

Health Plan Performance Improvement Project (PIP)

MCO Name: AmeriHealth Caritas

PIP Title: Improve Screening for Chronic Hepatitis C Virus (HCV) and Pharmaceutical Treatment Initiation

**PIP Implementation Period: January 1, 2020 –
December 31, 2021**

Submission Dates:

	Proposal / Baseline	Interim	Final
Version 1	3/11/2020	6/23/2020	12/31/2020
2021 Submission		6/30/2021	12/30/2021

MCO Contact Information

1. Principal MCO Contact Person

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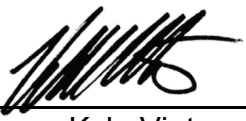
Attestation

Plan Name: AmeriHealth Caritas Louisiana

Title of Project: Improve Screening for Chronic Hepatitis C Virus (HCV) and Pharmaceutical Treatment Initiation

The undersigned approve this PIP and assure involvement in the PIP throughout the course of the project.

Medical Director Signature: Rodney Wise, MD.
First and Last Name: Rodney Wise, MD
Date: 12/30/2021

CEO Signature: 
First and Last Name: Kyle Viator
Date: 12/30/2021

Quality Director Signature: Rhonda Baird
First and Last Name: Rhonda Baird
Date: 12/30/2021

IS Director Signature (if applicable): Trampas Cranford
First and Last Name: Trampas Cranford
Date: 12/30/2021

Updates to the PIP

For Interim and Final Reports Only: Report all changes in methodology and/or data collection from initial proposal submission in the table below.

[EXAMPLES INCLUDE: ADDED NEW INTERVENTIONS, ADDED A NEW SURVEY, CHANGE IN INDICATOR DEFINITION OR DATA COLLECTION, DEVIATED FROM HEDIS® SPECIFICATIONS, REDUCED SAMPLE SIZE(S)]

Table 1: Updates to PIP

Change	Date of change	Area of change	Brief Description of change
Change 1	June 2020	<input checked="" type="checkbox"/> Project Topic <input checked="" type="checkbox"/> Methodology <input type="checkbox"/> Barrier Analysis / Intervention <input type="checkbox"/> Other	Under the guidance of USPSTF, AASLD, CDC, and LDH, the scope of the PIP changed from Increasing the Screening rates of the 1945-1965 cohort to include all Healthy Louisiana members aged 18-79 years old.
Change 2	February 2021	<input type="checkbox"/> Project Topic <input checked="" type="checkbox"/> Methodology <input type="checkbox"/> Barrier Analysis / Intervention <input type="checkbox"/> Other	ITM #1 – CM Outreach focused on utilizing the OPH listing of members with confirmed or probable HCV who are not receiving treatment
Change 3	April 2021	<input type="checkbox"/> Project Topic <input type="checkbox"/> Methodology <input checked="" type="checkbox"/> Barrier Analysis / Intervention <input type="checkbox"/> Other	Provider Education Update – LDH Hepatitis C Screening and Treatment Initiative, New CDC Guidelines, and Epclusa Treatment Option
Change 4	May 2021	<input type="checkbox"/> Project Topic <input type="checkbox"/> Methodology <input checked="" type="checkbox"/> Barrier Analysis / Intervention <input type="checkbox"/> Other	Enhanced Member Outreach – ACLA Newsletter to at-risk members, as well as a monthly texting campaign providing members with Screening and/or Treatment Information; Social Media Outreach – National Hepatitis Virus Month / Screening Day
Change 5	September 2020	<input type="checkbox"/> Project Topic <input type="checkbox"/> Methodology <input checked="" type="checkbox"/> Barrier Analysis / Intervention <input type="checkbox"/> Other	Enhanced Provider Outreach – Establish provider awareness of at-risk members associated with Provider Groups; QM virtual provider visits
Change 6	March 2021	<input type="checkbox"/> Project Topic <input type="checkbox"/> Methodology <input checked="" type="checkbox"/> Barrier Analysis / Intervention <input type="checkbox"/> Other	Publication of Universal Screening Gap in Care List to Provider Portal (NaviNet)

Abstract

For Final Report submission only. **Do not exceed 1 page.**

Title of Project: The Hepatitis C Virus (HCV) Performance Improvement Project

Rationale for Project/Objectives: The Hepatitis C Virus (HCV) Performance Improvement Project (PIP) is aimed to improve the Healthy Louisiana screening rate and initiation of HCV pharmaceutical treatment rate by ten percentage points. AmeriHealth Caritas Louisiana (ACLA) implemented a robust set of interventions to address two key intervention objectives:

- **Members:** Outreach and educate eligible members; and facilitate referrals and/or appointment scheduling with PCPs or HCV providers for screening and treatment.
- **Providers:** Educate providers on evidence-based recommendations and coordinate referrals for screening and treatment.

Methodology:

Performance Indicators

- **#1a (Universal Screening)** - The percentage of Healthy Louisiana enrollees ages 18-79 years, who were ever screened for HCV.
- **#1b (Birth Cohort Screening)** - The percentage of Healthy Louisiana enrollees for whom HCV screening is indicated by birth year between 1945 and 1965, and who were screened for HCV.
- **#2a (Non-Birth Cohort/Risk Factor Screening- ever screened)**- The percentage of Healthy Louisiana adults aged 18 and older for whom HCV screening is indicated by any one or more risk factors other than being born between 1945 and 1965, and who were ever screened for HCV.
- **#2b (Non-Birth Cohort/Risk Factor Annual Screening)**- The percentage of Healthy Louisiana adults aged 18 and older for whom HCV screening is indicated by any one or more risk factors other than being born between 1945 and 1965, and who were screened during the measurement year for HCV.
- **#3a (HCV Treatment Initiation-Overall)**-The percentage of the subset of adults with current or past drug use and with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per OPH listing, for whom pharmaceutical treatment for HCV was initiated.
- **#3b (HCV Treatment Initiation-Drug Users)** - The percentage of the subset of adults with current or past drug use and with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per OPH listing, for whom pharmaceutical treatment for HCV was initiated.
- **#3c (HCV Treatment Initiation-Persons with HIV)**- The percentage of the subset of adults ever diagnosed with HIV and with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per OPH listing, for whom pharmaceutical treatment for HCV was initiated.

Results:

ACLA's 2019 baseline data was calculated using the methodology listed above. Several internal departments collaborated to improve our baseline screening and treatment rates year to year. The target goal was a ten percentage point increase from the baseline rate for the seven performance indicators.

ACLA did not meet its 2021 target goals for all of the seven performance indicators; however, the goal was met for two of the performance indicators and four performance indicators demonstrated an increase from 2020 to 2021. Additionally, six of the seven performance indicators increased from the baseline year to the final reported year with two exceeding twenty percentage points.

Interventions:

Quality Management, Population Health Management (including Care Management), Enterprise Analytics and Provider Network Management Departments collaborated to initiate several member and provider focused interventions to increase our Hepatitis C Virus Screening and Treatment rates. Direct outreach was performed to members on the OPH HCV List received from LDH through Care Management and virtual provider visits with Quality and Provider Network Management. Additional member outreach was achieved utilizing the member newsletter and texting campaigns. Providers were outreached utilizing provider alerts and clinical practice guideline updates on the ACLA website.

Conclusions:

Considering the extraordinary barriers faced in 2021, e.g., the COVID-19 pandemic and various severe weather episodes, ACLA was able to make significant advances in our Hepatitis C Virus outreach, member screening rates, and Eplclusa treatment initiation. Based on the results, ACLA concludes that the Hepatitis C Virus PIP was successful. Performance indicators and intervention tracking measures were utilized to identify issues, resolve barriers, and promote best practices for eliminating Hepatitis C in Louisiana.

Next Steps:

The plan will continue to educate providers and members on the importance of appropriate treatment and screening guidelines, with an increased focus on treatment initiation in 2022.

Project Topic

To be completed upon Proposal submission. Do not exceed 2 pages.

Describe Project Topic and Rationale for Topic Selection

Describe how PIP Topic addresses your member needs and why it is important to your members:

The Hepatitis C Virus (HCV) is the most common blood-borne disease and the leading cause for liver transplant in the United States (LDH, 2019a). The prevalence of the anti-HCV antibody in the United States is approximately 1.6% in noninstitutionalized persons. According to data from 1999 to 2008, about three-fourths of patients in the United States living with HCV infection were born between 1945 and 1965, with a peak prevalence of 4.3% in persons aged 40 to 49 from 1999 to 2002 (USPSTF, 2013). HCV prevalence in Louisiana is estimated at 1.6% to 1.8%, with higher rates among urban residents, men and women aged 45-54 years (LA OPH, 2015). Louisiana ranks fifth in the U.S. for HCV/HIV co-infection; an estimated 6% of individuals with HCV in Louisiana are co-infected with HIV, and 18% of individuals with HIV as a result of intravenous drug use are also diagnosed with HCV co-infection (LA OPH, 2015).

- **Describe high-volume or high-risk conditions addressed:**

Many asymptomatic people are unaware that they are chronically infected with HCV, including those born between 1945 and 1965 (USPSTF, 2013). This contributes to significant delays in initiation of treatment and, as a result, can lead to clinical consequences. Therefore, the United States Preventative Services Task Force (USPSTF) recommends one-time HCV screening for all adults in this birth cohort (USPSTF, 2013). The USPSTF also recommends HCV screening for persons at high risk of chronic Hepatitis C infection (USPSTF, 2013):

- With Past or Current Injection Drug Use
- Persons Who Were Ever on Long-Term Hemodialysis
- Persons with a History of Incarceration
- Persons with HIV (AASLD/IDSA, 2018)

Persons born between 1945 and 1965 are more likely to be diagnosed with HCV infection, possibly because they received blood transfusions before the introduction of screening in 1992 or have a history of other risk for exposure decades earlier (USPSTF, 2013). A risk-based approach may miss detection of a substantial proportion of HCV-infected persons in the birth cohort because of a lack of patient disclosure or knowledge about prior risk status (USPSTF, 2013).

The most important risk factor for HCV infection is past or current injection drug use, with most studies reporting a prevalence of 50% or more (USPSTF, 2013). In 1998, the highest prevalence rates of the anti-HCV antibody occurred in persons with significant direct percutaneous exposure, such as injection drug users and persons with hemophilia (60% to 90%); persons with less significant percutaneous exposures involving smaller amounts of blood, such as patients receiving hemodialysis (10 to 30%), had more moderate prevalence rates (USPSTF, 2013).

- **Describe current research support for topic (e.g., clinical guidelines/standards):**

In February 2020, the American Association for the Study of Liver Diseases – Infectious Diseases Society of America (AASLD-IDSA) changed its recommendations regarding HCV Screening. These recommendations were updated to include recommended universal HCV screening for all adults aged 18 years or older followed by periodic testing for persons with ongoing risk behaviors and/or exposures (AASLD-IDSA, 2020). Independent studies using different modeling techniques demonstrated that one-time universal screening for adults 18 years or older is more cost effective compared to the birth-cohort screening (AASLD-IDSA, 2020). Along with this change, AASLD-IDSA still recommends risk-based HCV testing for members 18 years old with behaviors, exposures, or conditions or circumstances associated with an increased risk of HCV infection (AASLD-IDSA, 2020). Also periodic repeat HCV testing for persons with behavior, exposures, or conditions associated with an increased risk of HCV exposure, and annual HCV testing for all persons who inject drugs and for men with HIV infection who have unprotected sex with men (AASLD-IDSA, 2020).

- **Explain why there is opportunity for MCO improvement in this area (must include baseline and if available, statewide average/benchmarks):**

As of summer 2019, Healthy Louisiana enrollees, specifically our AmeriHealth Caritas Louisiana (ACLA) members, have access to safe and effective treatment for HCV. According to the baseline data for ACLA as of 12/31/2019, roughly 14,000 members fall within the 1945 – 1965 birth cohort. Through June 2019, only 629 of the 14,000 had been screened, and as of December 2019, roughly 1,100 ACLA members had been screened for the Hepatitis C Virus. Given the aim of this Performance Improvement Project (PIP) to improve the Healthy Louisiana HCV screening rate and initiation of HCV pharmaceutical treatment rate by ten percentage points from baseline to final measurement, there is immense potential for ACLA to improve in our HCV screening numbers when comparing the low number of members screened versus the high number of members who have not been screened.

For ACLA's members who are not within the 1945 to 1965 birth cohort but are at high risk for chronic HCV, e.g. persons with past or current drug use, persons ever on long-term hemodialysis, persons who were ever incarcerated, and persons ever diagnosed with HIV, the opportunity for improved screening rates is attainable. Our reports indicate a screening base of just over 10,000 members for whom HCV screening is indicated by any one or more of the aforementioned risk factors within the measurement year of 2019. Through June of 2019, roughly 600 members did receive a HCV screening, and a total of nearly 1,200 for the entire year.

Given these raw numbers mentioned above, ACLA members for whom HCV screening is indicated is roughly 25,000. Any improvements in our current screening rates will be beneficial for our at-risk members. Furthermore, with the current availability of pharmaceutical interventions provided by LDH, our Hepatitis C positive members have access to medication that was once not readily accessible. The authorized generic (AG) to which they have access is Epclusa, which has proven effective in curing 95% of persons living with HCV (LDH, 2019a). Epclusa is the preferred direct-acting antiviral (DAA) and does not require prior authorization unlike other available treatment regimens (LA Medicaid, 2019)

Aims, Objectives and Goals

Aim

Improve the Healthy Louisiana HCV screening rate and initiation of HCV pharmaceutical treatment rate by ten percentage points by implementing a robust set of interventions to address the following key intervention objectives:

1. **Member Intervention Objective:** Outreach and educate eligible members, and facilitate referrals to/schedule appointments with (I) PCPs for screening and (II) HCV providers (priority; per OPH database) or PCPs (per member preference) for treatment, with tailored interventions targeted to each of the following high risk subpopulations (which are not mutually exclusive, as enrollees may have multiple high risk characteristics)::
 - a. Beneficiaries born between the years 1945 and 1965
 - b. Current or past injection drug use
 - c. Persons ever on long term hemodialysis
 - d. Persons who were ever incarcerated
 - e. Persons with HIV infection
2. **Provider Intervention Objective:** Educate providers on evidence-based recommendations and availability of HCV specialty providers (USPSTF, 2013; AASLD/IDSA, 2018), and coordinate referrals for screening and treatment.

Table 2: Goals

Indicators	Baseline Rate ¹ Measurement Period: 1/1/19-12/31/19	Target Rate ²	Rationale for Target Rate ³
<p><u>Performance Indicator #1a</u> <u>(Universal Screening):</u></p> <p><i>The percentage of Healthy Louisiana enrollees ages 18-79 years {denominator} who were ever screened for HCV {numerator}.</i></p>	<p>N: 14,224 D: 91,922 R: 15.47%</p>	<p>R: 30.47%</p>	<p>15 Percentage Points For Maximum Proportion of Members That is Feasible</p>
<p><u>Performance Indicator #1b</u> <u>(Birth Cohort Screening):</u></p> <p><i>The percentage of Healthy Louisiana enrollees for whom HCV screening is indicated by birth year between 1945 and 1965 {denominator} and who were ever screened for HCV {numerator}.</i></p>	<p>N: 1,190 D: 13,956 R: 8.53%</p>	<p>R: 23.53%</p>	<p>15 Percentage Points For Maximum Proportion of Members That is Feasible</p>
<p><u>Performance Indicator #2a</u> <u>(Non-Birth Cohort/Risk Factor Screening- ever screened):</u></p> <p><i>The percentage of Healthy Louisiana adults aged 18 and older for whom HCV screening is indicated by any one or more risk factors other than being born between 1945 and 1965 {denominator} and who were ever screened for HCV {numerator}.</i></p>	<p>N: 1,137 D: 10,348 R: 10.99%</p>	<p>R: 25.99%</p>	<p>15 Percentage Points For Maximum Proportion of Members That is Feasible</p>
<p><u>Performance Indicator #2b</u> <u>(Non-Birth Cohort/Risk Factor Annual Screening):</u></p> <p><i>The percentage of Healthy Louisiana adults aged 18 and older for whom HCV screening is indicated by any one or more risk factors other than being born between 1945 and 1965 {denominator} and who were screened during the measurement year for HCV {numerator}.</i></p>	<p>N: 1,215 D: 11,717 R: 10.37%</p>	<p>R: 25.37%</p>	<p>15 Percentage Points For Maximum Proportion of Members That is Feasible</p>

Indicators	Baseline Rate ¹ Measurement Period: 1/1/19-12/31/19	Target Rate ²	Rationale for Target Rate ³
<p><u>Performance Indicator #3a</u> <u>(HCV Treatment Initiation-Overall):</u></p> <p><i>The percentage of all adults (ages 18 and older) with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per OPH listing {denominator} for whom pharmaceutical treatment for HCV was initiated {numerator}.</i></p>	<p>N: 495 D: 3,558 R: 13.91%</p>	<p>R: 28.91</p>	<p>15 Percentage Points For Maximum Proportion of Members That is Feasible</p>
<p><u>Performance Indicator #3b</u> <u>(HCV Treatment Initiation-Drug Users):</u></p> <p><i>The percentage of the subset of adults with current or past drug use and a confirmed or probable diagnosis of Chronic Viral Hepatitis C per OPH listing {denominator} for whom pharmaceutical treatment for HCV was initiated {numerator}.</i></p>	<p>N: 256 D: 1,981 R: 12.92%</p>	<p>R: 27.92%</p>	<p>15 Percentage Points For Maximum Proportion of Members That is Feasible</p>
<p><u>Performance Indicator #3c</u> <u>(HCV Treatment Initiation-Persons with HIV):</u></p> <p><i>The percentage of the subset of adults ever diagnosed with HIV and with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per OPH listing {denominator} for whom pharmaceutical treatment for HCV was initiated {numerator}.</i></p>	<p>N: 39 D: 226 R: 17.26%</p>	<p>R: 32.26%</p>	<p>15 Percentage Points For Maximum Proportion of Members That is Feasible</p>

¹ Baseline rate: the MCO-specific rate that reflects the year prior to when PIP interventions are initiated.

² Upon subsequent evaluation of performance indicator rates, consideration should be given to improving the target rate, if it has been met or exceeded at that time.

³ Indicate the source of the final goal (e.g., NCQA Quality Compass) and/or the method used to establish the target rate (e.g., 95% confidence interval).

Methodology

To be completed upon Proposal submission.

Performance Indicators

Table 3: Performance Indicators

Indicator	Description	Data Source	Eligible Population	Exclusion Criteria	Numerator	Denominator
<u>Performance Indicator #1a</u> <u>(Universal Screening)</u>	<i>The percentage of Healthy Louisiana enrollees ages 18-79 years {denominator} who were ever screened for HCV {numerator}.</i>	Administrative/ Claims/ Encounter data	All Healthy Louisiana enrollees ages 18-79 years	Healthy Louisiana adults with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per the Office of Public Health (OPH) listing	Number of Healthy Louisiana enrollees who were ever screened for HCV: CPT code 86803 OR CPT code 86804 OR CPT code 87520 OR CPT code 87521 OR CPT code 87522 OR HCPCS code G0472	Number of members in the eligible population less number of excluded members
<u>Performance Indicator #1b</u> <u>(Birth Cohort Screening)</u>	<i>The percentage of Healthy Louisiana enrollees for whom HCV screening is indicated by birth year between 1945 and 1965 {denominator} and who were screened for HCV {numerator}.</i>	Administrative/ Claims/ Encounter data	Healthy Louisiana enrollees born between 1945 and 1965	Healthy Louisiana adults with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per the Office of Public Health (OPH) listing	Number of Healthy Louisiana enrollees who were ever screened for HCV: CPT code 86803 OR CPT code 86804 OR CPT code 87520 OR CPT code 87521 OR CPT code 87522 OR HCPCS code G0472	Number of members in the eligible population less number of excluded members

Indicator	Description	Data Source	Eligible Population	Exclusion Criteria	Numerator	Denominator
<u>Performance Indicator #2a (Non-Birth Cohort/Risk Factor Screening- ever screened)</u>	<i>The percentage of Healthy Louisiana adults aged 18 and older for whom HCV screening is indicated by any one or more risk factors other than being born between 1945 and 1965 {denominator} and who were ever screened for HCV {numerator}.</i>	Administrative/ Claims/ Encounter data	<p>Healthy Louisiana adults aged 18 and older who were NOT born between 1945 and 1965, and who meet one or more of the following criteria:</p> <ul style="list-style-type: none"> a. Current or past injection drug use (ICD-9 or ICD-10 codes in Table A); OR b. Persons ever on long term hemodialysis (ICD-9 or ICD-10 codes in Table B); OR c. Persons who were ever incarcerated (ICD-9 or ICD-10 codes in Table C); OR <p>Persons ever diagnosed with HIV infection (ICD-9 or ICD-10 codes in Table d)</p>	Healthy Louisiana adults with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per the Office of Public Health (OPH) listing	Number of Healthy Louisiana enrollees who were ever screened for HCV: CPT code 86803 OR CPT code 86804 OR CPT code 87520 OR CPT code 87521 OR CPT code 87522 OR HCPCS code G0472	Number of members in the eligible population less number of excluded members

Indicator	Description	Data Source	Eligible Population	Exclusion Criteria	Numerator	Denominator
<u>Performance Indicator #2b (Non-Birth Cohort/Risk Factor Annual Screening)</u>	<i>The percentage of Healthy Louisiana adults aged 18 and older for whom HCV screening is indicated by any one or more risk factors other than being born between 1945 and 1965 {denominator} and who were screened during the measurement year for HCV {numerator}.</i>	Administrative/ Claims/ Encounter data	<p>Healthy Louisiana adults aged 18 and older who were NOT born between 1945 and 1965, and who meet one or more of the following criteria:</p> <ul style="list-style-type: none"> a. Current or past injection drug use (ICD-9 or ICD-10 codes in Table A); OR b. Persons ever on long term hemodialysis (ICD-9 or ICD-10 codes in Table B); OR c. Persons who were ever incarcerated (ICD-9 or ICD-10 codes in Table C); OR d. Persons ever diagnosed with HIV infection (ICD-9 or ICD-10 codes in Table d) 	Healthy Louisiana adults with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per the Office of Public Health (OPH) listing	Number of Healthy Louisiana enrollees who were screened during the measurement year for HCV: CPT code 86803 OR CPT code 86804 OR CPT code 87520 OR CPT code 87521 OR CPT code 87522 OR HCPCS code G0472	Number of members in the eligible population less number of excluded members

Indicator	Description	Data Source	Eligible Population	Exclusion Criteria	Numerator	Denominator
<u>Performance Indicator #3a</u> <u>(HCV Treatment Initiation-Overall)</u>	<i>The percentage of all adults (ages 18 and older) with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per OPH listing {denominator} for whom pharmaceutical treatment for HCV was initiated {numerator}.</i>	Administrative/ Claims/ Encounter data	Healthy Louisiana adults with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per the Office of Public Health (OPH) listing	None	Number of adults with a pharmaceutical claim for sofosbuvir/velpatasvir (the authorized generic (AG) of Epclusa ®) or other LDH-approved Hepatitis C Virus Direct Acting Antiviral Agent {DAA}	Number of members in the eligible population for Performance Indicator #3a
<u>Performance Indicator #3b</u> <u>(HCV Treatment Initiation-Drug Users)</u>	<i>The percentage of the subset of adults with current or past drug use and with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per OPH listing {denominator} for whom pharmaceutical treatment for HCV was initiated {numerator}.</i>	Administrative/ Claims/ Encounter data	Healthy Louisiana adults with current or past drug use (ICD-9 or ICD-10 codes in Appendix A) AND with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per the Office of Public Health (OPH) listing	None	Number of adults with a pharmaceutical claim for sofosbuvir/velpatasvir (the authorized generic (AG) of Epclusa ®) or other LDH-approved Hepatitis C Virus Direct Acting Antiviral Agent {DAA}	Number of members in the eligible population for Performance Indicator #3b
<u>Performance Indicator #3c</u> <u>(HCV Treatment Initiation-Persons with HIV)</u>	<i>The percentage of the subset of adults ever diagnosed with HIV and with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per OPH listing {denominator} for whom pharmaceutical treatment for HCV was initiated {numerator}.</i>	Administrative/ Claims/ Encounter data	Healthy Louisiana adults ever diagnosed with HIV (ICD-9 or ICD-10 codes in Appendix D) AND with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per the Office of Public Health (OPH) listing	None	Number of adults with a pharmaceutical claim for sofosbuvir/velpatasvir (the authorized generic (AG) of Epclusa ®) or other LDH-approved Hepatitis C Virus Direct Acting Antiviral Agent {DAA}	Number of members in the eligible population for Performance Indicator #3c

Data Collection and Analysis Procedures

Is the entire eligible population being targeted by PIP interventions? If not, why?

- The entire eligible population was targeted by PIP interventions.

Sampling Procedures

If sampling was employed (for targeting interventions, medical record review, or survey distribution, for instance), the sampling methodology should consider the required sample size, specify the true (or estimated) frequency of the event, the confidence level to be used, and the margin of error that will be acceptable.

- **Describe sampling methodology:** N/A

Data Collection

Describe who will collect the performance indicator and intervention tracking measure data (using staff titles and qualifications), when they will perform collection, and data collection tools used (abstraction tools, software, surveys, etc.). If a survey is used, indicate survey method (phone, mail, face-to-face), the number of surveys distributed and completed, and the follow-up attempts to increase response rate.

- **Describe data collection:**
 - AmeriHealth Caritas Louisiana's Enterprise Analytics (Informatics) Department collected data from claims/encounter files of all eligible members. Additional data sources may include the OPH HCV file distributed by LDH. Administrative data was collected as needed, quarterly, and annually. Intervention Tracking Measures (ITM) data was collected monthly utilizing claims/encounter data, clinical documentation software, and departmental tracking tools.

Validity and Reliability

Describe efforts used to ensure performance indicator and intervention tracking measure data validity and reliability. For medical record abstraction, describe abstractor training, inter-rater reliability (IRR) testing, quality monitoring, and edits in the data entry tool. For surveys, indicate if the survey instrument has been validated. For administrative data, describe validation that has occurred, methods to address missing data and audits that have been conducted.

- **Describe validity and reliability:**
 - Administrative data was collected by the Enterprise Analytics (Informatics) team. The process for verifying ITM data validity and reliability was conducted by quality associates within each department. Through the PDSA cycle, analysis was conducted to determine process improvements, strengths and opportunities.

Data Analysis

*Explain the data analysis procedures and, if statistical testing is conducted, specify the procedures used (note that hypothesis testing should only be used to test significant differences between **independent** samples; for instance, differences between health outcomes among sub-populations within the baseline period is appropriate). Describe the methods that will be used to analyze data, whether measurements will be compared to prior results or similar studies, and if results will be compared among regions, provider sites, or other subsets or benchmarks. Indicate when data analysis will be performed (monthly, quarterly, etc.).*

Describe how plan will interpret improvement relative to goal.

Describe how the plan will monitor intervention tracking measures (ITMs) for ongoing quality improvement (e.g., stagnating or worsening quarterly ITM trends will trigger barrier/root cause analysis, with findings used to inform modifications to interventions).

- **Describe data analysis procedures:**

- Procedures included an analysis of the comparability of baseline and re-measurement data, including factors that impacted validity. The results section of the PIP ensures that numerical data is accurate, clear, and easily understood. Interpretation involved looking at all possible explanations for results and factors that may have affected them. Historical circumstances were considered. Visual displays of data facilitate analysis and communicate results.

- **Describe how plan will interpret improvement relative to goal:**

- Data analysis will guide how well interventions are influencing performance indicator rates and outcomes. This data was assessed against established goals and drove decisions on effectiveness of change.

- **Describe how plan will monitor ITMs for ongoing QI:**

- ITMs were validated and monitored weekly and monthly as appropriate through trending, PDSA cycles, run charts, and other QI tools to analyze impact and effectiveness. The process for verifying ITM data validity and reliability was conducted by quality associates within each department.

(Tentative) PIP Timeline

Report the baseline, interim and final measurement data collections periods below.

Baseline Measurement Period:

Start date: 1/1/2019

End date: 12/31/2019

Submission of Proposal/Baseline Report Due: 2/3/2020

Interim/Final Measurement Period:

Start date: 1/1/2020

End date: 12/31/2020

Submission of 1st Quarterly Status Report for Intervention Period from 1/1/21-3/31/21 Due: 4/30/2021

Submission of 2nd Quarterly Status Report for Intervention Period from 4/1/21-6/30/21 Due: 7/31/2021

Submission of 3rd Quarterly Status Report for Intervention Period from 7/1/21-9/30/21 Due: 10/31/2021

Submission of Draft Interim Report Due: 12/10/2020

Submission of Final Interim Due: 12/31/2020

Final Measurement Period:

Start date: 1/1/2021

End date: 12/31/2021

Submission of Draft Final Report Due: 12/10/2021

Submission of Final Final Report Due: 12/31/2021

Barrier Analysis, Interventions, and Monitoring

Table 4: Alignment of Barriers, Interventions and Tracking Measures

Barrier 1: New Healthy Louisiana HCV Treatment Benefit May be Unknown to Enrollee. Method of Barrier Identification: IPRO HCV PIP Guidance Document.		2020				2021			
		Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4 INT
Intervention #1a to address barrier: Enhanced Case Management Outreach for HCV Treatment Initiation Planned Start Date: February 2020 Actual Start Date: February 2020	Intervention #1a tracking measure : N: # members with appointment scheduled with HCV specialist (in OPH database) or PCP for HCV treatment assessment/initiation D: # members with confirmed or probable HCV per OPH listing not receiving treatment	N: 28 D: 2,859 R: 0.98%	N: 9 D: 2,690 R: 0.33%	N: 56 D: 2,588 R: 2.16%	N: 9 D: 2,853 R: 0.32%	N: 11 D: 2,740 R: 0.40%	N: 22 D: 3,369 R: 0.65%	N: 10 D: 3,286 R: 0.30%	N: 6 D: 3,596 R: 0.17%
Intervention #1b to Address Barrier: Enhanced Case Management Outreach for HCV Screening / Treatment Initiation Planned Start Date: February 2020 Actual Start Date: February 2020	Intervention #1b Tracking Measure: N: # Members with Appointment Scheduled with HCV Specialist or PCP for HCV Screening / Treatment D: # Members from OPH List with Successful Contact by CM Outreach	N: 20 D: 76 R: 26.32%	N: 7 D: 18 R: 38.89%	N: 50 D: 288 R: 17.36%	N: 9 D: 148 R: 6.08%	N: 11 D: 124 R: 8.87%	N: 22 D: 176 R: 12.50%	N: 10 D: 111 R: 9.01%	N: 6 D: 20 R: 30.00%

Barrier 2: Asymptomatic Enrollees May Not Know They are Infected with HCV. Method of Barrier Identification: IPRO HCV PIP Guidance Document.		2020				2021			
		Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4
Intervention #2a to address barrier: Enhanced Case Management Outreach for HCV Screening Planned Start Date: July 2020 Actual Start Date: July 2020	Intervention #2a Tracking Measure: N: # members with appointment scheduled with PCP for HCV screening D: # members at risk for HCV per MCO claims/encounter data	N: N/A D: N/A R: N/A	N: N/A D: N/A R: N/A	N: 13 D: 12,277 R: 0.11%	N: 9 D: 12,610 R: 0.07%	N: 11 D: 13,135 R: 0.08%	N: 22 D: 13,623 R: 0.16%	N: 10 D: 14,068 R: 0.07%	N: 6 D: 14,004 R: 0.04%
Barrier 3: Providers May Not be Aware That Epclusa Does Not Require Prior Authorization. Method of Barrier Identification: CM Outreach Feedback / Analysis		2020				2021			
		Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4
Intervention #3a to address barrier: Provider education regarding SOFOSBUVIR-VELPATASVIR 400-100 (AG Epclusa: Preferred) prescription. Planned Start Date: January 2020 Actual Start Date: January 2020	Intervention #3a Tracking Measure: N: # members with SOFOSBUVIR-VELPATASVIR 400-100 (AG Epclusa: Preferred) dispensed D: # members with any DAA dispensed	N: 501 D: 508 R: 98.62%	N: 344 D: 354 R: 97.18%	N: 340 D: 358 R: 94.97%	N: 306 D: 316 R: 96.84%	N: 319 D: 324 R: 98.46%	N: 307 D: 313 R: 98.08%	N: 251 D: 254 R: 98.82%	N: 82 D: 84 R: 97.62%

Intervention #3b to Address Barrier DAA Treatment Initiation of OPH Confirmed / Probable Members Planned Start Date: February 2020 Actual Start Date: February 2020	N: # Members who Received First DAA Treatment Medication in 2020 D: # Members with Confirmed / Probable HCV on OPH List Not Receiving Treatment **This measure was retired for 2021.	N: 141 D: 2,859 R: 4.93%	N: 101 D: 2,690 R: 3.75%	N: 120 D: 2,588 R: 4.64%	N: 118 D: 2,451 R: 4.81%	Retired Measure	Retired Measure	Retired Measure	Retired Measure
Barrier 4: Member Unaware of Hepatitis C Virus Screening / Treatment Initiative Method of barrier identification: QM / CM Outreach Feedback / Analysis		2020				2021			
		Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4
Intervention #4a to address barrier: Enhanced Member Outreach to Increase Awareness of HCV Screening / Treatment Initiative via Mailed Member Newsletter Planned Start Date: July 2020 Actual Start Date: September 2020	Intervention #4a tracking measure: N: # of ACLA Members who were screened for Hepatitis C after Newsletter Mailed Date D: # of Members Mailed Newsletter with Hepatitis C Virus Screening / Treatment Education This measure was retired in 2020, one-time educational mailer completed.	N: N/A D: R:	N: N/A D: R:	N: N/A D: R:	N: 175 D: 10,895 R: 1.61%	Retired Measure	Retired Measure	Retired Measure	Retired Measure
Intervention #4b to address barrier: Enhanced Member Outreach to Increase Awareness of HCV Screening / Treatment Initiative via Texting Campaign Planned Start Date: July 2020 Actual Start Date: September 2020	Intervention #4b tracking measure: N: # of ACLA Members who were screened for Hepatitis C after Texting Campaign Began D: # of Members on the Texting Campaign Distribution List <ul style="list-style-type: none"> Text Messaging Campaign began Q4 2020. To account for claims lag, denominators are representative of the members who received a text message during the 3rd Month of the previous Quarter. 	N: N/A D: R:	N: N/A D: R:	N: N/A D: R:	N: 1 D: 69,837 R: 0.00%	N: 1,408 D: 67,412 R: 2.09%	N: 3,771 D: 69,933 R: 5.39%	N: 3,357 D: 70,033 R: 4.79%	N: 895 D: 70,444 R: 1.27%

Results

To be completed upon Baseline, Interim and Final Report submissions. The results section should present project findings related to performance indicators. ***Do not*** interpret the results in this section.

Table 5: Results

Indicator	Baseline Period 1/1/2019 – 12/31/2019	Interim Period 1/1/2020 – 12/31/2020	2021 Final Period 1/1/2021 – 11/30/21	Target Rate ¹
Performance Indicator #1a (Universal Screening): <i>The percentage of Healthy Louisiana enrollees ages 18-79 years {denominator} who were ever screened for HCV {numerator}.</i>	N: 14,224 D: 91,922 R: 15.47%	N: 18,590 D: 96,560 R: 19.25%	N: 23,809 D: 102,912 R: 23.14%	Rate: 30.47%
Performance Indicator #1b (Birth Cohort Screening): <i>The percentage of Healthy Louisiana enrollees for whom HCV screening is indicated by birth year between 1945 and 1965 {denominator} and who were ever screened for HCV {numerator}.</i>	N: 1,190 D: 13,956 R: 8.53%	N: 3,703 D: 14,261 R: 25.97%	N: 3,761 D: 13,123 R: 28.66%	Rate: 23.53%
Performance Indicator #2a (Non-Birth Cohort/Risk Factor Screening- ever screened): <i>The percentage of Healthy Louisiana adults aged 18 and older for whom HCV screening is indicated by any one or more risk factors other than being born between 1945 and 1965 {denominator} and who were ever screened for HCV {numerator}.</i>	N: 1,137 D: 10,348 R: 10.99%	N: 3,868 D: 12,610 R: 30.67%	N: 4,844 D: 14,004 R: 34.59%	Rate: 25.99%

Indicator	Baseline Period 1/1/2019 – 12/31/2019	Interim Period 1/1/2020 – 12/31/2020	2021 Final Period 1/1/2021 – 11/30/21	Target Rate ¹
<p><u>Performance Indicator #2b (Non-Birth Cohort/Risk Factor Screening-Annual Screening):</u> <i>The percentage of Healthy Louisiana adults aged 18 and older for whom HCV screening is indicated by any one or more risk factors other than being born between 1945 and 1965 {denominator} and who were screened during the measurement year for HCV {numerator}.</i></p>	<p>N: 1,215 D: 11,717 R: 10.37%</p>	<p>N: 1,134 D: 12,610 R: 8.99%</p>	<p>N: 2,033 D: 14,004 R: 14.52%</p>	<p>Rate: 25.37%</p>
<p><u>Performance Indicator #3a (HCV Treatment Initiation-Overall):</u> <i>The percentage of all adults (ages 18 and older) with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per the Office of Public Health (OPH) listing {denominator} for whom pharmaceutical treatment for HCV was initiated {numerator}.</i></p>	<p>N: 495 D: 3,558 R: 13.91%</p>	<p>N: 686 D: 3,793 R: 18.09%</p>	<p>N: 648 D: 4,924 R: 13.16%</p>	<p>Rate: 28.91%</p>
<p><u>Performance Indicator #3b (HCV Treatment Initiation-Drug Users):</u> <i>The percentage of the subset of adults with current or past drug use and with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per the Office of Public Health (OPH) listing {denominator} for whom pharmaceutical treatment for HCV was initiated {numerator}.</i></p>	<p>N: 256 D: 1,981 R: 12.92%</p>	<p>N: 394 D: 2,243 R: 17.57%</p>	<p>N: 406 D: 3,019 R: 13.45%</p>	<p>Rate: 27.92%</p>

Indicator	Baseline Period 1/1/2019 – 12/31/2019	Interim Period 1/1/2020 – 12/31/2020	2021 Final Period 1/1/2021 – 11/30/21	Target Rate ¹
Performance Indicator #3c (HCV Treatment Initiation- Persons with HIV): <i>The percentage of the subset of adults ever diagnosed with HIV and with a confirmed or probable diagnosis of Chronic Viral Hepatitis C per the Office of Public Health (OPH) listing {denominator} for whom pharmaceutical treatment for HCV was initiated {numerator}.</i>	N: 39 D: 226 R: 17.26%	N: 57 D: 216 R: 26.39%	N: 57 D: 264 R: 21.59%	Rate: 32.26%

¹ Upon subsequent evaluation of quarterly rates, consideration should be given to improving the target rate, if it has been met or exceeded at that time.

Discussion

To Be Completed Upon Interim / Final Report Submission –

The discussion section is for explanation and interpretation of the results.

Discussion of Results - 2021

Interpret the performance indicator rates for each measurement period, i.e., describe whether rates improved or declined between baseline and interim, between interim and final and between baseline and final measurement periods.

- **Performance Indicator #1a (Universal Screening)**: Universal screening increased 3.78 percentage points from the 2019 baseline MY to the 2020 interim MY (15.47% to 19.25%), and 3.89 percentage points from the 2020 interim MY to the final MY (19.25% to 23.14%). There was a 7.67 percentage point increase from the baseline to the final MY (15.47% to 23.14%). This measure showed sustained improvement across all measurement years, however, the target goal of 30.47% was not achieved.
- **Performance Indicator #1b (Birth Cohort Screening)**: Birth Cohort screening increased 17.44 percentage points from the 2019 baseline MY to the 2020 interim MY (8.53% to 25.97%), and 2.69 percentage points from the 2020 interim MY to the final MY (25.97% to 28.66%). There was a 20.13 percentage point increase from the baseline to the final MY (8.53% to 28.66%). This measure showed sustained improvement across all measurement years, the target goal of 23.53% was achieved.
- **Performance Indicator #2a (Non-Birth Cohort/Risk Factor Screening- ever screened)**: Non-Birth Cohort/Risk Factor Screening-ever screened increased 19.68 percentage points from the 2019 baseline MY to the 2020 interim MY (10.99% to 30.67%), and 3.92 percentage points from the 2020 interim MY to the final MY (30.67% to 34.59%). There was a 23.60 percentage point increase from the baseline to the final MY (10.99% to 34.59%). This measure showed sustained improvement across all measurement years, the target goal of 25.99% was achieved.
- **Performance Indicator #2b (Non-Birth Cohort/Risk Factor Screening- Annual Screening)**: Non-Birth Cohort/Risk Factor Screening-annual screening decreased 1.38 percentage points from the 2019 baseline MY to the 2020 interim MY (10.37% to 8.99%), and increased 5.53 percentage points from the 2020 interim MY to the final MY (8.99% to 14.52%). There was a 4.15 percentage point increase from the baseline to the final MY (10.37% to 14.52%). The target goal of 25.37% was not achieved for any of the measurement years.
- **Performance Indicator #3a (HCV Treatment Initiation-Overall)**: HCV Treatment Initiation-Overall increased 4.18 percentage points from the 2019 baseline MY to the 2020 interim MY (13.91% to 18.09%), and decreased 4.93 percentage points from the 2020 interim MY to the final MY (18.09% to 13.16%). There was a 0.75 percentage point decrease from the baseline to the final MY (13.91% to 13.16%). The target goal of 28.91% was not achieved for any of the measurement years.
- **Performance Indicator #3b (HCV Treatment Initiation-Drug Users)**: HCV Treatment Initiation-Drug Users increased 4.65 percentage points from the 2019 baseline MY to the 2020 interim MY (12.92% to 17.57%), and decreased 4.12 percentage points from the 2020 interim MY to the final MY (17.57% to 13.45%). There was a 0.53 percentage point increase from the baseline to the final MY (12.92% to 13.45%). The target goal of 27.92% was not achieved for any of the measurement years.

- **Performance Indicator #3c (HCV Treatment Initiation-Persons with HIV):** HCV Treatment Initiation-Persons with HIV increased 9.13 percentage points from the 2019 baseline MY to the 2020 interim MY (17.26% to 26.39%), and decreased 4.80 percentage points from the 2020 interim MY to the final MY (26.39% to 21.59%). There was a 4.33 percentage point increase from the baseline to the final MY (17.26% to 21.59%). The target goal of 32.26% was not achieved for any of the measurement years.

- **Explain and interpret the results by reviewing the degree to which objectives and goals were achieved.** Use your ITM data to support your interpretations.

Similar to our 2020 performance indicator results, the performance indicators that improved the most during the 2021 PIP were the 1945-1965 birthing cohort and the High Risk-Factors populations. While it is not the only contributing factor, it is important to note that the majority of ACLA's interventions were focused towards these two populations which may contribute to the significant improvement. Care Management outreach from the OPH HCV list provided by LDH included those most at risk, due to either their age or previous behavior. In addition, member newsletter mailings were sent to this birthing cohort, and the texting campaign targeted that same cohort. Performance Indicator #2a (Non-Birth Cohort/Risk Factor Screening- ever screened) measure showed sustained improvement across all measurement years, the target goal of 25.99% was achieved.

- **PIP Highlights:**

One of the most significant member barriers in this PIP was low adherence to medication treatment for HCV. The plan addressed this barrier by implementing the interventions listed below:

- Eplclusa/DAA Medication Adherence Texting Campaign to notify members of lapse in fill of HCV treatment medication (began Q4, 2021). 188 members have been texted thus far.
- Monthly texting campaign providing members with screening and/or treatment information (began Q4, 2020). 82,545 members received a text and 9,432 members were screened for HCV (11.43%) after the texting campaign began.
- Increased social media presence advocating for screening and treatment.

The top provider barrier for this PIP was provider awareness of screening/treatment guidelines for HCV. The plan addressed this barrier by implementing the interventions listed below:

- Virtual provider visits outlining screening and treatment initiative (82 completed by Quality Management)
- Publication of OPH List available in the Provider Portal
- HCV screening care gaps available in the Provider Portal
- Provider updates distributed and published providing education on LDH initiatives, new CDC Guidelines, and Eplclusa treatment option

It is important to note that for Performance Indicator #3C (HCV treatment overall, the rate did increase from the Baseline MY to Final MY (for the Final MY, claims are through 11/30/21).

Provider and Member Feedback:

Provider feedback is received by the plan via various methods including the Provider Satisfaction Survey and Quality Committee Meetings. The HCV PIP, as well as the HCV quarterly rates, are reviewed by the Committee and input is received from both internal and external providers. Provider feedback indicated that they were not aware of assigned members who are positive for HCV. Education was provided on utilization of the HCV screening care gap report to confirm test results as well as screening/treatment guidelines. During "Provider Outreach Visits", providers reported that they were very appreciative of the HCV screening care gap report that was added to the Provider Portal.

Member feedback is received from the member during direct member outreach by an ACLA Case Manager. Feedback is received on a variety of subjects including: social determinant of health needs, questions on Hepatitis medication, screening and treatment options. Member feedback is also received through the annual Member Satisfaction Survey. In addition, during ACLA's Member Advisory Council (MAC) meeting, members are asked to contribute to the development of health education programs to improve the member's quality of care.

- **What factors were associated with success or failure?** For example, in response to stagnating or declining ITM rates, describe any findings from the barrier analysis triggered by lack of intervention progress, and how those findings were used to inform modifications to interventions.

ACLA found direct contact with either members or providers was a factor most associated with success. This is evident in our care management success rate in making screening/treatment appointments compared to successful contact with members not engaged in care management.

When comparing the appointment rate with the OPH list and the successful contact amount, we saw tremendous rate increases when live contact was made with members. Similarly, we also found our greatest failure to be the results of the care management outreach due to the large amount of members who were Unable to Contact and those that chose to opt-out.

Limitations

As in any population health study, there are study design limitations for a PIP. Address the limitations of your project design, i.e., challenges identified when conducting the PIP (e.g., accuracy of administrative measures that are specified using diagnosis or procedure codes are limited to the extent that providers and coders enter the correct codes; accuracy of hybrid measures specified using chart review findings are limited to the extent that documentation addresses all services provided).

- **Were there any factors that may pose a threat to the internal validity the findings?** Definition and examples: *internal validity means that the data are measuring what they were intended to measure. For instance, if the PIP data source was meant to capture all children 5-11 years of age with an asthma diagnosis, but instead the PIP data source omitted some children due to inaccurate ICD-10 coding, there is an internal validity problem.*

Threats to the internal validity of the findings include care management/case management process due to the limitations of episodic documentation and data abstractions from the ACLA's integrated care management software. Also, the administrative measure accuracies that are specified using diagnosis or procedure codes are limited to the extent that providers and coders enter the correct codes and accuracy of the Office of Public Health file data.

- **Were there any threats to the external validity the findings?** Definition and examples: *external validity describes the extent that findings can be applied or generalized to the larger/entire member population, e.g., a sample that was not randomly selected from the eligible population or that includes too many/too few members from a certain subpopulation (e.g., under-representation from a certain region).*

The retrieval of data was affected by several factors. Care management outreach experienced several Unable to Contact members, as well as member opt-outs. Further, the COVID-19 Pandemic impacted our outreach for the year, and the various extreme weather events in Louisiana pulled our focus away from HCV Outreach and prioritized overall member care during these stressful events.

- **Describe any data collection challenges.**

Definition and examples: *data collection challenges include low survey response rates, low medical record retrieval rates, difficulty in retrieving claims data, or difficulty tracking case management interventions.*

The plan faced numerous challenges with data collection for process measures focused on case management / care management outreach. Limitations relative to the episodic documentation and data abstraction from the plan's integrated care management software may have resulted in under-represented Case Management / Care Management member interactions.

Next Steps

This section is completed for the Final Report. For each intervention, summarize lessons learned, system-level changes made and/or planned, and outline next steps for ongoing improvement beyond the PIP timeframe.

Table 6: Next Steps

Description of Intervention	Lessons Learned	System-Level Changes Made and/or Planned	Next Steps
Enhanced Case Management Outreach for HCV Screening and Treatment Initiation	The need to improve Care management/Case management process measure data accuracy due to limitations of episodic documentation and data abstractions from the plan's integrated care management software	System Changes are current being evaluated to improve data retrieval and accuracy	Monitor the data being received to ensure all data is complete and accurate
Enhanced Case Management Outreach for HCV Screening and Treatment Initiation	Unable to contact members are at risk for missing appointments, miss Case Management interventions well as educational opportunities	No system changes	Continue member outreach to the unable to contact population Increase Community Education face to face outreach to unable to contact members as available
CM Outreach – OPH List	Direct Contact is Pivotal	Explore New Ways to Achieve Contact with Members – Community Health Centers; Mobile Screening Options	Collaborate Further Within ACLA and Other MCOs on Direct Contact Interventions
Texting Campaigns/ Increased social media presence advocating for screening and treatment	Texting campaigns/social media posts are a good way outreach to members to promote education on HCV screening/treatment	No System Changes	Continue to promote HCV screening and treatment information via text/social media
Educational Outreach to Providers	Providers are not aware of the preferred medication regimen for HCV as well as they are not aware of screening and treatment guidelines	No System Changes	Continue to educate Providers on the importance of HCV screening and treatment guidelines
Provider Education – CP and Provider Newsletter	Direct Contact is Pivotal. Education is Available Through Email/ Website Only to Those Who Access Them	More Involvement with Provider Network Management on LDH Initiation/Screening/Treatment	Collaborate Further Within ACLA and Other MCOs to Initiate More Active Provider Interventions

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United States Preventive Services Task Force. Screening for Hepatitis C Virus Infection in Adults: U.S. Preventive Services Task Force Recommendation Statement. *Ann Intern Med.* 2013;159:349-357.

Table A: Current or past injection drug use (any one or more of diagnosis codes or diagnosis code combinations in this table, not restricted to place of service and not restricted to principal or primary diagnosis; note: a limitation of this measure is that ICD-9 and 10 codes do not specify injection vs. other route)

ICD-9 code or code combination	ICD-10 code or code combination	Description
	F11-	Opioid related disorders (Hyphen indicates that all codes within F11 should be included. This applies to all other ICD-10 and ICD-9 codes with hyphens that are listed in this table, as well.)
304.0-		Opioid dependence
304.7-		Opioid combined with other drug dependence
	F14-	Cocaine related disorders
304.2-		Cocaine dependence
	F15-	Other stimulant related disorders
304.4-		Amphetamine and other psychostimulant dependence
V69.8 AND 304.91		(other problems related to life style) AND (unspecified drug dependence continuous)
	Z72.89 AND F19.20	(other problems related to life style) AND (other psychoactive substance abuse, uncomplicated)

Table B. Persons ever on long term hemodialysis (any one or more of diagnosis codes in this table, not restricted to place of service and not restricted to principal or primary diagnosis)

ICD-9 code	ICD-10 code	Description
	Z49-	Encounter for care involving renal dialysis (Hyphen indicates that all codes within Z49 should be included. This applies to all other ICD-10 and ICD-9 codes with hyphens that are listed in this table, as well.)
	Z99.2	Dependence on renal dialysis

ICD-9 code	ICD-10 code	Description
V4511		Dependence on renal dialysis
V560 or V561 or V562 or V5631 or V5632 or V568		Encounter for care involving renal dialysis

Table C. Persons who were ever incarcerated (any one or more of diagnosis codes in this table, not restricted to place of service and not restricted to principal or primary diagnosis)

ICD-9 code	ICD-10 code	Description
	Z65.1	Imprisonment and other incarceration
	Z65.2	Problems related to release from prison

Table D. Persons ever diagnosed with HIV infection. (any one or more of diagnosis codes in this table, not restricted to place of service and not restricted to principal or primary diagnosis)

ICD-9 code	ICD-10 code	Description
	B20	Human immunodeficiency virus (HIV) disease
042		Human immunodeficiency virus (HIV) disease
	Z21	Asymptomatic human immunodeficiency virus (HIV) infection status
V08		Asymptomatic human immunodeficiency virus (HIV) infection status

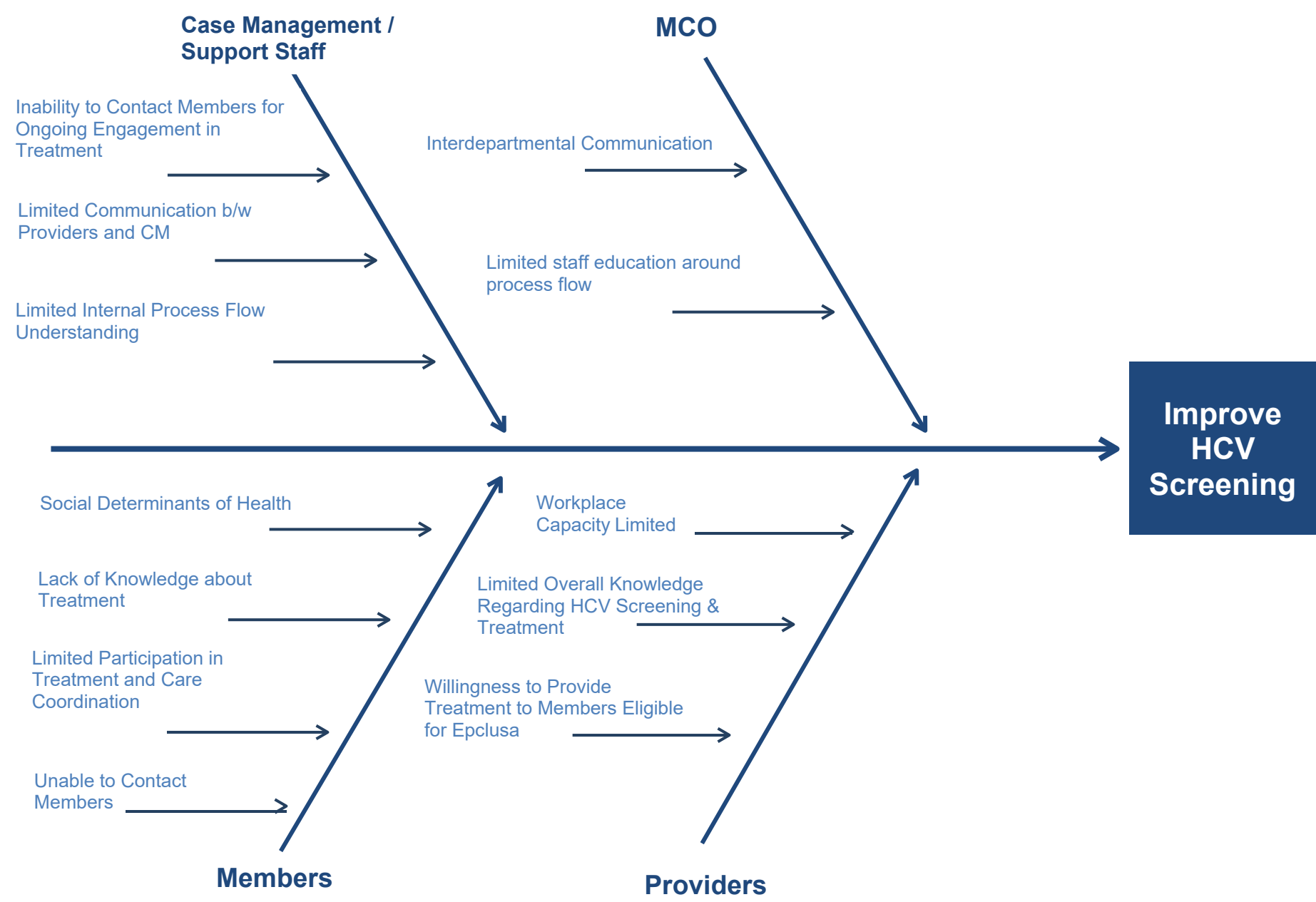
Glossary of PIP Terms

Table 7: PIP Terms

PIP Term	Also Known as...	Purpose	Definition
Aim	<ul style="list-style-type: none"> • Purpose 	To state what the MCO is trying to accomplish by implementing their PIP.	An aim clearly articulates the goal or objective of the work being performed for the PIP. It describes the desired outcome. The Aim answers the questions “How much improvement, to what, for whom, and by when?”
Barrier	<ul style="list-style-type: none"> • Obstacle • Hurdle • Road block 	To inform meaningful and specific intervention development addressing members, providers, and MCO staff.	Barriers are obstacles that need to be overcome in order for the MCO to be successful in reaching the PIP Aim or target goals. The root cause (s) of barriers should be identified so that interventions can be developed to overcome these barriers and produce improvement for members/providers/MCOs. A barrier analysis should include analyses of both quantitative (e.g., MCO claims data) and qualitative (such as surveys, access and availability data or focus groups and interviews) data as well as a review of published literature where appropriate to root out the issues preventing implementation of interventions.
Baseline rate	<ul style="list-style-type: none"> • Starting point 	To evaluate the MCO’s performance in the year prior to implementation of the PIP.	The baseline rate refers to the rate of performance of a given indicator in the year prior to PIP implementation. The baseline rate must be measured for the period before PIP interventions begin.
Benchmark rate	<ul style="list-style-type: none"> • Standard • Gauge 	To establish a comparison standard against which the MCO can evaluate its own performance.	The benchmark rate refers to a standard that the MCO aims to meet or exceed during the PIP period. For example, this rate can be obtained from the statewide average, or Quality Compass.
Goal	<ul style="list-style-type: none"> • Target • Aspiration 	To establish a desired level of performance.	A goal is a measurable target that is realistic relative to baseline performance, yet ambitious, and that is directly tied to the PIP aim and objectives.
Intervention tracking measure	<ul style="list-style-type: none"> • Process Measure 	To gauge the effectiveness of interventions (on a quarterly or monthly basis).	Intervention tracking measures are monthly or quarterly measures of the success of, or barriers to, each intervention, and are used to show where changes in PIP interventions might be necessary to improve success rates on an ongoing basis.

PIP Term	Also Known as...	Purpose	Definition
Limitation	<ul style="list-style-type: none"> • Challenges • Constraints • Problems 	To reveal challenges faced by the MCO, and the MCO's ability to conduct a valid PIP.	Limitations are challenges encountered by the MCO when conducting the PIP that might impact the validity of results. Examples include difficulty collecting/ analyzing data, or lack of resources / insufficient nurses for chart abstraction.
Performance indicator	<ul style="list-style-type: none"> • Indicator • Performance Measure (terminology used in HEDIS) • Outcome measure 	To measure or gauge health care performance improvement (on a yearly basis).	Performance indicators evaluate the success of a PIP annually. They are a valid and measurable gauge, for example, of improvement in health care status, delivery processes, or access.
Objective	<ul style="list-style-type: none"> • Intention 	To state how the MCO intends to accomplish their aim.	Objectives describe the intervention approaches the MCO plans to implement in order to reach its goal(s).

Appendix A: Fishbone (Cause and Effect) Diagram



Appendix B: Priority Matrix

Which of the Root Causes Are . . .	Very Important	Less Important
Very Feasible to Address	<ul style="list-style-type: none"> AmeriHealth Caritas Louisiana Interdepartmental Communication Regarding Initiative and Screening / Treatment Options Provider Education on Appropriate Evidence-Based Practices via ACLA Clinical Practice Guidelines Member Education and Awareness of New LDH Initiative 	<ul style="list-style-type: none"> Face to Face Provider Trainings
Less Feasible to Address	<ul style="list-style-type: none"> Member Unable to Contact Providers Unwilling to Initiate Treatment Protocol Fee Schedule Discrepancies Regarding RNA Testing and Reimbursement 	<ul style="list-style-type: none"> Locating Transient At-Risk Members Differentiating Between IV Drug Users and Drug Users

Appendix C: Strengths, Weaknesses, Opportunities, and Threats (SWOT) Diagram

	Positives	Negatives
INTERNAL under your control	<p>build on STRENGTHS</p> <p>Examples:</p> <ul style="list-style-type: none"> ▪ Medical Economics (Informatics) Reports Accessibility ▪ Case Management Linkage to At-Risk Members ▪ Ability to Outreach At-Risk Members on a Large Scale Communication Basis 	<p>minimize WEAKNESSES</p> <p>Examples:</p> <ul style="list-style-type: none"> ▪ Compliance Regarding Communicating with Members
EXTERNAL not under your control, but can impact your work	<p>pursue OPPORTUNITIES</p> <p>Examples:</p> <ul style="list-style-type: none"> ▪ Member Outreach Opportunities via Health-Fairs and Community Health Center Screening Events ▪ Provider Education Through Provider Network Management with Appropriate Evidence-Based Practice Guidelines 	<p>protect from THREATS</p> <p>Examples:</p> <ul style="list-style-type: none"> ▪ IV Drug User & Drug User Differentiation ▪ Unable to Contact Members ▪ Provider Participation ▪ Limited Workforce Capacity

Appendix D: Driver Diagram

Aims	Primary Drivers	Secondary Drivers	Specific Ideas for Interventions to Test/ Implement (Change Concepts)
Aim 1. Increase the HCV screening rates among Healthy Louisiana adults at risk for HCV by 10 percentage points from CY 2019 to CY 2020.	PCPs screen the following high risk Healthy Louisiana adults for HCV antibody: a. Beneficiaries born between the years 1945 and 1965 b. Beneficiaries with Current or past injection drug use c. Beneficiaries ever on long term hemodialysis d. Persons who were ever incarcerated e. Beneficiaries with HIV infection	Educate PCPs about evidence-based guidelines (EBGs) for HCV screening: -U.S. Preventive Service Task Force Guidelines -American Association for the Study of Liver Diseases (AASLD)/ Infectious Diseases Society of America (IDSA). -Office of Public Health streamlined test and treat strategy (forthcoming) -Medicaid reimbursable CPT/HCPCS codes	-Notify providers regarding Provider Portal access to HCV EBGs -Medical Director and Provider Relations face-to-face Outreach for Education -Incorporate USPSTF and AASLD/IDSA HCV screening guidelines into Clinical Practice Guideline repository -Disseminate Office of Public Health streamlined test and treatment strategy (forthcoming) -Develop and disseminate billing guidelines for HCV screening and Medicaid reimbursement - Encourage providers to participate in OPH-provided HCV treatment training [this covers screening as well]
		Identify adult members at risk for HCV	-Utilize HCV PIP specifications to identify at risk members using historical and current claims -Develop PCP lists of members eligible for screening -Develop Care Coordinator lists of members eligible for HCV screening
		Inform PCPs of their patients who are at risk/ eligible for screening	-Distribute to each PCP their listing of eligible members with instructions to contact patients to schedule an appointment for HCV screening
		Educate at risk members about HCV screening	-Care Coordinators Outreach, educate and council members at risk who are eligible for HCV screening
		Refer at risk members to PCPs and facilitate appointment scheduling for HCV screening	-Care Coordinators refer and schedule appointments with PCPs for HCV screening

Aims	Primary Drivers	Secondary Drivers	Specific Ideas for Interventions to Test/ Implement (Change Concepts)
Aim 2. Increase the HCV pharmaceutical treatment initiation rate among Healthy Louisiana adults ever diagnosed with HCV by 10 percentage points from CY 2019 to CY 2020.	HCV Providers identified in the OPH database (e.g., gastroenterologists, infectious disease specialists) and/or PCPs prescribe LDH-approved Hepatitis C Virus Direct Acting Antiviral Agent {DAA} for beneficiaries diagnosed with HCV	Educate PCPs about evidence-based guidelines (EBGs) for HCV diagnosis and treatment: -Office of Public Health streamlined test and treat guideline -American Association for the Study of Liver Diseases (AASLD)/ Infectious Diseases Society of America (IDSA).	-Provider Portal notification regarding access to HCV EBGs -Medical Director and Provider Relations face-to-face Outreach for Education -Incorporate the Office of Public Health streamlined test and treat guideline into Clinical Practice Guideline repository -Educate providers that prior authorization is not required for Epclusa generic for any Medicaid member -Develop and disseminate billing guidelines for HCV DAA agents and Medicaid reimbursement -Disseminate existing LDH resources to providers, including (1) the DAA Agent Medication Therapy Worksheet, (2) the HCV Treatment Agreement for Louisiana Medicaid Recipients, and (3) the Louisiana Medicaid Hepatitis C Direct-Acting Antiviral (DAA) Agents criteria, and (4) Office of Public Health (OPH) streamlined test and treatment guideline. - Encourage providers to participate in OPH-provided HCV treatment training
		Foster collaboration between PCPs, behavioral health and HCV specialists	-Develop and implement new processes to facilitate communication and coordinate care between PCPs, behavioral health and HCV providers listed in the OPH database (e.g., gastroenterologists, infectious disease specialists)
		Identify all members diagnosed with HCV	-Utilize the Office of Public Health listing of members with probable or confirmed HCV PIP to identify members with HCV diagnosis -Collaborate with OPH to develop PCP-specific listings of their patients who are potential candidates for HCV treatment -Develop Care Coordinator lists of members with HCV diagnosis for referral to PCPs for treatment
		Inform PCPs of their patients with HCV	-Distribute to each PCP their listing of members with HCV for medical assessment of appropriate treatment and/or referral to/ coordination with HCV specialist for treatment
		Educate and refer members with HCV for treatment assessment	-Care Coordinators Outreach, educate, refer and schedule member's appointment with HCV provider on OPH listing or PCP for treatment assessment.

Appendix E: Plan-Do-Study-Act Worksheet

	Pilot Testing	Measurement #1	Measurement #2
Intervention #1: Enhanced Case Management Outreach for HCV Screening and Treatment			
Plan: Document the plan for conducting the intervention.	<ul style="list-style-type: none"> Telephonic Outreach to Members with Confirmed / Probable HCV Who are Not Receiving Treatment to Assist with Appointment Scheduling 	<ul style="list-style-type: none"> Screening Appointments Scheduled by Case Management with HCV Specialist or PCP 	<ul style="list-style-type: none"> Treatment Appointments Scheduled by Case Management with HCV Specialist or PCP
Do: Document implementation of the intervention.	<ul style="list-style-type: none"> Implementation Began February 2020 	<ul style="list-style-type: none"> Implementation Began February 2020 	<ul style="list-style-type: none"> Implementation Began February 2020
Study: Document what you learned from the study of your work to this point, including impact on secondary drivers.	<ul style="list-style-type: none"> Direct Contact with Member Made Largest Impact on Success or Failure of Intervention 	<ul style="list-style-type: none"> Direct Contact with Member Made Largest Impact on Success or Failure of Intervention 	<ul style="list-style-type: none"> Direct Contact with Member Made Largest Impact on Success or Failure of Intervention
Act: Document how you will improve the plan for the subsequent phase of your work based on the study and analysis of the intervention.	<ul style="list-style-type: none"> 	<ul style="list-style-type: none"> Improve Contact Rate by Exploring Other Options – Health Clinics; Mobile Testing Units 	<ul style="list-style-type: none"> Improve Contact Rate by Exploring Other Options – Health Clinics; Mobile Testing Units
Intervention #2: Enhanced Member Outreach to Provide Education Regarding Hepatitis C Virus Initiative			
Plan: Document the plan for conducting the intervention.	<ul style="list-style-type: none"> Member Newsletter and Texting Campaign Outreach 	<ul style="list-style-type: none"> Screening Appointments Made After Receiving Text 	<ul style="list-style-type: none"> Screening Appointment Made After Receiving Newsletter
Do: Document implementation of the intervention.	<ul style="list-style-type: none"> 	<ul style="list-style-type: none"> 3rd Party Texting Initiative Based off of ACLA's Med Informatics At-Risk Report 	<ul style="list-style-type: none"> ACLA-Originated Delivery of HCV Newsletter
Study: Document what you learned from the study of your work to this point, including impact on secondary drivers.	<ul style="list-style-type: none"> 	<ul style="list-style-type: none"> 60-Day Claims Lag Affecting Current Results – Will Have Monthly Assessment Moving Forward 	<ul style="list-style-type: none"> 60-Day Claims Lag Affecting Current Results – Will Have Monthly Assessment Moving Forward
Act: Document how you will improve the plan for the subsequent phase of your work based on the study and analysis of the intervention.	<ul style="list-style-type: none"> 	<ul style="list-style-type: none"> We Expect to See a Screening Rate Increase in Contacted Members – Will Assess Further 	<ul style="list-style-type: none"> We Expect to See a Screening Rate Increase in Contacted Members – Will Assess Further