Role Of Physician Specialties To Close Gaps in the Care Cascade of Hepatitis C: Evidence From Paid Claims in the United States From 2010 To 2016

Darshan Mehta¹, Jeffrey McCombs¹, Yuri Sanchez Gonzalez², Steven Marx², Sammy Saab³

¹Schaeffer Center for Health Policy and Economics, University of Southern California; ²Health Economics and Outcomes Research, AbbVie Inc.; ³David Geffen School of Medicine, UCLA

BACKGROUND

- The Centers for Disease Control and Prevention [CDC] estimate the prevalence of chronic hepatitis C virus (HCV) in the United States (US) to be 1.3% (1.2%–2.4%), equivalent to 2.7–3.9 million individuals,¹ but only 10– 50 % are currently diagnosed.²
- It is recommended that all positive HCV antibody (AB) tests be followed by a HCV RNA blood test to determine the presence of active infection.
- National estimates indicate that only 27% of the population screened positive for HCV exposure has a confirmatory RNA test,² posing a significant gap in the cascade of care for hepatitis C.
- Limited referrals, linkage to care and access to treatment constitute other important gaps in the HCV care cascade.
- Despite the availability of novel oral direct-acting antiviral (DAA) regimens post 2013 that provide cure rates of up to 100% and improve patient reported outcomes,³⁻⁹ a national study estimated only 16% of patients with HCV were prescribed treatment.²
- Prior studies on key gaps in the care cascade were either conducted during the pre – DAA era⁹⁻¹¹ or restricted to specific individual centers, geographies⁹⁻¹³ or ethnic groups.¹⁴
- Furthermore, the role of physician specialties for closing key gaps in the HCV care cascade is poorly understood.

OBJECTIVES

• This study assesses AB screening rates, AB+ detection rates and linkage to care rates by physician specialty, and evaluates key determinants of HCV screening, detection, linkage to care and treatment in the US.

METHODS

Data source:

• Claims data from Optum Clinformatics[®] Data Mart, a de-identified database from the US, analysed over 2010–2016.

HCV AB Screening

- HCV screening was identified by paid claims for Current Procedural Terminology (CPT) codes 86803, 86804, or G0742.
- Screened patients having a diagnosis of HCV during the pre-index period were excluded from the analysis.
- Physician specialty for screening was based on the physician's order or bill for the index AB test.
- A cohort of unscreened patients was also analyzed using a randomly selected pool of patients without AB test.
- Index date was defined as the date of first observed AB test for the screened cohort and a randomly selected date for the unscreened cohort.
- All patients were required to be at least 20 years of age at index date and have at least 6 months of continuous enrollment pre-index.

HCV Diagnosis and Linkage to Care

- Screened patients who tested AB+ were identified and their access to treatment was assessed.
- The HCV diagnosis analysis focused on the subset of patients who received HCV AB screening and had non-missing lab data for the AB screening tests [LOINC codes (13955–0, 48159–8, 5198–7)] along with relevant test names.
- In addition to the earlier requirement of 6 months of pre-index data, patients in this analysis were further required to have minimum 6 months of continuous enrollment following the screening index date.
- Patients with missing or ambiguous test results were excluded from the analysis. The positive test results were identified if the numeric results were greater than 0.9¹⁵ or had character results mentioned as positive.
- Linkage to care and access to treatment was defined by presence of drug prescription following confirmed diagnosis. NDC codes and drug names were used to identify presence of treatment during the post-index period.

Multivariate analysis:

• Logistic regression was used to estimate the effect of physician specialty and time trend on the likelihood of being linked to care and receiving treatment, controlling for patient characteristics.

abbvie





- **HCV Screening**

HCV Diagnosis: AB+ Testing

- health care costs.
- like HIV and HBV.

Referral, Linkage to Care and Treatment

Table 1. Patient Characteristic At Baseline

Sample Size [N] Age [mean] Baby boomer popul Gender: Female^b

Race Asian

Black Hispanic

White Unknown

Missing

Medical cost prior 6 m Pharmacy cost prior Visits per month

FIB 4 Sample Size Mean FIB - 4 score F0-F1 F2 F3–F4

Table 2. Comorbidities at Baseline

Sample Size [N] Comorbidities Cardiovascular dise Type 2 diabetes me Nephritis, nephroti syndrome, nephros Chronic kidney dise Inflammatory bowe disease Fatigue Fibromyalgia Depression

Gastroesophageal disease

Mixed Cryoglobuli

Cirrhosis Hepatic compensat

Obesity

Coronary arterial di Cardiac arrest

HIV

HBV

Presented at the European Association for the Study of the Liver (EASL) • April 11–15, 2018 • Paris, France

RESULTS – Descriptive Statistics

 Baseline demographic characteristics are displayed in Table 1 and key comorbidities are displayed in Table 2.

• The demographic characteristics of screened and unscreened individuals were similar with some notable exceptions.

 Screened patients were more likely to be female, consumed more health care per month prior to their screening date, had fewer doctor visits and a higher risk of extrahepatic manifestations (EHMs) related to HCV such as cardiovascular disease (CVD), type 2 diabetes mellitus (T2DM), nephritis and chronic kidney disease (CKD); hepatic comorbidities like

cirrhosis and hepatic compensation; and coinfections like HIV and HBV.

• Compared to AB- patients, AB+ patients at baseline were more likely to be male and baby boomers, and have more doctor visits and greater

• A higher proportion of AB+ patients were in F2 stage or higher and had EHMs related to HCV such as CVD, T2DM, nephritis and CKD; hepatic comorbidities like cirrhosis, and hepatic compensation; and coinfections

• Treated patients were more likely to be males and less likely to have EHMs related to HCV such as CVD, T2DM, nephritis and CKD, and coinfections like HIV and HBV.

However, treated patients exhibited a higher proportion of hepatic related comorbidities like cirrhosis, hepatic compensation, suggesting prioritization of treatment to advanced liver disease stages.

	Likelihood of Screening		Likelihood of AB + Test		Likelihood of HCV treatment		
	Screened	Not Screened	AB+	AB-	Treated	Not Treated	
	1,056,583	1,243,581	12,578	507,490	2,399	5,853	
	46.3	46.5	51.5	43.2	56.1	54.8	
ation ^a	34.7%	35.5%	55.5%	29.1%	74.9%	61.7%	
	61.9%	51.3%	48.0%	64.1%	38.7%	44.0%	
	6.6%	3.7%	4.9%	7.5%	2.9%	6.7%	
	11.5%	7.3%	15.2%	12.6%	17.5%	16.2%	
	11.5%	9.5%	11.8%	13.4%	10.3%	12.8%	
	59.7%	58.8%	58.2%	56.6%	4.3%	3.8%	
	5.0%	4.6%	4.5%	5.0%	62.3%	57.0%	
	5.8%	16.1%	5.6%	4.9%	2.7%	3.5%	
nos.	\$6,173.49	\$4,674.87	\$7,266.14	\$4,272.89	\$4,157.31	\$8,114.14	
mos.	\$2 <i>,</i> 768.99	\$938.19	\$5,068.31	\$2,556.87	n.a.	n.a.	
	2.0	2.7	2.3	1.8	2.2	2.5	
FIB-4 Data for Sub-Sample with Laboratory Data							
	254,450		6,252	213,206	1,006	1,609	
	1.13		1.88	1.07	2.67	2.18	
	80.5%	_	57.1%	82.3%	37.1%	52.8%	
	16.6%	-	31.3%	15.4%	40.6%	32.5%	
	2.9%	_	11.6%	2.3%	22.3%	14.7%	

^aRest of the proportion represents non baby boomer population; ^bRest of the proportion represents male population

			-			
	Likelihood of Screening		Likelihood of AB + Test		Likelihood of HCV treatment	
	Not					Not
	Screened	Screened	AB+	AB-	Treated	Treated
	1,056,583	1,243,581	12,578	507,490	2,399	5,853
ase	6.9%	4.0%	8.3%	5.1%	11.1%	14.7%
llitus	12.9%	5.5%	16.9%	10.3%	22.4%	26.1%
c is	3.2%	1.0%	2.9%	1.7%	4.5%	9.8%
ase	2.9%	0.7%	2.4%	1.6%	3.9%	7.8%
el	1.3%	0.8%	1.2%	1.4%	1.6%	1.8%
	3.6%	1.2%	3.8%	3.4%	4.1%	4.6%
	4.7%	2.3%	4.8%	4.4%	5.4%	6.5%
	6.6%	3.9%	8.5%	5.8%	11.8%	11.9%
eflux	8.4%	5.1%	9.8%	7.4%	17.7%	15.8%
emia	0.0%	0.0%	0.1%	0.0%	0.3%	0.1%
	1.2%	0.3%	3.0%	0.8%	21.2%	13.3%
ion	1.5%	0.4%	3.8%	1.1%	23.8%	14.9%
	6.8%	3.6%	6.5%	6.4%	8.8%	8.5%
sease	2.2%	1.2%	2.4%	1.4%	4.0%	4.7%
	0.1%	0.1%	0.1%	0.0%	0.0%	0.3%
	5.5%	0.3%	7.3%	6.4%	7.9%	9.4%
	0.7%	0.1%	1.2%	0.7%	4.4%	9.1%

RESULTS

- Among the 1,056,583 total patients screened, most HCV AB tests were ordered by primary care physicians (PCPs; 29.7%), obstetricians/ gynaecologists (OB/GYNs; 19.1%) and nurses/physician assistants (PAs; 5.3%). (Figure 1)
- The mean AB+ detection rate across all studied specialties was 3% of screened patients, higher than the national AB+ prevalence of 1.7%.¹⁶ (Figure 2)
- While gastroenterologists, oncologists, and infectious disease their tests resulted in the highest observed AB+ detection rate (3.9%–5.3%).
- Linkage to care and treatment was low among AB+ patients at 13.8% overall, with hospitalist and gastroenterologists having the highest treatment rate of diagnosed patients (19.9%–25.0%). (Figure 3)
- Among the top three specialties ordering AB screening, OB/GYNs had the lowest rate of treated patients after AB+ diagnosis at 4.7%.

Figure 1: Proportion of Patients Screened By Physician Specialty



Figure 2: Proportion of Patients Testing AB+ By Physician Administering AB Test

PCPs: Primary care physicians; OB/GYN: Obstetrician/ Gynaecologist; PA: Physician Assistant; IDS: Infectious Disease Specialis



PCPs: Primary care physicians; OB/GYN: Obstetrician/ Gynaecologist; PA: Physician Assistant; IDS: Infectious disease Specialist; Note: Green dotted line represents sample average AB+ detection rate of 3%; Black dashed line represents national average of 1.7%¹⁶

specialists did not account for more than 3% of total patients screened,

Figure 3: Proportion of Patients Receiving Treatment By Index **Screening Physician Specialty**



PCPs: Primary care physicians; OB/GYN: Obstetrician/ Gynaecologist; PA: Physician Assistant; IDS: Infectious disease Specialist; Note: Dotted line represents sample average treatment rate of 13.8%; Dashed orange graph represents national average treatment rate of 16%

RESULTS – Logistic Regression

- Logistic regressions showed that being screened by gastroenterologists significantly increased the odds of treatment (odds ratio [OR]: 1.56) compared to PCP screening. (Table 3)
- On the other hand being screened by OB/GYN resulted in reduced odds of being treated (OR: 0.49) as compared to PCPs.
- Males (OR: 1.24) and baby boomers (OR: 2.23) were more likely to receive HCV treatment.
- The availability of novel DAAs in 2014 increased the treatment rate for AB+ patients (OR: 1.25).

Table 3: Logistic regression on likelihood of linkage to care and treatment among diagnosed HCV patients

Effect	Odds ratio	95% Wald Confidence Limits	
Age Categories (Ref: < 30)			
30 –45	1.314	1.004	1.719
45–65	2.237	1.761	2.841
>65	1.727	1.296	2.301
Physician specialty (Ref: Primary care Physician)			
OB/GYN	0.493	0.353	0.688
Nurse/PAs	0.805	0.604	1.074
Rheumatologist	0.593	0.383	0.918
Hospital	1.7	0.947	3.051
Gastroenterologist	1.567	1.251	1.964
Nephrologist	1.231	0.741	2.046
Infectious diseases specialist	1.184	0.784	1.79
Hematologist/oncologist	1.235	0.747	2.044
Pediatrician	0.777	0.305	1.984
Others/unknown	1.037	0.855	1.257
Other specialties	1.044	0.842	1.294
Time (Ref: 2010)			
2011–2013	1.102	0.884	1.375
≥2014	1.256	1.01	1.562
Gender (Ref: Female)			
Male	1.24	1.113	1.381
Comorbidities			
HIV	1.127	0.919	1.382
HBV	0.771	0.482	1.231
OB/GYN: Obstetrician and Gynaecologist: PA: Physician assis	tant		

THU-112

DISCUSSION

- Results from our study indicate, PCPs and OB/GYNs were more likely to administer an AB screening test, however they were less likely to prescribe hepatitis C treatment.
- This finding is consistent with prior studies indicating that only 28–32% of patients have been connected to specialty care.¹⁷
- Konerman et al. identifies PCPs as champions for attaining HCV eradication,¹⁷ given that treatment upon diagnosis could close important gaps in referral and linkage to care.
- However, results from our study show that treatment rates among PCPs remain low.
- With the recent introduction of simplified pangenotypic and short-duration treatments, the ability to prescribe could be expanded to PCPs, and OB/GYNs to improve treatment upon diagnosis.

LIMITATIONS

- Inclusion criteria rely on accurate identification of labs, lab results, CPT and ICD-9 codes, which have well known limitations.
- It is possible that not every antibody lab was captured, and therefore there may have been some patients whose first documented positive AB test was prior to 2010. We attempted to minimize this possibility by excluding any patient who had a hepatitis C ICD-9 or detectable viral load documented prior to their first positive AB test.
- Lastly, this study used only data from single large commercial payer, and thus the results may not be generalizable to other health systems and/or populations.

CONCLUSIONS

- PCPs and OB/GYNs are the gatekeepers for HCV screening accounting for nearly half of total AB tests.
- In spite of high AB+ detection rates across specialties, treatment rates remain low
- Increased efforts are needed to improve linkage to care and treatment, especially in PCP and OB/GYN settings.

DISCLOSURES AND CONFLICTS OF INTEREST

Design and study conduct for the study was approved by AbbVie, Inc. AbbVie Inc. participated in the interpretation of data, and review and approval of the abstract. All authors contributed to the development of the publication and maintained control over the final content.

Jeffrey McCombs: Received funds from AbbVie Inc. as a part of fellowship agreement between AbbVie and University of Southern California (USC) Sammy Saab: Consultant to and serves on speaker bureau for AbbVie, BMS, Gilead, Janssen, Merck

Darshan Mehta: Financially supported for graduate research work by AbbVie Inc. as a part of fellowship agreement between AbbVie Inc. and University of Southern California (USC) Steven Marx and Yuri Sanchez Gonzalez are employees of AbbVie Inc. and may own stocks and/or options of the

ACKNOWLEDGEMENTS

Editorial support for poster development was provided by Robert Dawson and funded by AbbVie

REFERENCES

company.

- https://www.cdc.gov/hepatitis/statistics/index.htm. Assessed August 2017 2. Yehia BR. Schranz AJ. Umscheld CA et al. The Treatment Cascade for Chronic Hepatitis C Virus Infection in the United States: A Systematic Review and Meta-Analysis. *PLoS ONE* 9(7): e101554. doi:10.1371/journal.pone.0101554.
- Saab S, Mehta DA, Hudgens S, et al. Effect of hepatitis C treatment with ombitasvir/paritaprevir/ritonavir + dasabuvir regimen on patient's health related quality of life: analysis of Phase 3a and phase 3b clinical trials. Journal of Hepatology. 66 (1), S736.
- 4. Younossi ZM, Stepanova M, Henry L, et al. Minimal impact of sofosbuvir and ribavirin on health related quality of life in Chronic Hepatitis C (CH-C). J Hepatol. 2014;60:741-747 Younossi Z, Stepanova M, Marcellin P, et al. Ledipasvir (LDV) and Sofosbuvir (SOF) Combination Improves Patient-
- Reported Outcomes (PRO) During Treatment Of Chronic Hepatitis C (CH-C) Patients: Results From The ION-1 Clinical Trial. Presented at International Liver Congress, London, England 2014. . Younossi et al. Treatment With Ledipasvir and Sofosbuvir Improves Patient-Reported Outcomes: Results From
- the ION-1,-2, and -3 Clinical Trials. HEPATOLOGY, Vol. 61, No. 6, 2015.
- Younossi ZM, Stepanova M, Henry L, et al. Effects of sofosbuvir-based treatment, with and without interferon on outcome and productivity of patients with chronic hepatitis C. *Clin Gastroenterol Hepatol*. 2013;12:1349-57. 8. Liu, Y., L. Larsen, S. Zeuzem, et al. "P0873: Ombitasvir/paritaprevir/ritonavir and dasabuvir with ribavirin (RBV) has minimal impact on health-related quality of life (HRPRO OUTCOMES) compared with placebo during 12-
- week treatment in treatment-naïve adults with chronic Hepatitis C (CHC)." Journal of Hepatology. 2015;62:S668. 9. Liu, Y., L. Larsen, M. Bourliere, et al. "P0856: Ombitasvir/paritaprevir/ritonavir and dasabuvir with ribavirin (RBV) has mild impact on Health-Related Quality of Life (HRPRO outcomes) compared with placebo during 12-week treatment in treatment-experienced adults with chronic hepatitis C (CHC)." Journal of Hepatology. 2015;62:S661.
- 10. Groom H, Dieperink E, Nelson DB, et al. Outcomes of a Hepatitis C Screening Program at a Large Urban VA Medical Center. J Clin Gastroenterol. 2008;42:97-106. 11. Young KL, Huang W, Horsburgh CR, et al. Eighteen- to 30-year-olds more likely to link to hepatitis C virus care: an
- opportunity to decrease transmission. Journal of Viral Hepatitis. 2016;23:274-281. 12. Putaka B, Mullen K, Birdi S, et al. The disposition of hepatitis C antibody-positive patients in an urban hospital. Journal of Viral Hepatitis. 2009;16:814-821.
- 13. Falade Nwulia O, Mehta SH, Lasola J, et al. Public health clinic-based hepatitis C testing and linkage to care in baltimore. *Journal of Viral Hepatitis*. 2016;23:366-374.
- 14. Schaeffer S, Khalili M. Resaons for HCV non-treatment in underserved African Americans: Implications for treatment with new therapeutics. Ann Hepatol. 2015;14(2):234-242. 15. http://www.dynacare.com/dos/dos2 show.php?uc=12618&db=h.
- 16. Rosenberg E et al. 2017. Estimation of State-Level Prevalence of Hepatitis C Virus Infection, US States and District of Columbia, 2010. Clin Infect Dis. 2017; Jun 1:64(11):157-1581.
- 17. Konerman et al. Hepatitis C Treatment and Barriers to Eradication. *Clin Transl Gastroenterol*. 2016;Sep;7(9):e193.