

Development of an Interdisciplinary Telehealth Model of Provider Training and Comprehensive Care for Hepatitis C and Opioid Use Disorder in a High-Burden Region

Jacqueline E. Sherbuk,^{1,®} Terry Kemp Knick,¹ Chelsea Canan,¹ Patrice Ross,¹ Bailey Helbert,¹ Eleanor Sue Cantrell,² Charlene Joie Cantrell,² Rachel Stallings,³ Nicole Barron,³ Diana Jordan,³ Kathleen A. McManus,¹ and Rebecca Dillingham¹

¹Division of Infectious Diseases and International Health, University of Virginia, Charlottesville, Virginia, USA, ²Lenowisco Health Department, Wise, Virginia, USA, and ³Virginia Department of Health, Division of Disease Prevention, Richmond, Virginia, USA

Background: Hepatitis C virus (HCV) and the opioid epidemic disproportionately affect the Appalachian region. Geographic and financial barriers prevent access to specialty care. Interventions are needed to address the HCV-opioid syndemic in this region.

Methods: We developed an innovative, collaborative telehealth model in Southwest Virginia featuring bidirectional referrals from and to comprehensive harm reduction (CHR) programs and office-based opioid therapy (OBOT), as well as workforce development through local provider training in HCV management. We aimed to (1) describe the implementation process of provider training and (2) assess the effectiveness of the telehealth model by monitoring patient outcomes in the first year.

Results: The provider training model moved from a graduated autonomy model with direct specialist supervision to a 1-day workshop with parallel tracks for providers and support staff followed by monthly case conferences. Forty-four providers and support staff attended training. Eight providers have begun treating independently. For the telehealth component, 123 people were referred, with 62% referred from partner OBOT or CHR sites; 103 (84%) attended a visit, 93 (76%) completed the treatment course, and 61 (50%) have achieved sustained virologic response. Rates of sustained virologic response did not differ by receipt of treatment for opioid use disorder.

Conclusions: Providers demonstrated a preference for an in-person training workshop, though further investigation is needed to determine why only a minority of those trained have begun treating HCV independently. The interdisciplinary nature of this program led to efficient treatment of hepatitis C in a real-world population with a majority of patients referred from OBOTs and CHR programs.

Keywords. Hepatitis C virus; Injection drug use; Opioid use disorder; Substance use disorder; Opioid treatment; Healthcare delivery; Nurse navigator; Public health.

Hepatitis C virus (HCV) disproportionately affects people living in the Appalachian region of the United States [1]. Infections with HCV have risen rapidly in this region, particularly among young people [2], driven by the opioid epidemic and injection drug use [3]. People who use drugs and those living in rural regions face significant barriers to HCV treatment and limited access to services [4–9], including in Southwest Virginia [10], which has resulted in very low HCV treatment uptake among rural Appalachian people who inject drugs (PWID) [11, 12].

The Journal of Infectious Diseases® 2020;222(S5):S354–64

© The Author(s) 2020. Published by Oxford University Press for the Infectious Diseases Society of America. This is an Open Access article distributed under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs licence (http://creativecommons.org/licenses/ by-nc-nd/4.0/), which permits non-commercial reproduction and distribution of the work, in any medium, provided the original work is not altered or transformed in any way, and that the work is properly cited. For commercial re-use, please contact journals.permissions@oup.com D0I: 10.1093/infdis/jiaa141

The presence of direct-acting antivirals (DAAs) has changed the landscape of HCV treatment. DAAs are well tolerated, highly effective, and require as few as 8 weeks of treatment, a stark contrast to earlier interferon-based regimens associated with frequent toxicities, limited effectiveness, and prolonged courses [13]. With the development of DAAs, national goals have been established for the elimination of HCV by 2030 [14]. Unfortunately, rather than moving toward elimination, new HCV infection rates continue to rise [15]. Modeling studies have demonstrated that even if access to DAAs were unlimited and unrestricted, goals for the elimination of HCV will not be achieved without targeting treatment to PWID [16] and incorporating comprehensive harm reduction (CHR) programs along with HCV treatment [17-19]. Although rural Appalachian PWID frequently interact with the health care system [12], they are rarely prescribed medication for HCV [12, 20], owing in part to provider reluctance to treat PWID [21].

Optimal HCV treatment requires care integrated with substance use disorder (SUD) treatment, access to CHR and social services, and close patient follow-up in a setting that is accessible to and

Correspondence: Jacqueline E. Sherbuk, PO Box 801379, Charlottesville, VA 22908 (jes2nk@ hscmail.mcc.virginia.edu).

supportive of vulnerable populations [22]. Treatment for HCV and opioid use disorder can be initiated concurrently, with high uptake of treatment for opioid use disorder, and high rates of sustained virologic response (SVR) [23]. Given the presence of simple and effective HCV treatment through DAAs, there has been a push to shift HCV care from specialists based at academic medical centers to primary care providers (PCPs) [24]. With appropriate training, PCPs can achieve HCV treatment outcomes similar to, if not better, than specialists [25, 26]. However, treatment by PCPs has been slow to gain traction [22, 27, 28]. PCPs continue to report barriers to treating HCV, including feeling uncomfortable treating HCV and lacking the requisite treatment knowledge [27, 29]. In addition, the ability of PCPs to provide comprehensive care may be limited, particularly in rural settings where only 3% of rural PCPs are waivered to prescribe buprenorphine for treatment of opioid use disorder [30]. Clinicians who provide office-based opioid therapy (OBOT) also report barriers to providing HCV therapy, despite regular contact with a high-risk and high-prevalence population [31]. New strategies to approach the HCV-opioid syndemic are needed.

The University of Virginia (UVA) is a tertiary care medical center that serves as a referral center for a large portion of the state. Following the traditional model of specialists providing HCV treatment, UVA has successfully treated patients through the gastroenterology and infectious disease specialty clinics with a high overall SVR rate [32] compared with other health systems in the Southeastern United States [33]. However, this success accounted only for those patients who had received diagnoses and been connected to our health system. State epidemiologic data demonstrate that the highest rates of HCV occur in Southwest Virginia [10], a region hundreds of miles distant from the UVA health system yet within our referral region (Figure 1). Unfortunately, the high-burden Southwestern region also has the state's lowest rates of HCV treatment [10] owing to a dearth of specialty providers. Therefore, a clear need exists for increased access to HCV and SUD care in Southwest Virginia.

We developed an interdisciplinary telehealth model of comprehensive care for HCV and SUD in Southwest Virginia with provider training for HCV treatment in a high-burden region (Figure 2). Through partnerships with specialists, PCPs, the Virginia Department of Health (VDH), local health departments with CHR services, clinics, and medication-assisted treatment (MAT) programs, we aim to provide patient-centered care through telehealth and to develop the capacity for local treatment of the HCV-opioid syndemic. Here we (1) describe the implementation of the model of care and (2) evaluate the effectiveness of the telehealth model in successfully treating HCV.

METHODS

Objectives

Our objectives were (1) to implement a comprehensive model of care for people with HCV that overcame existing barriers

and included care for SUD, access to CHR services, and connection to routine medical care through telehealth and provider training in HCV treatment and (2) to evaluate the effectiveness of the model in successfully treating HCV, and to assess for disparities in outcomes based on receipt of MAT, presence of an on-site nurse navigator, and Medicaid enrollment. This study was determined to be a quality improvement project, and it met criteria for nonhuman subject research by the UVA Health Sciences Research Institutional Review Board.

Model of Care Development

We developed a telehealth program of HCV care based on the foundation of a pre-existing HIV telehealth model for patients in Southwest Virginia who would not otherwise have access to specialty care. The Karen S. Rheuban Center for Telehealth at UVA (https://uvahealth.com/services/telemedicine) has provided telehealth services to rural Virginia for >2 decades. The robust network of telehealth access points and favorable state statutes relative to provision of and reimbursement for telehealth services [34] make telehealth an appealing option for delivery of HCV care to Southwest Virginia. Successful implementation of telehealth treatment of HCV has been previously demonstrated in the United States through the Extension for Community Healthcare Outcomes (ECHO) program [35]. The UVA team includes a portion of an infectious disease physician's time and a full-time clinical nurse coordinator. The infectious disease physician provides HCV treatment through telehealth and leads provider training. The nurse coordinator is responsible for receiving referrals for telehealth treatment, coordinating patient laboratory studies, imaging, and paperwork required to obtain medication, and guiding patients through the entire treatment process. The nurse coordinator performs community outreach and education to identify providers interested in providing HCV treatment and leads the training of clinician support staff.

Increasing Capacity Through Provider Training

The aim of provider training was to increase access to HCV care by training local PCPs on HCV treatment, with priority given to those who provide OBOT and those in highest need regions. After initial training, ongoing support and mentoring is delivered through monthly calls where clinical updates or difficult cases are discussed. The training model was developed iteratively, based on participant feedback. Provider training workshops and continuing medical education credit are funded by VDH.

Direct Patient Care Through Telehealth

Telehealth access points are located at the local health department and community health centers. The local health department has a colocated CHR program that includes the state's first syringe services program and an on-site public health nurse who provides patient navigation services. Both the health

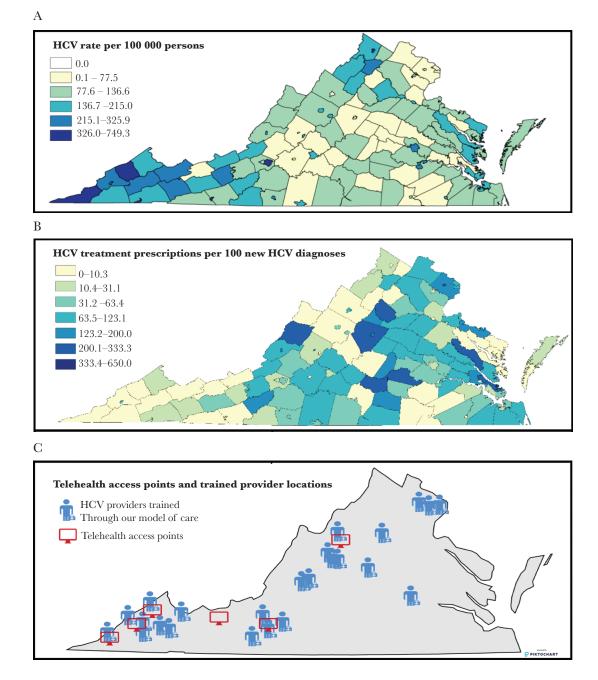


Figure 1. Distribution of hepatitis C virus (HCV) cases and treatment within Virginia and the development of access to HCV treatment through telehealth and provider training in areas or highest need. *A*, HCV case rate by city/county in Virginia for 2018, Virginia Department of Health, accessed August 2019. *B*, HCV treatment prescriptions per 100 new HCV diagnoses based on All-Payer Claims Databases for 2014 [10]. *C*, Access to HCV across the state of Virginia provided by our model of care through telehealth access points (n = 6) and providers (n = 36) who have completed HCV training through our program.

department and a local federally qualified health center have providers who provide OBOT and who wanted to receive training to treat HCV. Existing partnerships with specialty pharmacies were expanded to include provision of HCV treatment.

The only requirement for referral is a positive HCV antibody test and referral to a participating site (ClinicalTrials.gov NCT 21167). After referral, the nurse coordinator assists in managing required laboratory testing and paperwork, as well as performing liver staging using a mobile FibroScan (https://echosens.us/), because there is no local access to radiographic staging of liver disease. A VDH-sponsored program covers laboratory testing, because local health departments and other community-based screening programs rarely have financial support for confirmatory HCV viral load testing. The type of medication prescribed is influenced by the patients' insurance coverage and other medical conditions. For patients who are uninsured, the nurse

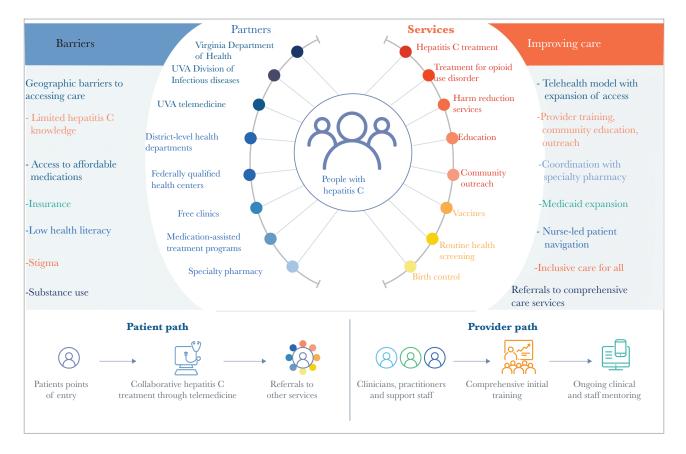


Figure 2. Model of comprehensive care for hepatitis C virus (HCV) and opioid use disorder (interdisciplinary telehealth model including provider training in a high-burden region). Abbreviation: UVA, University of Virginia.

coordinator assists in applying for Medicaid; if patients are determined to be noneligible, the nurse coordinator works with pharmaceutical patient assistance programs to ensure medication access. Medicaid was expanded in Virginia in January 2019 to include adults who earn <138% of the federal poverty level. Medicaid in Virginia covers DAAs regardless of liver fibrosis stage [36], although staging is still required for medication approval, given its importance for determining type and length of therapy and need for additional referrals.

Patients found to have advanced liver disease are referred to a hepatologist for comprehensive liver care [13]. All patients are offered hepatitis A and B vaccinations as well as referrals to additional services, including CHR, MAT, screening for sexually transmitted infections, pre-exposure prophylaxis for HIV, and family planning. Our model of care requires a total of 3 patient visits: the first is a nurse visit for completion of diagnostic studies; the second, a telehealth visit with the HCV provider; and the third, a follow-up nurse visit with viral load testing at 12 weeks after completion of therapy (SVR at 12 weeks [SVR12]), which serves as a determination of cure.

Effectiveness of the Telehealth Model

To monitor the effectiveness of this treatment model, we developed a cascade of care comprising the following steps: (1) referral for treatment, (2) attendance of initial telehealth specialist appointment, (3) completion of all diagnostic testing, including staging of liver disease, (4) prescription of medication, (5) insurance approval of medication, (6) medication started by the patient, (7) medication course completed, (8) posttreatment HCV viral load measured ≥ 12 weeks after treatment completion, and (9) achievement of SVR12.

Data Collection and Statistical Analyses

The population evaluated is all patients with active HCV referred to the telehealth treatment program during the program's initial 12 months (1 June 2018 to 31 May 2019). Follow-up data were measured through 15 February 2020. As part of routine clinic care, the nurse coordinator developed a clinical database tracking the cascade steps and date of completion, used to determine time to completion for each step. The following patient characteristics were obtained from the patient database: demographic variables, including age, sex, race, and ethnicity; insurance coverage; enrollment in and type of MAT; telehealth location; genotype and fibrosis score; and treatment, including type of medication prescribed and duration.

As part of the evaluation of the model, we aimed to compare progress along the cascade and time to completion of cascade steps by receipt of MAT, presence of an on-site nurse navigator, and Medicaid status, hypothesizing that each of these factors could influence HCV care. Proportions were compared using χ^2 tests with Fisher exact tests if indicated. Times to step completion were compared using the Mann-Whitney test, owing to nonnormal distribution.

RESULTS

Implementation—Provider Training

The provider training model required modification based on feedback from clinic teams (Figure 3). Initially, based on discussions with local PCPs, we developed a graduated autonomy model of training, similar to previously implemented models [37]. However, this model was unsuccessful owing to the practical challenges of coordinating schedules for observing and performing visits with the specialist, PCPs, and patients. Providers also indicated a strong preference for in-person as opposed to on-line asynchronous instruction. Therefore, we moved to a 1-day training workshop focusing on provider training. Although the workshop was well received, providers expressed concern about their ability to implement a new treatment program without trained support staff, leading to the third phase. In the current iteration, providers and support staff follow parallel tracks during a 1-day training session. Providers gain in-depth knowledge about the clinical management of HCV, while support staff learn the practical details related to the process of accessing medication and patient navigation.

Thirty-six providers and 8 support staff have completed HCV training workshops. They will serve patients in high-need regions of the state (Figure 1). Eight providers have begun treating patients independently, one located at the health department providing CHR services. Initial feedback suggests that enrollment of patients referred from CHR may be increasing owing to the flexible scheduling permitted by an on-site provider.

Patient Outcomes

A total of 123 referred patients with active HCV were included in analyses. An additional 8 patients were referred and found to have negative HCV viral loads indicative of spontaneous clearance and were therefore not included. The mean age was 40.5 years (standard deviation, 10.3 years), with the majority of patients <40 years old (Table 1). Nearly all patients were white and non-Hispanic. The most common insurance was Medicaid (63%), and a quarter of patients were uninsured. Thirty-one of 96 patients (33%) who underwent liver staging had advanced liver fibrosis. Seventy-six patients (62%) were enrolled in MAT, most commonly treated with buprenorphine-naltrexone. Most patients were prescribed glecaprevir-pibrentasvir for an 8-week duration.

Of the 123 referred patients, 103 (84%) linked to care by attending an appointment, 98 (80%) completed all required diagnostic testing, and 93 (76%) were prescribed medication

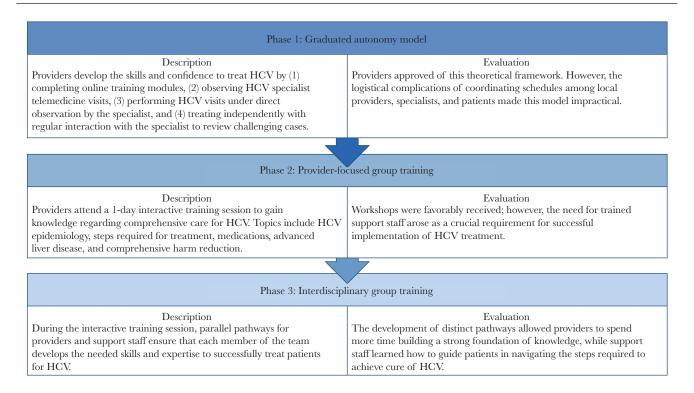


Figure 3. Iterative model of hepatitis C virus (HCV) provider training.

Table 1.	Hepatitis	C	Virus	Telehealth	Model	Patient	Characteristics
(N = 123)							

Characteristic	Patients, No. (%) ^a
Age, mean (SD), y	40.5 (10.3)
Age group, y	
20–29	18 (15)
30–39	46 (37)
40–49	37 (30)
50–59	14 (11)
≥60	8 (7)
Sex	
Female	57 (46)
Male	66 (54)
Race/ethnicity	
White, non-Hispanic	122 (99)
Black, non-Hispanic	1 (1)
Insurance status	
Medicaid	78 (63)
Medicare	8 (7)
Private	6 (5)
Uninsured	31 (25)
MAT	
Any MAT	76 (62)
Buprenorphine-naloxone	65 (53)
Methadone	5 (4)
Naltrexone (extended release)	5 (4)
Buprenorphine	1 (1)
None	47 (38)
Telemedicine location	
Health departments	
FQHC	67 (54)
Free clinic	44 (36)
HCV genotype	12n(10)
1	84 (68)
2	15 (12)
3	19 (15)
Unable to determine ^b	5 (4)
Fibrosis score (n = 96)	
F0/F1	59 (61)
F2	6 (6)
F3	17 (18)
F4	14 (15)
Medication prescribed ^c (n = 93)	
Glecaprevir-pibrentasvir	73 (78)
Ledipasvir-sofosbuvir	19 (21)
Sofosbuvir-velpatasvir	2 (2)
Treatment duration ^{c} (n = 93)	
8 wk	74 (80)
12 wk	19 (20)

Abbreviations: FQHC, federally qualified health center; HCV, hepatitis C virus; MAT, medication-assisted treatment; SD, standard deviation.

^aData represent no. (%) of patients unless otherwise specified

^bGenotype could not be determined if viral load was below threshold required for genotype testing.

^cThe type of medication prescribed and the duration of treatment were determined by the patients' insurance coverage and other medical conditions.

(Figure 4). Of the 5 patients who completed diagnostic testing but were not prescribed medication, 2 were incarcerated, 2 were lost to follow-up, and 1 declined to apply for Medicaid to access

treatment. Medication was approved, initiated, and completed for all those prescribed treatment. Of the 93 patients who completed treatment, 63 have obtained a posttreatment viral load measurement. For 8 patients, posttreatment SVR measurement is not due yet. SVR was achieved in 61 patients, representing 50% of all referred patients and 97% of those evaluated for SVR. There were 2 patients with virologic failure.

Patients receiving MAT progressed along the cascade at rates equivalent to those of patients not receiving MAT, and times to completion of individual steps were similar as well (Table 2). Patients receiving care at a telehealth access location with an on-site nurse navigator had a 50% reduction in time from referral to initial visit compared with those without an on-site nurse navigator (median [interquartile range], 11 [3–27.5] days vs 22 [12–46] days, respectively; P = .001). Progression along the cascade and time to completion of individual steps was similar for patients receiving Medicaid compared with all others.

DISCUSSION

This telehealth program in a state with Medicaid expansion and a favorable statutory environment for telehealth led to efficient linkage to HCV care, appropriate clinical evaluation, and prescription and procurement of medication among a real-world population with previously limited access to HCV care and high rates of SUD. The results of the first year of this program demonstrate that HCV treatment through telehealth can successfully reach high-risk populations when a comprehensive collaboration is developed that includes (1) coordination with the state health department to target programming and funding to the areas most affected by the HCV-opioid syndemic; (2) appropriate clinical staffing, including nurse coordinators; and (3) partnerships with local health departments, SUD treatment centers, and safety net clinics.

Overall linkage to care occurred at a high rate, particularly for a real-world model of care targeting a population that historically has faced geographic barriers and limited access to HCV care [32, 38–41]. Telehealth technology combined with a full-time nurse coordinator was able to overcome these barriers by decreasing geographic isolation and facilitating care coordination. However, we continue to lose patients at each step in the cascade of care. As calls escalate to simplify the cascade of care [22, 42], a streamlined cascade could simplify the process for patients and expand the pool of providers willing and able to treat patients for HCV. In this clinical cohort, the highest dropoff occurs at time of SVR measurement, resulting in patients who are unaware of treatment outcome and incomplete data at a state level relative to progress addressing the HCV epidemic. Despite laboratory cost coverage, physical distance and poor access to transportation limit patients' ability to return for laboratory testing. This drop-off is not unique to our study [8, 43, 44], suggesting a need for improved access to SVR12 testing. Ensuring Medicaid coverage for transport to a laboratory

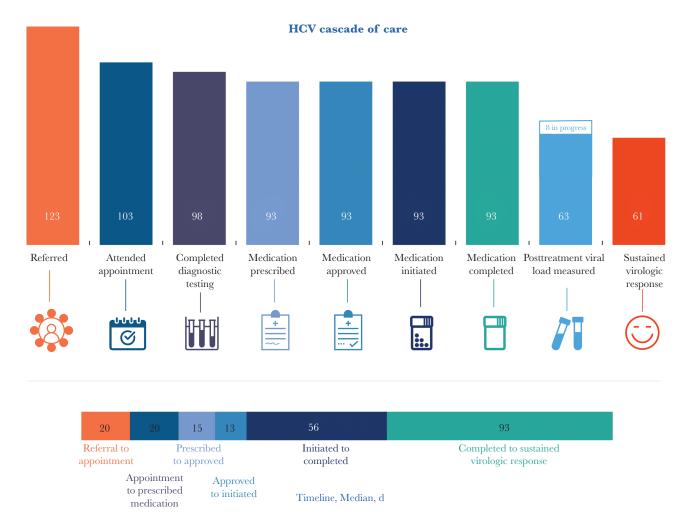


Figure 4. Hepatitis C virus (HCV) telehealth model cascade of care. *Top*, Cascade of care displays the numbers of patients (total N = 123) who completed each step required for treatment of HCV. *Bottom*, Timeline presents the median number of days to step completion. The median time from appointment to completion of diagnostic testing is combined with time from completion of diagnostic testing to prescription of medication, because some participants completed the diagnostic testing before their scheduled appointment.

appointment or coverage of mobile phlebotomy could address transportation barriers. Dried blood spot HCV testing may offer an option for shipping samples that could overcome transportation barriers [45].

The majority of our patients are enrolled in MAT programs. These patients had rates of progression along the cascade similar to those of patients not receiving MAT. This findings support recent studies demonstrating that patients receiving MAT have success in HCV treatment comparable to that of patients not receiving MAT [46]. Although we offer referrals to and receive referrals from a CHR program, patients who participated in CHR were not specifically tracked.

The presence of an on-site nurse navigator at a local health department led to a 50% reduction in time from referral to the initial telehealth visit. Although we saw rates of progression along the cascade similar to that of patients without a local nurse navigator, this early finding may suggest a benefit to local nurse navigators. Public health settings and local health departments are well situated to provide comprehensive patient care for people with HCV in high-burden rural areas. In addition to trained health department-based PCPs who can deliver on-site HCV treatment along with vaccination services, family planning, pre-exposure prophylaxis, and sexually transmitted infection screening, treatment, and prevention, there are public health nurses who can facilitate patient navigation. Prior qualitative studies have demonstrated that engaging PWID in HCV care requires more than a single visit [47], and a local nurse presence provides an opportunity to develop a therapeutic relationship.

Insurance prior authorizations have been a major barrier to HCV treatment in the DAA era, with frequent insurance denial of treatment requiring appeal [48], diversion of clinical staff time to insurance paperwork [49], and delays between time of prescription and approval [50]. Together, these logistical barriers to medication access place a significant burden on staff that may

Table 2. Progression Along the Cascade of Care Stratified by Receipt of Medication-Assisted Treatment, Presence of On-Site Nurse Navigator at Telehealth Location, and Medicaid Insurance

	Patient				
Patient Category	Attended Visit	Completed Medication	SVR	Time From Referral to Initial Visit, Median (IQR), d	
All patients (N = 123)	103 (84)	93 (76)	61 (50)	17.5 (5–33)	
MAT					
Yes (n = 76)	65 (86)	57 (75)	39 (51)	17.5 (6–33)	
No (n = 47)	38 (81)	36 (76)	22 (47)	17 (5–34)	
On-site nurse navigator					
Yes (n = 57)	49 (86)	46 (81)	26 (46)	11 (3–27.5) ^a	
No (n = 66)	54 (82)	47 (71)	35 (53)	22 (12–46) ^a	
Medicaid					
Yes (n = 78)	64 (83)	61 (79)	41 (53)	16 (5–31)	
No (n = 45)	39 (85)	32 (70)	20 (43)	20 (7–37)	

Abbreviations: IQR, interquartile range; MAT, medication assisted treatment; SVR, sustained virologic response.

 $^{a}P = .001$ for time from referral to initial visit based on presence of on-site nurse navigator. No differences were seen in time from referral to initial visit based on MAT or Medicaid status. No significant differences were seen in rates of completion of cascade steps based on receipt of MAT, on-site nurse navigator, or Medicaid status, using χ^2 tests to compare proportions.

limit clinics' ability to institute and scale up HCV treatment [49]. Historically, this burden has fallen most heavily on Medicaid patients [51] and may contribute to the observation that Medicaid patients have been less likely to link to care and receive treatment for HCV [39, 52]. In our model, progression along the cascade was similar for those on Medicaid compared with other types of insurance, probably owing to the full-time telehealth nurse coordinator who is expert at navigating the process of obtaining DAAs. Further reduction of HCV treatment restrictions for Medicaid patients [36] could alleviate the paperwork burden, streamline care, and enhance equity of access to HCV treatment. Reduced paperwork will allow skilled nursing or navigator time to be targeted to coordinating care with patients.

Improvements in access to comprehensive HCV treatment in high-burden areas require improved workforce capacity to deliver DAAs. However, as noted above, training medical providers is not sufficient for the development of local treatment programs. Support staff involvement is critical. Peers, people with prior experience with HCV treatment and/or substance use, may be an additional source of patient support, especially in areas where peer services are supported through state Medicaid programs [53]. However, at this time, results related to the success of peer support programs for HCV care are mixed [11, 31, 54].

We acknowledge limitations in the evaluation of our model. To truly eliminate HCV in the Southwest Virginia, efforts aimed at increasing testing would be needed. This model begins with patients whose HCV is already diagnosed. In addition, data reported are preliminary findings. We have presented all available data. However, many patients are continuing to progress along the cascade, and we anticipate ultimate cure rates to be higher than presented. Moreover, policy issues, including telehealth laws and reimbursement, Medicaid expansion, and the commitment of VDH to invest in this pilot program have led to the success of our model. Replication of this model elsewhere requires engagement with these policy issues on community and state levels.

A telehealth model can successfully provide HCV treatment and increase the workforce to move toward a goal of reducing disparities in HCV treatment access. This is a key opportunity for academic institutions in states with favorable policy environments.

Notes

Acknowledgments. First and foremost, we acknowledge our patients. We also thank Art Van Zee, Melissa Keene, Michelle Hilgart, Kimberly Rogers, Charles Lewis, Latasha Tolliver, Steve Murphy, Jessica Clark, Nicole Bissell, Cindy Foley, Devan Kaufman, as well as our partner organizations in this project: Virginia Department of Health Division of Disease Prevention, Lenowisco Health Department, the Stone Mountain federally qualified health center, and the University of Virginia Specialty Pharmacy.

Financial support. This work was supported by the National Institute of Allergy and Infectious Diseases, National Institutes of Health (grants T32 AI007046-43 and K08AI136644) and by Gilead (CHIME grant).

Supplement sponsorship. This supplement is sponsored by the Centers for Disease Control and Prevention.

Potential conflicts of interest. K. A. M. reports stock ownership in Gilead Sciences. R. D. provides consulting services to Warm Health Technologies, on activities unrelated to the current work. All other authors report no potential conflicts. All authors have submitted the ICMJE Form for Disclosure of Potential Conflicts of Interest. Conflicts that the editors consider relevant to the content of the manuscript have been disclosed.

References

1. 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:e186371.

- Zibbell JE, Iqbal K, Patel RC, et al. Increases in hepatitis C virus infection related to injection drug use among persons aged ≤30 years—Kentucky, Tennessee, Virginia, and West Virginia, 2006–2012. MMWR Morb Mortal Wkly Rep 2015; 64:453–8.
- 3. 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 **2018**; 108:175–81.
- Hachey DM, Holmes JT, Aubuchon-Endsley NL. Hepatitis C treatment cascade in a federally qualified health center. J Community Health 2020; 45:264–8.
- Schranz AJ, Barrett J, Hurt CB, Malvestutto C, Miller WC. Challenges facing a rural opioid epidemic: treatment and prevention of HIV and hepatitis C. Curr HIV/AIDS Rep 2018; 15:245–54.
- 6. Harris M, Rhodes T. Hepatitis C treatment access and uptake for people who inject drugs: a review mapping the role of social factors. Harm Reduct J **2013**; 10:7.
- Barua S, Greenwald R, Grebely J, Dore GJ, Swan T, Taylor LE. Restrictions for Medicaid reimbursement of sofosbuvir for the treatment of hepatitis C virus infection in the United States. Ann Intern Med 2015; 163:215–23.
- 8. Read P, Lothian R, Chronister K, et al. Delivering direct acting antiviral therapy for hepatitis C to highly marginalised and current drug injecting populations in a targeted primary health care setting. Int J Drug Policy **2017**; 47:209–15.
- 9. Falade-Nwulia O, Irvin R, Merkow A, et al. Barriers and facilitators of hepatitis C treatment uptake among people who inject drugs enrolled in opioid treatment programs in Baltimore. J Subst Abuse Treat **2019**; 100:45–51.
- Virginia Department of Health Office of Epidemiology. Virginia hepatitis C epidemiologic profile (2016).
 2016.
- 11. Høj SB, Jacka B, Minoyan N, Artenie AA, Bruneau J. Conceptualising access in the direct-acting antiviral era: an integrated framework to inform research and practice in HCV care for people who inject drugs. Int J Drug Policy 2019; 72:11–23.
- 12. Stephens DB, Young AM, Havens JR. Healthcare contact and treatment uptake following hepatitis C virus screening and counseling among rural Appalachian people who use drugs. Int J Drug Policy **2017**; 47:86–94.
- 13. American Association for the Study of Liver Diseases– Infectious Diseases Society of America. Recommendations for testing, managing, and treating hepatitis C. http://www. hcvguidelines.org. Accessed 30 September 2019.
- 14. National Academies of Sciences Engineering and Medicine. A national strategy for the elimination of hepatitis B and C:

phase two report. Washington DC: The National Academies Press; **2017**.

- Centers for Disease Control and Prevention. National viral hepatitis progress report. 2019. https://www.cdc.gov/hepatitis/policy/NationalProgressReport.htm?deliveryName=U SCDC_291-DM9332. Accessed 27 September 2019.
- Scott N, Doyle JS, Wilson DP, et al. Reaching hepatitis C virus elimination targets requires health system interventions to enhance the care cascade. Int J Drug Policy 2017; 47:107–16.
- Pitcher AB, Borquez A, Skaathun B, Martin NK. Mathematical modeling of hepatitis c virus (HCV) prevention among people who inject drugs: a review of the literature and insights for elimination strategies. J Theor Biol 2019; 481:194–201.
- Martin NK, Vickerman P, Grebely J, et al. Hepatitis C virus treatment for prevention among people who inject drugs: modeling treatment scale-up in the age of direct-acting antivirals. Hepatology **2013**; 58:1598–609.
- Platt L, Minozzi S, Reed J, et al. Needle syringe programmes and opioid substitution therapy for preventing hepatitis C transmission in people who inject drugs. Cochrane Database Syst Rev 2017; 9:CD012021.
- Tsui JI, Miller CM, Scott JD, Corcorran MA, Dombrowski JC, Glick SN. Hepatitis C continuum of care and utilization of healthcare and harm reduction services among persons who inject drugs in Seattle. Drug Alcohol Depend 2019; 195:114–20.
- 21. Asher AK, Portillo CJ, Cooper BA, Dawson-Rose C, Vlahov D, Page KA. Clinicians' views of hepatitis C virus treatment candidacy with direct-acting antiviral regimens for people who inject drugs. Subst Use Misuse **2016**; 51:1218–23.
- 22. Lazarus J V, Pericàs JM, Picchio C, et al. We know DAAs work, so now what? simplifying models of care to enhance the hepatitis C cascade. J Intern Med **2019**; 286:503–25.
- 23. Rosenthal ES, Silk R, Mathur P, et al. Concurrent initiation of hepatitis C and opioid use disorder treatment in people who inject drugs. Clin Infect Dis **2020**. doi:10.1093/cid/ ciaa105. Published 3 February 2020.
- 24. Guss D, Sherigar J, Rosen P, Mohanty SR. Diagnosis and management of hepatitis C infection in primary care settings. J Gen Intern Med **2018**; 33:551–7.
- 25. Kattakuzhy S, Gross C, Emmanuel B, et al. Expansion of treatment for hepatitis C virus infection by task shifting to community-based nonspecialist providers: a nonrandomized clinical trial. Ann Intern Med] **2017**; 167:311–8.
- 26. Wade AJ, Doyle JS, Gane E, et al. Outcomes of treatment for hepatitis C in primary care compared to hospital-based care: a randomised controlled trial in people who inject drugs. Clin Infect Dis **2020**; 70:1900–6.

- 27. Thomson M, Konerman MA, Choxi H, Lok ASF. Primary care physician perspectives on hepatitis C management in the era of direct-acting antiviral therapy. Dig Dis Sci **2016**; 61:3460–8.
- 28. Falade-Nwulia O, McAdams-Mahmoud A, Irvin R, et al. Primary care providers knowledge, attitude and practices related to hepatitis C screening and treatment in the oral direct acting antiviral agents era. J Community Med Health Educ **2016**; 6:481.
- 29. Naghdi R, Seto K, Klassen C, et al. A hepatitis C educational needs assessment of Canadian healthcare providers. Can J Gastroenterol Hepatol **2017**; 2017:5324290.
- Havens JR, Walsh SL, Korthuis PT, Fiellin DA. Implementing treatment of opioid-use disorder in rural settings: a focus on HIV and hepatitis C prevention and treatment. Curr HIV/AIDS Rep 2018; 15:315–23.
- 31. Litwin AH, Drolet M, Nwankwo C, et al. Perceived barriers related to testing, management and treatment of HCV infection among physicians prescribing opioid agonist therapy: the C-SCOPE Study. J Viral Hepat 2019; 26:1094–104.
- 32. Sherbuk JE, McManus KA, Rogawski McQuade ET, Knick T, Henry Z, Dillingham R. Hepatitis C within a single health system: progression along the cascade to cure is higher for those with substance misuse when linked to a clinic with embedded support services. Open Forum Infect Dis **2018**; 5:ofy202.
- 33. 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 **2020**; 27:13–19.
- Public Health Institute Center for Connected Health Policy. State telehealth laws & reimbursement policies. Center for Connected Health Policy, Public Health Institute; 2019.
- 35. Arora S, Thornton K, Murata G, et al. Outcomes of treatment for hepatitis C virus infection by primary care providers. N Engl J Med **2011**; 364:2199–207.
- 36. National Viral Hepatitis Roundtable and Center for Health Law and Policy Innovations at Harvard Law School. Hepatitis C: state of Medicaid access. 2019 https:// stateofhepc.org/. Accessed 27 September 2019.
- Sokol R, Early J, Barner A, et al. Implementation of a multidisciplinary, team-based model to treat chronic hepatitis C in the primary care setting: lessons learned. Healthcare 2018; 6:205–9.
- 38. Sacks-Davis R, Doyle JS, Rauch A, et al. Linkage and retention in HCV care for HIV-infected populations: early data from the DAA era. J Int AIDS Soc 2018; 21(suppl 2):e25051.
- 39. Zuckerman A, Douglas A, Nwosu S, Choi L, Chastain C. Increasing success and evolving barriers in the hepatitis C cascade of care during the direct acting antiviral era. PLoS One 2018; 13:e0199174.
- 40. Reau N, Marx S, Manthena S, Strezewski J, Chirikov V. National examination of HCV linkage to care in the United

States (2013–2016) based on large real-world dataset. Hepatology **2018**; 68:892A–3A.

- 41. Sherbuk JE, McManus KA, Kemp Knick T, Canan CE, Flickinger T, Dillingham R. Disparities in hepatitis C linkage to care in the direct acting antiviral era: findings from a referral clinic with an embedded nurse navigator model. Front Public Health 2019; 7:362.
- 42. Mulligan K, Sullivan J, Yoon L, Chou J, Nuys K Van. Evaluating HCV screening, linkage to care, and treatment across insurers. Am J Manag Care **2018**; 24:e257–64.
- 43. Read P, Gilliver R, Kearley J, et al. Treatment adherence and support for people who inject drugs taking direct-acting antiviral therapy for hepatitis C infection. J Viral Hepat **2019**; 26:1301–10.
- 44. Coyle C, Moorman AC, Bartholomew T, et al. The hepatitis C virus care continuum: linkage to hepatitis C virus care and treatment among patients at an urban health network, Philadelphia, PA. Hepatology **2019**; 70:476–86. 2019; 26:1301-10.
- 45. Vázquez-Morón S, Ardizone Jiménez B, Jiménez-Sousa MA, Bellón JM, Ryan P, Resino S. Evaluation of the diagnostic accuracy of laboratory-based screening for hepatitis C in dried blood spot samples: a systematic review and metaanalysis. Sci Rep **2019**; 9:7316.
- Latham NH, Doyle JS, Palmer AY, et al. Staying hepatitis C negative: a systematic review and meta-analysis of cure and reinfection in people who inject drugs. Liver Int **2019**; 39:2244–60.
- 47. Coupland H, White B, Bates A, Park JN, Iversen J, Maher L. Engaging people who inject drugs in hepatitis C virus testing and prevention through communitybased outreach, in Sydney, Australia. Drug Alcohol Rev 2019; 38:177-84.
- 48. Do A, Mittal Y, Liapakis A, et al. Drug authorization for sofosbuvir/ledipasvir (Harvoni) for chronic HCV infection in a real-world cohort: a new barrier in the HCV care cascade. PLoS One **2015**; 10:e0135645.
- 49. Millman AJ, Ntiri-Reid B, Irvin R, et al. Barriers to treatment access for chronic hepatitis C virus infection: a case series. Top Antivir Med **2017**; 25:110–3.
- 50. Ma J, Non L, Amornsawadwattana S, Olsen MA, Garavaglia Wilson A, Presti RM. Hepatitis C care cascade in HIV patients at an urban clinic in the early direct-acting antiviral era. Int J STD AIDS **2019**; 30:834–42.
- 51. Younossi ZM, Bacon BR, Dieterich DT, et al. Disparate access to treatment regimens in chronic hepatitis C patients: data from the TRIO network. J Viral Hepat 2016; 23:447–54.
- 52. Wong RJ, Jain MK, Therapondos G, et al. Race/ethnicity and insurance status disparities in access to direct acting antivirals for hepatitis C virus treatment. Am J Gastroenterol **2018**; 113:1329–38.

- 53. Virginia Department of Behavioral Health and Developmental Services. Peer services supplement, peer support services and family support partners. 2017. http:// www.dbhds.virginia.gov/assets/doc/recovery/providermanual-supplement.pdf. Accessed 10 March 2020.
- 54. Ward KM, Falade-Nwulia O, Moon J, et al. A randomized controlled trial of cash incentives or peer support to increase HCV treatment for persons with HIV who use drugs: the CHAMPS study. Open Forum Infect Dis **2019**; 6:ofz166.