leston J, Terrault NA, et al. An evaluation of hepatitis C virus telehealth services serving tribal communities: patterns of usage, evolving needs, and barriers. *J Public Health Manag Pract.* 2019;25(Suppl 5):S97–S100. doi:10.1097/PHH.000000000001061

Struble K, Chan-Tack K, Qi K, Naeger LK, Birnkrant D. Benefit-risk assessment for sofosbuvir/velpatasvir/voxilaprevir based on patient population and hepatitis C virus genotype: U.S. Food and Drug Administration's evaluation. *Hepatology*. 2018;67(2):482–491. doi:10.1002/hep.29601

Stvilia K, Spradling PR, Asatiani A, et al. Progress in testing for and treatment of hepatitis C virus infection among persons who inject drugs—Georgia, 2018. *MMWR Morb Mortal Wkly Rep.* 2019;68(29):637–641. <u>doi:10.15585/mmwr.mm6829a2</u>

Su F, Ioannou GN. The impact of direct-acting antiviral therapy for hepatitis C on hepatocellular carcinoma risk. *Curr Hepatol Rep.* 2018;17(4):377–384. doi:10.1007/s11901-018-0424-8

Tapper EB, Parikh ND, Green PK, et al. Reduced incidence of hepatic encephalopathy and higher odds of resolution associated with eradication of HCV infection. *Clin Gastroenterol Hepatol.* Published online October 4, 2019;18(5):1197–1206.E7. doi:10.1016/j.cgh.2019.09.033

Tayob N, Christie I, Richardson P, et al. Validation of the hepatocellular carcinoma early detection screening (HES) algorithm in a cohort of veterans with cirrhosis. *Clin Gastroenterol Hepatol.* 2019;17(9):1886–1893.E5. <u>doi:10.1016/j.cgh.2018.12.005</u>

Tayob N, Richardson P, White DL, et al. Evaluating screening approaches for hepatocellular carcinoma in a cohort of HCV related cirrhosis patients from the Veteran's Affairs health care system. *BMC Med Res Methodol.* 2018;18(1):1. doi:10.1186/s12874-017-0458-6

Trickey A, Sood A, Midha V, et al. Clustering of hepatitis C virus antibody positivity within households and communities in Punjab, India. *Epidemiol Infect.* 2019;147:e283. doi:10.1017/s0950268819001705

Tsertsvadze T, Gamkrelidze A, Chkhartishvili N, et al. Three years of progress toward achieving hepatitis C elimination in the country of Georgia, April 2015–March 2018. *Clin Infect Dis.* Published online September 29, 2019:ciz956. doi:10.1093/cid/ciz956

Wang L, Cao D, Wang L, et al. HCV-associated exosomes promote myeloid-derived suppressor cell expansion via inhibiting miR-124 to regulate T follicular cell differentiation and function. *Cell Discov*. 2018;4:51. <u>doi:10.1038/s41421-018-0052-z</u>

Wei B, Ji F, Yeo YH, et al. Systematic review and meta-analysis: real-world effectiveness of direct-acting antiviral therapies in chronic hepatitis C genotype 3 in Asia. *BMJ Open Gastroenterol.* 2018;5(1):e000209. doi:10.1136/bmjgast-2018-000209

Wei B, Ji F, Yeo YH, et al. Real-world effectiveness of sofosbuvir plus ribavirin for chronic hepatitis C genotype 2 in Asia: a systematic review and meta-analysis. *BMJ Open Gastroenterol.* 2018;5(1):e000207. doi:10.1136/bmjgast-2018-000207

Wilder J, Choi SS, Moylan CA. Evaluation for fibrosis after cure of hepatitis C—reply. *JAMA*. 2019;321(15):1535. doi:10.1001/jama.2019.0873

Woodrell CD, Hansen L, Schiano TD, Goldstein NE. Palliative care for people with hepatocellular carcinoma, and specific benefits for older adults. *Clin Ther*. 2018;40(4):512–525. doi:10.1016/j.clinthera.2018.02.017



Woodward EN, Matthieu MM, Uchendu US, Rogal S, Kirchner JE. The health equity implementation framework: proposal and preliminary study of hepatitis C virus treatment. *Implement Sci.* 2019;14(1):26. <u>doi:10.1186/s13012-019-0861-y</u>

Yakovchenko V, Bolton RE, Drainoni ML, Gifford AL. Primary care provider perceptions and experiences of implementing hepatitis C virus birth cohort testing: a qualitative formative evaluation. *BMC Health Serv Res.* 2019;19(1):236. doi:10.1186/s12913-019-4043-z

Yakovchenko V, Gifford AL, Matthews KL, Greenstone CL, Tsai J, McInnes DK. Improving VHA's approach to community care: lessons learned from an imperfect hepatitis C choice program. *Med Care.* 2018;56(3):274–276.

Yee HS, Burton MJ, Belperio PS, Morgan TR. The Veterans Affairs hepatitis C treatment considerations. *Am J Gastroenterol.* 2019;114(2):185–188. doi:10.1038/s41395-018-0231-4

Yin S, Barker L, Teshale EH, Jiles RB. Rising trends in emergency department visits associated with hepatitis C virus infection in the United States, 2006–2014. *Public Health Rep.* Published online October 2, 2019;134(6):685–694. doi:10.1177/0033354919878437

Younossi ZM, Stepanova M, Asselah T, et al. Hepatitis C in patients with minimal or no hepatic fibrosis: the impact of treatment and sustained virologic response on patient-reported outcomes. *Clin Infect Dis.* 2018;66(11):1742–1750. doi:10.1093/cid/cix1106

Zamor PJ, Vierling J, Ghalib R, et al. Elbasvir/grazoprevir in Black adults with hepatitis C virus infection: a pooled analysis of phase 2/3 clinical trials. *Am J Gastroenterol.* 2018;113(6):863–871. doi:10.1038/s41395-018-0053-4

Zhao J, Dang X, Zhang P, et al. Insufficiency of DNA repair enzyme ATM promotes naive CD4 T-cell loss in chronic hepatitis C virus infection. *Cell Discov*. 2018;4:16. doi:10.1038/s41421-018-0015-4

Zou B, Yeo YH, Le MH, et al. Prevalence of viremic hepatitis C virus infection by age, race/ethnicity, and birthplace and disease awareness among viremic persons in the United States, 1999-2016. *J Infect Dis*. Published online September 27, 2019;221(3):408–418. doi:10.1093/infdis/jiz479

Zou WY, Choi K, Kramer JR, et al. Risk of hepatocellular cancer recurrence in hepatitis C virus+ patients treated with directacting antiviral agents. *Dig Dis Sci.* Published online April 30, 2019;64(11):3328–3336. <u>doi:10.1007/s10620-019-05641-3</u>

Hepatitis E

Averhoff F, Lazarus JV, Sergeenko D, et al. Excellence in viral hepatitis elimination—lessons from Georgia. *J Hepatol*. Published online July 26, 2019;71(4):645–647. doi:10.1016/j.jhep.2019.06.026

Landry ML, Kamili S, Jain D. Subacute liver failure due to autochthonous hepatitis E virus infection in an elderly man in the United States. *Hum Pathol (N Y).* 2018;12:68–70. <u>doi:10.1016/j.ehpc.2018.02.007</u>

GOAL 3: REDUCE VIRAL HEPATITIS HEALTH DISPARITIES

Hepatitis A

Foster M, Ramachandran S, Myatt K, et al. Hepatitis A virus outbreaks associated with drug use and homelessness—California, Kentucky, Michigan, and Utah, 2017. *MMWR Morb Mortal Wkly Rep.* 2018;67(43):1208–1210. doi:10.15585/mmwr.mm6743a3

Peak CM, Stous SS, Healy JM, et al. Homelessness and hepatitis A—San Diego County, 2016–2018. *Clin Infect Dis.* Published online August 15, 2019;71(1):14–21. doi:10.1093/cid/ciz788

Wilson E, Hofmeister MG, McBee S, et al. Notes from the field: hepatitis A outbreak associated with drug use and homelessness—West Virginia, 2018. *MMWR Morb Mortal Wkly Rep*. 2019;68(14):330–331. doi:10.15585/mmwr.mm6814a4

Hepatitis B

Le MH, Yeo YH, Cheung R, Henry L, Lok AS, Nguyen MH. Chronic hepatitis B prevalence among foreign-born and U.S.-born adults in the United States, 1999–2016. *Hepatology*. Published online June 22, 2019;71(2):431–443. doi:10.1002/hep.30831

Tang AS, Lyu J, Wang S, He Q, Pong P, Harris AM. Disparities in hepatitis B virus infection and immunity among New York City Asian American patients, 1997 to 2017. *Am J Public Health*. 2018;108(Suppl 4):S327–335. doi:10.2105/AJPH.2018.304504

Hepatitis C



Abara WE, Trujillo L, Broz D, et al. Age-related differences in past or present hepatitis C virus infection among people who inject drugs—National HIV Behavioral Surveillance, 8 US cities, 2015. *J Infect Dis.* Published online March 27, 2019;220(3):377–385. doi:10.1093/infdis/jiz142

Barnett PG, Joyce VR, Lo J, et al. Effect of interferon-free regimens on disparities in hepatitis C treatment of US veterans. *Value Health*. 2018;21(8):921–930. doi:10.1016/j.jval.2017.12.025

Byrne T, Troszak L, Midboe AM, et al. A novel measure to assess variation in hepatitis C prevalence among homeless and unstably housed veterans, 2011–2016. *Public Health Rep.* 2019;134(2):126–131. doi:10.1177/0033354918821071

Hofmeister MG, Edlin BR, Rosenberg ES, et al. Reply to HCV prevalence estimates among incarcerated persons [letter to the editor regarding HEP-18-1268.R1]. *Hepatology*. Published online April 4, 2019;70(2):759–760. doi:10.1002/hep.30635

Liang TJ, Ward JW. Hepatitis C in injection-drug users—a hidden danger of the opioid epidemic. *N Engl J Med.* 2018;378(13):1169–1171. doi:10.1056/NEJMp1716871

Mera J, Reilley B, Leston J, Stephens D. In a critical state: ongoing barriers to treatment for hepatitis C virus (HCV). *Am J Med*. 2019;132(5):547–549. doi:10.1016/j.amjmed.2018.10.031

Momin B, Mera J, Essex W, et al. Implementation of liver cancer education among health care providers and community coalitions in the Cherokee Nation. *Prev Chronic Dis.* 2019;16:E112. <u>doi:10.5888/pcd16.180671</u>

Ngo-Metzger Q, Mabry-Hernandez I, Heslin KC, Weiss AJ, Mummert A, Bierman AS. *Characteristics of Inpatient Stays Involving Hepatitis C, 2005–2014: HCUP Statistical Brief #232.* Agency for Healthcare Research and Quality. Published online December 5, 2017. Accessed August 19, 2020. www.hcup-us.ahrq.gov/reports/statbriefs/sb232-Hepatitis-C-Hospital-Stays-Trends.pdf

Reilley B, Leston J, Doshani M, et al. Assessing disparities in the rates of HCV diagnoses within American Indian or Alaska Native populations served by the U.S. Indian Health Service, 2005–2015. *J Community Health*. Published online May 28, 2018;43(6):1115–1118. doi:10.1007/s10900-018-0528-7

Reilley B, Miller M, Hudson M, Haverkate R, Leston J. Hepatitis C drug prescriptions and Medicaid policies--four states, Indian health care system, USA 2018. *Int J Equity Health*. 2019;18(1):190. doi:10.1186/s12939-019-1101-4

Zibbell JE, Asher AK, Patel RC, et al. Increases in acute hepatitis C virus infection related to a growing opioid epidemic and associated injection drug use, United States, 2004 to 2014. *Am J Public Health*. Published online January 10, 2018;2:e1–7. doi:10.2105/AJPH.2017.304132

GOAL 4: COORDINATE, MONITOR, AND REPORT ON IMPLEMENTATION OF VIRAL HEPATITIS ACTIVITIES

Hepatitis B

Popping S, Bade D, Boucher C, et al. The global campaign to eliminate HBV and HCV infection: International Viral Hepatitis Elimination Meeting and core indicators for development towards the 2030 elimination goals. *J Virus Erad.* 2019;5(1):60–66. Accessed August 19, 2020. <u>https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6362901/</u>

Hepatitis C

Longmire AG, Sims S, Rytsareva I, et al. GHOST: global hepatitis outbreak and surveillance technology. *BMC Genomics*. 2017;18(Suppl 10):916. doi:10.1186/s12864-017-4268-3

Ly KN, Miniño AM, Liu SJ, et al. Deaths associated with hepatitis C virus infection among residents in 50 states and the District of Columbia, 2016–2017. *Clin Infect Dis.* Published online October 5, 2019:ciz976. doi:10.1093/cid/ciz976

Popping S, Bade D, Boucher C, et al. The global campaign to eliminate HBV and HCV infection: International Viral Hepatitis Elimination Meeting and core indicators for development towards the 2030 elimination goals. *J Virus Erad.* 2019;5(1):60–66. Accessed August 19, 2020. <u>https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6362901/</u>

Popping S, El-Sayed M, Feld J, et al. Report from the International Viral Hepatitis Elimination Meeting (IVHEM), 17–18 November 2017, Amsterdam, the Netherlands: gaps and challenges in the WHO 2030 hepatitis C elimination framework. *J Virus Erad.* 2018;4(3):193–195. Accessed August 19, 2020. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6038125/





APPENDIX 3: ABBREVIATIONS

AAPI	Asian Americans and Pacific Islanders
AASLD	American Association for the Study of Liver Diseases
AETC	AIDS Education and Training Centers (HHS/HRSA)
AHRQ	Agency for Healthcare Research and Quality (HHS)
CDC	Centers for Disease Control and Prevention (HHS)
CDER	Center for Drug Evaluation and Research (HHS/FDA)
CMS	Centers for Medicare & Medicaid Services (HHS)
CNHS	Cherokee Nation Health Services
DAA	Direct-acting antiviral
DOJ	U.S. Department of Justice
DVH	Division of Viral Hepatitis (HHS/CDC)
ECHO	Extensions for Community Health Outcomes (HHS/HRSA)
EHR	Electronic health record
FBOP	Federal Bureau of Prisons (DOJ)
FDA	U.S. Food and Drug Administration (HHS)
FOA	Funding Opportunity Announcement
GHOST	Global Hepatitis Outbreak and Surveillance Technology
HAB	HIV/AIDS Bureau (HHS/HRSA)
HAV	Hepatitis A virus
HBeAG	Hepatitis B e antigen
HBsAg	Hepatitis B surface antigen
HBV	Hepatitis B virus
HCV	Hepatitis C virus
HDV	Hepatitis D virus
HEV	Hepatitis E virus
HHS	U.S. Department of Health and Human Services
HIT	Hepatic Innovation Team
HRSA	Health Resources and Services Administration (HHS)
HUD	U.S. Department of Housing and Urban Development
IDSA	Infectious Diseases Society of America
IDU	Injection drug use
IHS	Indian Health Service (HHS)
MAI	Minority AIDS Initiative (HHS/SAMHSA)
MAI-CoC	Minority AIDS Initiative Continuum of Care (HHS/SAMHSA)
MAT	Medication-assisted treatment



MMWR	Morbidity and Mortality Weekly Report
MSM	Men who have sex with men
NASEM	National Academies of Sciences, Engineering, and Medicine
NCI	National Cancer Institute (HHS/NIH)
NHANES	National Health and Nutrition Examination Survey
NHIS	National Health Interview Survey
NHLBI	National Heart, Lung, and Blood Institute (HHS/NIH)
NIAAA	National Institute on Alcohol Abuse and Alcoholism (HHS/NIH)
NIAID	National Institute of Allergy and Infectious Diseases (HHS/NIH)
NICHD	<i>Eunice Kennedy Shriver</i> National Institute of Child Health and Human Development (HHS/NIH)
NIDA	National Institute on Drug Abuse (HHS/NIH)
NIDDK	National Institute of Diabetes and Digestive and Kidney Diseases (HHS/NIH)
NIH	National Institutes of Health (HHS)
NIMHD	National Institute on Minority Health and Health Disparities (HHS/NIH)
NIS-Child	National Immunization Survey - Children
NNDSS	National Notifiable Disease Surveillance System
NVAC	National Vaccine Advisory Committee
NVPO	National Vaccine Program Office (HHS)
NVSS	National Vital Statistics System
OASH	Office of the Assistant Secretary for Health (HHS)
OIDP	Office of Infectious Disease and HIV/AIDS Policy (HHS/OASH)
ОМН	Office of Minority Health (HHS)
ONC	Office of the National Coordinator for Health Information Technology (HHS)
OPA	Office of Population Affairs (HHS/OASH)
OSG	Office of the Surgeon General (HHS/OASH)
OWH	Office on Women's Health (HHS/OASH)
PLWH	People living with HIV
PWH	People with HIV
RHA	Regional Health Administrator (HHS/OASH)
RWHAP	Ryan White HIV/AIDS Program (HHS/HRSA)
SAMHSA	Substance Abuse and Mental Health Services Administration (HHS)
SSP	Syringe services program
STI	Sexually transmitted infection
SUD	Substance use disorder
USPSTF	U.S. Preventive Services Task Force
VA	U.S. Department of Veterans Affairs
VHA	Veterans Health Administration (VA)
VHIG	Viral Hepatitis Implementation Group



VISN Veterans Integrated Services Network (VA)