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2018-04-25

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### **Repository Citation**

Martin SA, Bosse J, Wilson A, Losikoff P, Chiodo L. (2018). Under one roof: identification, evaluation, and treatment of chronic hepatitis C in addiction care. Open Access Publications by UMMS Authors. https://doi.org/10.1186/s13722-018-0111-7. Retrieved from https://escholarship.umassmed.edu/oapubs/3436



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### COMMENTARY

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# Under one roof: identification, evaluation, and treatment of chronic hepatitis C in addiction care

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#### Abstract

For over a decade, the vast majority of new hepatitis C virus (HCV) infections have been among young people who inject drugs (PWID). Well-characterized gaps in chronic HCV diagnosis, evaluation, and treatment have resulted in fewer than 5% of PWID receiving HCV treatment. While interferon-based treatment may have intentionally been foregone during part of this time in anticipation of improved oral therapies, the overall pattern points to deficiencies and treatment exclusions in the health care system. Treatment for HCV with all-oral, highly effective direct-acting antiviral medication for 12 weeks or less is now the standard of care, putting renewed focus on effective delivery of care. We describe here both the need for and process of chronic HCV care under the roof of addiction medicine.

**Keywords:** Opioid use disorder, Chronic hepatitis C, Cirrhosis, Hepatitis C epidemiology, Hepatitis C treatment, Treatment cascade, Project ECHO, Continuity of care

#### Background

Two profound changes in chronic hepatitis C virus (HCV) infection have occurred over the past decade. The first is the availability of medications that allow an all-oral highly effective cure. These direct-acting antiviral (DAA) agents have few adverse effects and enable treatment completion in as little as 8–12 weeks, with success rates exceeding 90% [1]. They are sorely needed given that HCV is now the leading cause of infectious death in the United States [2] and the most common cause of liver transplant [3].

The second change is the development of a bimodal chronic HCV demographic, with American Baby Boomers joined by young people who inject drugs (PWID). Among new acute HCV cases in 2014, more than two-thirds (68.2%) reported injection drug use [4]. HCV incidence had been in decline until early 2000s, a success enabled by identification of the virus in 1989. Subsequent blood product screening prior to transfusion, begun in

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1990, virtually eliminated this mode of HCV transmission by 1992. Needle exchange efforts and a decrease in injection drug use further helped HCV incidence to decline to below 1 per 100,000 by the early 2000s. Between 2010 and 2015 this trend reversed, with acute HCV infections rising nearly threefold—a figure likely underestimated by a factor of 15 [2, 5].

Patients who are not able to obtain HCV treatment risk transmitting the virus to others, which in turn, leads to higher incidence of disease and subsequent illness. Doubling the number of patients treated each year, however, could decrease HCV prevalence to fewer than 100,000 cases by 2030 [6]. This approach, similar to that pursued for HIV, has been termed treatment (or cure) as prevention (CasP) [7–9]. Given the impact of HCV on individuals who are infected and their families, the health care system, and larger society, minimizing barriers and maximizing access to more easily tolerated evidence-based treatment for younger persons who inject drugs is imperative. Yet PWID are often relatively isolated, even stigmatized, by standard health care [10].

The HCV treatment cascade (see Table 1) describes the clinical course, from initially evaluating HCV risk to finally achieving sustained virologic response (SVR).



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Cascade step	Outpatient addiction treatment site advantages	Cascade step eliminated
1. Patients are diagnosed and aware of their infection	Standardized screening of all patients and counseling with results	
2. Linkage to care (Access to outpatient care)	HCV care remains at the site	✓
3. Confirmatory testing for HCV RNA	Standardized reflex testing	✓
4. Liver disease evaluation, including liver biopsy (Note: Liver biopsy is not required for all patients)	Blood-based biomarker testing, a recommended alternative to liver biopsy, can be readily performed on site	1
5. Prescribed HCV treatment	Prescribed on site with coordination from specialty pharmacies	1
<ol> <li>Achieved a sustained virologic response (SVR), also referred to as being cured</li> </ol>	SVR testing done on site	1

Table 1 Mechanisms to address gaps in treatment cascade [16, 23]

There are substantial impediments at each stage [11, 12]. Two recent cohort studies examined this younger population—people with opioid use disorder (OUD) in medication-assisted treatment who have chronic HCV. Both found < 3% of patients initiated HCV treatment [13, 14].

Many barriers to treatment can be removed through treatment at a trusted place where HCV testing occurs. This would allow for immediate counseling and treatment evaluation after a positive result [15]. Given the high risk for HCV with the use of injection drugs, sites for outpatient treatment of OUD are a logical starting point to provide this range of HCV services. In 2017, this approach was strongly supported by the American Society of Addiction Medicine, which stated that "integration of service delivery, addressing the unique needs of addiction patients including HCV treatment, is strongly encouraged" [16]. HCV treatment in settings of OUD care is successful in Australia [17], Switzerland [18], and Italy [19]. Recently this success was repeated in Connecticut in a primary care setting embedded in an opioid treatment center [20].

## HCV and medication assisted treatment care integration

Over the past year, we have been learning from the pilot provision of HCV treatment as part of meeting our patients' health needs—independent of primary care and eliminating multiple steps of the usual HCV treatment cascade (Table 1). CleanSlate Addiction Treatment Centers offers dedicated treatment for opioid and alcohol use disorders in eight states, with clinicians that include social workers, advanced practice clinicians, and physicians. Patients in treatment for OUD at CleanSlate have a median age of 36 years old, with 26% under the age of 30 and 64% under the age of 40. As such, they represent the demographic profile of the younger person with OUD and an elevated risk of chronic HCV infection. As the treatment cascade makes clear, caring for a group of patients with a high burden of chronic HCV does not inevitably lead to its cure; the combination of an effective treatment strategy together with longitudinal care is especially helpful. The average length of treatment in our buprenorphine program is over one year (SD = 1.3, range 0.2-5.5). This timeframe is well beyond the 12 weeks of HCV treatment needed and allows providers more time with their patients to develop therapeutic relationships. Additional time spent with patients allows providers to assess patients' stability and motivation for treatment and provide education, which, in the context of the therapeutic relationship, can maximize adherence to treatment.

All patients who initiate substance use disorder treatment at CleanSlate are screened for blood-borne pathogens: Hepatitis B (HBV), Human Immunodeficiency Virus (HIV), and HCV testing that reflexes to a HCV viral count for all HCV antibody-positive specimens. By screening all patients we have found a 25–30% prevalence of positive HCV antibody; 75% of these patients with positive HCV antibody are subsequently found to have chronic HCV. All addiction providers are trained in HCV pre- and post-test counseling, including educating patients about DAA. Standard of care has been to refer all HCV infected patients to community providers for further evaluation and treatment [21].

In 2016, we began a pilot program in one of our clinics offering treatment for chronic HCV infection together with addiction treatment. Results of our pilot found that co-location of HCV treatment in our addiction clinic reduces the care cascade attrition found in PWID populations. In our pilot site sample, of the approximately 740 patients who were screened, 167 (22.5%) patients were found to have chronic HCV. All are in the process of being offered treatment. As of September 1, 2017, 72 (43%) entered DAA treatment. Among those who entered treatment, 72% received HCV treatment at CleanSlate and 28% chose to receive care via community treatment. Of the patients receiving care at our site

who have reached 12 weeks after treatment completion (N = 48), 4 have been lost to follow-up and 2 did not complete treatment. One patient had a treatment failure at 12 weeks post-treatment. Our on-site rate of cure is 85% (41 of 48) using intention-to-treat analysis and 98% (41 of 42) among those for whom we have SVR data. In a similar CleanSlate without co-located care site, only 10% entered DAA care in the community.

Three HCV treatment model strategies have been critical for success; each lends itself to scaling and sustainability. First, we have in-house expertise of an infectious disease specialist with interest in both addiction care and HCV. This expert mentors advanced practice clinicians to independently treat HCV and also directs clinical care for patients with different severities of cirrhosis and associated complications. This expertise can be remotely offered to other sites. Though many sites for OUD treatment do not have infectious disease specialists, our sense is that partnerships with them allow a novel, effective way to meet the needs of a high-risk population and, by extension, contribute to cure as prevention.

Second, just as our clinical experience with OUD registries lent itself to HCV treatment, we have repurposed staff expertise with prior authorizations, medication management, and treatment adherence. Laboratory staff have been especially helpful in facilitating both our screening and treatment protocols.

Third, we are using the model of a learning, capacitybuilding organization, with clinicians (advanced practice clinicians and physicians) taking part in an HCV Project ECHO and developing new skills in HCV care [22]. Over the course of several months of ECHO participation including presenting de-identified cases and listening to others' cases—we were able to develop advanced practice clinician skills to provide treatment independently. The ECHO has also been a place where we can share and learn about newer treatments and approaches in this rapidly-developing field.

#### Conclusion

People with chronic HCV currently face fragmented care provision, including diagnosis and cure of chronic HCV. Addiction treatment programs are uniquely situated to either use their existing program infrastructures or expand current programming to increase access to HCV care and treatment. By recognizing the unique, trusted, ongoing clinical role we play in many patients' lives, we can help their health beyond addiction.

#### Authors' contributions

Each author contributed to the conception, writing, and revising of this manuscript. All authors read and approved the final manuscript..

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#### Acknowledgements

Not applicable

#### **Competing interests**

We describe in this manuscript a model of care in place at CleanSlate Addiction Treatment Centers. It is not intrinsic to our organization, and we make the case that it can—and should—be implemented at many other sites of care for substance use disorders.

#### Availability of data and materials

Not applicable.

#### Consent for publication

Not applicable.

#### Ethics approval and consent to participate

Not applicable.

#### Funding

Funding for the development of this manuscript was provided by the CleanSlate Research and Education Foundation.

#### **Publisher's Note**

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

#### Received: 20 July 2017 Accepted: 14 February 2018 Published online: 25 April 2018

#### References

- American Association for the Study of Liver Diseases, Infectious Diseases Society of America. HCV guidance: recommendations for testing, managing, and treating hepatitis C. http://www.hcvguidelines.org/. Accessed 3 Jul 2017.
- Centers for Disease Control and Prevention (CDC). Hepatitis C kills more Americans than any other infectious disease. 2016. Available at: https://www.cdc.gov/media/releases/2016/p0504-hepc-mortality.html. Accessed 20 July 2017.
- Clark P, Muir A. Overcoming barriers to care for hepatitis C. N Engl J Med. 2012;366:2436–8.
- Division of Viral Hepatitis, Centers for Disease Control and Prevention (CDC). U.S. 2014 surveillance data for viral hepatitis. 2016. https://www.cdc.gov/hepatitis/statistics/2014surveillance/index. htm#tabs-1170600-11.
- Hepatitis Awareness Month and Testing Day–May 2017. MMWR Morb Mortal Wkly Rep. 2017;66:465.
- Razavi H, Elkhoury AC, Elbasha E, Estes C, Pasini K, Poynard T, et al. Chronic hepatitis C virus (HCV) disease burden and cost in the United States. Hepatology. 2013;57:2164–70.
- Stopka TJ, Goulart MA, Meyers DJ, Hutcheson M, Barton K, Onofrey S, et al. Identifying and characterizing hepatitis C virus hotspots in Massachusetts: a spatial epidemiological approach. BMC Infect Dis. 2017;17:294.
- Martin NK, Vickerman P, Grebely J, Hellard M, Hutchinson SJ, Lima VD, 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.

- Robaeys G, Grebely J, Mauss S, Bruggmann P, Moussalli J, De Gottardi A, et al. Recommendations for the management of hepatitis C virus infection among people who inject drugs. Clin Infect Dis. 2013;57(suppl 2):S129–37.
- Zeremski M, Zibbell JE, Martinez AD, Kritz S, Smith BD, Talal AH. Hepatitis C virus control among persons who inject drugs requires overcoming barriers to care. World J Gastroenterol. 2013;19:7846.
- Yehia BR, Schranz AJ, Umscheid CA, Lo Re V. The treatment cascade for chronic hepatitis C virus infection in the United States: a systematic review and meta-analysis. PLoS ONE. 2014;9:3–9.
- Holmberg SD, Spradling PR, Moorman AC, Denniston MM. Hepatitis C in the United States. N Engl J Med. 2013;368:1859–61.
- Carey KJ, Huang W, Linas BP, Tsui JI. Hepatitis C virus testing and treatment among persons receiving buprenorphine in an office-based program for opioid use disorders. J Subst Abuse Treat. 2016;66:54–9.
- Krans EE, Zickmund SL, Rustgi VK, Park SY, Dunn SL, Schwarz EB. Screening and evaluation of hepatitis C virus infection in pregnant women on opioid maintenance therapy: a retrospective cohort study. Subst Abuse. 2015;37(1):88–95.
- 15. Laraque F, Varma JK. A public health approach to hepatitis C in an urban setting. Am J Public Health. 2017;107:922–6.
- American Society of Addiction Medicine. Policy Statement: Hepatitis C Infection. 2017. Available at: https://www.asam.org/advocacy/ find-a-policy-statement/view-policy-statement/public-policy-statements/2017/04/11/hepatitis-c. Accessed 17 Feb 2018.

- Alavi M, Grebely J, Micallef M, Dunlop AJ, Balcomb AC, Day CA, et al. Assessment and treatment of hepatitis C virus infection among people who inject drugs in the opioid substitution setting: ETHOS study. Clin Infect Dis. 2013;57(2):S62–9.
- Schitz A, Moser S, Marchart K, Haltmayer H, Gschwantler M. Direct observed therapy of chronic hepatitis C with interferon-free all-oral regimens at a low-threshold drug treatment facility—a new concept for treatment of patients with borderline compliance receiving opioid substitution therapy. Am J Gastroenterol. 2016;111:903–5.
- 19. Grassi A, Ballardini G. Hepatitis C in injection drug users: it is time to treat. World J Gastroenterol. 2017;23:3569.
- Butner JL, Gupta N, Fabian C, Henry S, Shi JM, Tetrault JM. Onsite treatment of HCV infection with direct acting antivirals within an opioid treatment program. J Subst Abuse Treat. 2017;75:49–53.
- Losikoff P, Gomes L, Coonan B, Mendenhall A, Hewitt T, Kwapien T, et al. Poster: treatment of hepatitis C infection in an office-based opiate treatment (OBOT) clinic. American Association for the Study of Liver Diseases. 2017.
- 22. Arora S, Thornton K, Murata G, Deming P, Kalishman S, Dion D, et al. Outcomes of treatment for hepatitis C virus infection by primary care providers. N Engl J Med. 2011;364:2199–207.
- Division of Viral Hepatitis, Centers for Disease Control and Prevention. National viral hepatitis action plan for 2017–2020. 2017. https://www.cdc. gov/hepatitis/hhs-actionplan.htm. Accessed 15 June 2017.

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