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Title: Developing a primary care-initiated hepatitis C treatment pathway in Scotland: A qualitative study

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Abstract:

Background: The ease of contemporary hepatitis C virus (HCV) therapy has prompted a global drive towards simplified and decentralised treatment pathways. In some countries, primary care has become an integral component of community-based HCV treatment provision. However, in the UK, the role of primary care providers remains largely focused on testing and diagnosis alone.

Aim: To develop a primary care-initiated HCV treatment pathway for people who use drugs, and recommend theory-informed interventions to help embed that pathway into practice.

Design and Setting: A qualitative design informed by behaviour change theory. Semi-structured interviews were undertaken with key stakeholders (n=38) primarily from two large urban conurbations in Scotland.

Method: Analysis was three-stage. First, we outlined a broad pathway structure, and then specified sequential pathway steps. Second, thematic data were aligned to pathway steps, and significant barriers and enablers identified. Third, the theoretical domains framework and behaviour change wheel were employed to systematically develop ideas to enhance pathway implementation, which stakeholders then appraised.

Results: The proposed pathway structure spans broad, overarching challenges to primary-care initiated HCV treatment. The theory-informed recommendations align to influences on different behaviours at key pathway steps, and focus on relationship building, routinisation, education, combating stigmas, publicising the pathway, and treatment protocol development.

Conclusion: This study provides the first practicable pathway for primary care-initiated HCV treatment in Scotland, and the UK. It positions primary care providers as an integral part of community-based HCV treatment, providing workable solutions to ingrained barriers to care.

Keywords:

General Practice; Hepatitis C; Primary Health Care; Qualitative Research; Scotland;

Therapeutics

How this fits in:

Historically, GPs were rarely involved in the treatment of HCV, their role more commonly restricted to viral testing and diagnosis. Contemporary drug therapy for HCV has allowed reconsideration of this status quo, and offers potential for GPs to initiate HCV treatment in primary care. This study provides a way forward, detailing a practicable theory-informed pathway and recommendations for primary care-initiated HCV treatment in the UK.

Introduction:

An estimated 58 million people were living with the hepatitis C virus (HCV) in 2019¹, equating to approximately 0.7% of the global population. In the UK, recent estimates suggest 118,000 individuals are chronically infected with HCV, with injecting drug use the most important risk factor for acquisition². While the number of people accessing HCV treatment is rising, the rate of increase has slowed in recent years, suggesting it is becoming harder to find, diagnose and treat people living with HCV as the pool of infection gradually decreases².

Historically, HCV treatment presented an arduous and challenging ordeal. Suboptimal toxic medications offered limited success and necessitated intensive monitoring and support³. As such, these drugs were the sole domain of specialist practitioners in secondary and centralised care. Over the last decade a sea change has occurred, with a barrage of novel drugs coming to market that target specific steps within the HCV life cycle. Collectively, these Direct Acting Antivirals (DAAs) now offer a safe, simple and effective cure with just 8-12 weeks of treatment⁴. The relative ease of contemporary HCV drug therapy has spurred a global drive towards decentralisation, expanding access by relocating the nexus of care firmly within the community. The World Health Organisation (WHO) endorses this move, calling for simplified and streamlined HCV treatment pathways to be integrated into existing healthcare systems, producing a plurality of community-based provision⁵.

Globally, this push for decentralised care has encouraged a raft of HCV treatment pathways in diverse settings, including harm reduction services, prisons, community pharmacies, homeless facilities as well as within general practice. Such pathways have repeatedly demonstrated increased uptake and comparable cure rates to treatment initiated in more traditional, hospital-based settings, illustrating their viability and acceptability as loci of care⁶.

⁷. While many community-based treatment pathways continue to rely on specialist

practitioners to initiate therapy (in an outreach model of provision), some have prioritised partial or comprehensive task-shifting to non-specialists, with no reduction in treatment efficacy⁸. The importance of some degree of task-shifting is underpinned by arguments for enabling access in areas of inadequate healthcare infrastructure⁹, and concerns that numbers of existing specialists are limited¹⁰. Task-shifting is also supported by recent challenges to 'medical speciality protectionism' voiced by the European Association for the Study of the Liver¹¹.

In the UK, while primary care has witnessed effective interventions to increase HCV testing and diagnosis¹², the role of most general practitioners (GPs) in HCV treatment remains limited to specialist referral, or hosting specialist-led outreach clinics. Linkage from diagnosis to treatment is a known bottleneck in the HCV care cascade¹³, and while other countries have helped alleviate barriers to HCV treatment by more comprehensive integration of GPs into models of DAA provision¹⁴, the UK is yet to follow suit. We have previously identified overarching challenges to primary care provision of HCV treatment in Scotland¹⁵, and here we move this work forward by articulating what a practicable, acceptable and sustainable model of primary care-initiated HCV treatment could look like.

We aim to develop a primary care-initiated HCV treatment pathway for the UK, and recommend theory-informed intervention elements to embed that pathway into practice.

Methods

The research team united academics and clinicians, offering a blend of expertise and perspectives in HCV, general practice, behaviour change theory and research methodology.

As principal investigator, DW is an early career researcher and former HCV nurse specialist, supported by LE, professor of public health and PF, professor of psychology. HJ, KD and MQ are all experienced clinicians, HJ and MQ working in general practice and KD in secondary care. ES is an experienced research fellow.

Theoretical Framework

Increasing evidence suggests public health interventions grounded in theory may be more effective than those which lack a theoretical basis¹⁶. We therefore conceptualised the implementation of primary-care initiated HCV treatment as requiring the development of a complex multi-actor/agent behaviour change intervention. We ask what behaviours need to change, who and where are they being performed, and how do these behaviours interact?¹⁷.

To avoid constraints imposed by any single behavioural theory, we employed the Theoretical

Domains Framework (TDF) to explore cognitive, affective, social and environmental influences on behaviour. The TDF is an integrated meta-theoretical framework which combines 33 theories of behaviour or behaviour change into 14 domains and 84 theoretical constructs¹⁸. It aims to make identification of the determinants of implementation behaviours more comprehensible, and is now widely used in intervention development studies, particularly within primary healthcare in developed nations¹⁹. The TDF also has established links with the Behaviour Change Wheel (BCW), a framework for classifying the *types* of intervention required to effectively alter specific behaviours²⁰. Both the TDF and the BCW are employed within this study.

Participants and Setting

The concept of primary-care initiated HCV treatment has implications for diverse stakeholders, and a purposive maximum variation sample was therefore used. This comprised GPs currently providing care for people who use drugs; people who use, or have used drugs, and are living with HCV (hereafter referred to as people living with HCV [PLHCV]); HCV specialists (doctors, nurses and pharmacists) working in hepatology and/or infectious diseases; staff from agencies providing support for PLHCV; and representatives from community pharmacies and NHS procurement. The study was primarily located within

two Scottish health boards with the highest number of new HCV diagnoses in 2018²¹, although recruitment of GPs widened to six additional Scottish health boards following study inception. The impact of the COVID-19 pandemic meant data collection was suspended in March 2020, with the study subsequently suspended in its entirety from May until August 2020.

Recruitment

GPs were recruited in two ways. First, an advertisement was placed in a monthly newsletter sent to all GP practices by the NHS Research Scotland Primary Care Network, inviting GPs to contact the study team. Second, GP practices within areas of social deprivation in two NHS boards were identified and invited to participate using publicly available email addresses. HCV specialists and community pharmacists were recruited through the existing networks of DW and KD, who purposively identified key individuals. Support agencies and NHS procurement were also approached directly, and the most appropriate potential participants within these organisations collaboratively identified. Interested respondents from all groups were given further information about the study, and time to consider their involvement prior to interviews being arranged at a time and place convenient for the interviewee.

Recruitment of PLHCV was led by the Scottish Drugs Forum (SDF), who work with and support a wide network of people who use drugs, many of whom are living with HCV. Staff from SDF approached PLHCV, who, if interested, were provided with further details of the study and given time to consider participation. Other avenues of recruitment for PLHCV were abandoned early in the study, and prior to any participants being recruited, to prevent additional pressure on NHS services during the COVID-19 pandemic.

Data Generation

Data were generated through semi-structured interviews. Two phases of interviews were compelled by the study suspension, with the first phase between October 2019 and March 2020, and the second phase between October 2020 and January 2021. Interviews with care providers were conducted by ES, predominantly in person during phase one, and using teleconferencing software in phase two. Interviews with PLHCV took place exclusively during phase two and were conducted remotely by SDF peer researchers, all of whom had lived experience of drug use and HCV. These interviews were facilitated by a member of SDF staff during a three-way conference call. Topic guides, specific to each participant group, were used to focus conversation. The topic guide for PLHCV was revised in

consultation with SDF peer researchers prior to use. In addition, examples of published primary care-initiated HCV treatment pathways were used as stimulus material in phase one interviews²²⁻²³, with the developing pathway also presented for discussion during interviews in phase two (i.e. key steps of the pathway helped to focus data generation). All interviews were audio-recorded, transcribed verbatim and pseudonymised. Finally, participants were asked if they would be willing to participate in a further focus group at a later date.

Data Analysis: Pathway Development

Initially, data from first phase interviews with care providers were thematically analysed to identify fundamental components of a primary care-initiated HCV treatment pathway, and explore broad, overarching challenges to its success. This stage of analysis has been described in detail elsewhere, and findings previously reported¹⁵. For clarity, figure 1 illustrates where this preliminary work sits within the process of pathway and recommendation development described here. An initial, simple pathway structure grew from this early analysis, which was iteratively revised, refined and expanded by DW during phase two interviews in light of emerging insights from the data. During this time, DW, ES, LE and PF met regularly to interrogate the developing pathway and specify the component parts as a series of behavioural steps, detailing *who* was enacting *what* behaviour at each

individual point. Individual steps were further specified as 'essential' or 'potential', and the evolving pathway regularly examined and critiqued by the wider study team, challenging its logic and lucidity. Eleven steps within the pathway were then collaboratively identified, that either significantly diverged from routine clinical practice, or were deemed 'hotspots' critical to success. In this way, key elements of a potential pathway were iteratively developed.

This stage ended with a clear set of sequential behavioural steps that together provided the basic elements of a potential primary care pathway for HCV treatment (figure 2).

Subsequent analysis fleshed out how these steps could be operationalised.

Data Analysis: Recommendation Development

The behavioural steps specified within the pathway comprised focal points for recommendation development to make the pathway work and be fit for purpose. Using a coding framework, ES and DW deductively coded relevant data to each of the 11 steps, identified key barriers and facilitators at each point, and ranked them reflecting their prominence within the data. *NVivo v. 12* software was used to manage this process. These barriers and facilitators were then mapped to domains of the TDF, to categorise influences on the stated behaviour. Next, established links between the TDF and BCW were utilised to identify precisely *how* these influences could be changed. Each domain of the TDF aligns to

specific intervention functions (i.e. *how* interventions might change behaviours) which link to a taxonomy of behaviour change techniques¹⁷. For example, if the data suggested a knowledge deficit represented a key barrier at a particular pathway step, this would be coded to the theoretical domain '*knowledge*' using the TDF, the BCW would then suggest the intervention function '*education*' which could be operationalised with the behaviour change technique '*providing instructions on how to perform a behaviour*'. This process was regularly audited by PF, an HCPC Registered Health Psychologist, who interrogated the initial TDF mapping, and provided robust review of the alignment to intervention functions and classified behaviour change techniques. From here, through robust and repeated discussions, the research team collaboratively developed detailed recommendations that specified how the pathway could be implemented. This stage ended with a rich set of theoretically informed and evidence-based recommendations for how to operationalise the proposed pathway. Subsequent work made sure these were pragmatic and likely to be sustainable.

Data Analysis: Stakeholder Review

Three focus groups were convened in May 2021 to provide expert review of both pathway and recommendations with a particular focus on if they could be delivered and were sustainable within current provision. Participants who had previously expressed interest

were approached, and a self-selecting sample asked to participate in one of three, two-hour, online-focus groups, facilitated by DW and ES using teleconferencing software. The APEASE criteria (Affordability, Practicability, Effectiveness, Acceptability, Side-effects and Equity)¹⁷ were used to guide discussion, with participants asked to reject, modify or accept the proposed recommendations. Focus groups were audio-recorded for reference, and any subsequent modifications made to the pathway and/or recommendations reviewed by the study team.

Results

Thirty-eight participants were interviewed. They comprised 11 GPs, 10 PLHCV; nine HCV specialists (four consultants, three specialist nurses and two specialist pharmacists); five staff from agencies providing support for PLHCV; two individuals employed by NHS procurement with insight into DAA reimbursement; and one community pharmacist. Twelve participants subsequently took part in one of the three focus groups, which comprised six GPs, two PLHCV, two HCV specialists (one consultant and one specialist pharmacist) and two staff from HCV support agencies.

Pathway

Figure 2 shows the final pathway developed, with 20 individual behavioural steps detailing 'who' needs to do 'what' within each sequential element. It expands on the basic building blocks of DAA treatment, but maintains three essential elements: the provision of drug (steps 6, 6a, 6b and 7), the assessment of liver fibrosis (steps 6 and 9) and opportunities for the provision of holistic support (steps 5, 5a, 5b, 6c and 8)¹⁵. While GPs were the initial focus of this study, the more generic term Primary Care Provider (PCP) has been purposefully used within the pathway, recognising that GPs are not the only practitioners located in GP surgeries who could initiate HCV treatment in primary care. Figure 2 also distinguishes steps characterised as 'essential' or 'potential' and shows which were designated 'hotspots' in particular need of operationalisation. These eleven hotspots incorporated both essential and potential steps, and covered the patient journey from attending an initial appointment with a PCP, to a post-treatment test of cure and liver fibrosis assessment. Steps 3a and 4b both focused on the utilisation of dried blood spot testing and/or venepuncture by PCPs, and were therefore combined within the analysis.

Detailed recommendations for pathway implementation

While pathway structure could help alleviate broad, overarching challenges previously identified¹⁵, the behavioural specification of the pathway allowed specific barriers and

facilitators to individual actions at each hotspot to be addressed. Supplementary box 1 provides an overview of 21 key barriers and facilitators identified across the eleven pathway hotspots, alongside illustrative quotes. The identified barriers and facilitators related to multiple behaviours by different actors across the pathway stages. An overview of the final recommendations is provided in supplementary box 2, detailed by each pathway step, following the TDF and BCW analysis and subsequent stakeholder appraisal. Of the 82 interventions originally suggested, stakeholders rejected 25, and modified eight of the remaining 57.

While diverse, the suggested recommendations could be categorised into one of six broad areas of intervention. First, interventions designed to foster and build professional and therapeutic relationships. For example, prioritising continuity of carer for PLHCV; developing and utilising GP clusters and managed care networks; and creating connections between GP surgeries and third sector partners. Second, interventions to help routinise and habituate the pathway into PCP's everyday practice. Included here were prompts within existing IT systems, and the establishment of clear and simple communication channels between PCPs and HCV specialist services. The third area of intervention was education. Specifically, training for PCPs on the pathway, HCV care, and the benefits of dried blood

spot testing, alongside optimising opportunities to educate PLHCV about the advantages of pathway engagement. Here, informal education was key, with important roles for the third sector, peer support workers and community link workers engaging in recurrent ad hoc conversations with PLHCV. Fourth, interventions to reduce the impact of social stigmas on pathway engagement, including offering HCV treatment within routine clinics, rather than identifiable 'HCV clinics' running on specific days and times. Fifth, a focus on interventions to publicise the existence of the pathway to PLHCV, promoted by third sector partners and other health and social care professionals embedded within the community. Finally, interventions to protocolise the pathway into local contexts, including plans for audit and review. For the interested reader, we provide full details of our analyses that underpin the findings in supplementary box 3.

Discussion

Summary

The decentralisation of HCV treatment into the community is a fundamental component of global and national HCV elimination plans^{2,5}. In some countries, GPs have been an integral cog in the decentralisation wheel for a number of years¹⁴, but in the UK, optimising the GP role has received little attention to date. This study is the first to provide a tangible way

forward, offering a detailed pathway for HCV treatment initiation by GPs and other PCPs in the UK. The pathway tackles previously identified challenges to primary-care initiated HCV treatment¹⁵, while offering theory-informed recommendations to help overcome specific behavioural barriers, and embed the pathway into practice.

Strengths and Limitations

The study has a number of strengths. Participants were drawn from diverse stakeholders, offering contrasting perspectives and insights from both service providers and service users. Also, while the study suspension due to COVID-19 was unplanned, it enhanced the iterative nature of pathway development by offering time for in-depth engagement with phase one interviews. Finally, effective public health initiatives are grounded in an understanding of health behaviours and the contexts in which they occur¹⁶. By defining the pathway and recommendations in behavioural terms, and utilising established tools that translate behavioural theory into actionable recommendations (including wide stakeholder engagement), this study bridges the divide between theoretical and applied research. There are also a number of limitations. Our participants were a self-selecting sample, who may represent a particularly motivated group of individuals. The COVID-19 pandemic and associated lockdowns arrived in Scotland six months after the start of recruitment. This

impacted recruitment and data generation options, particularly for PLHCV, who were entirely drawn from people engaging with SDF services, and who had the means to participate in a remote interview. Finally, we are conscious that the pathway and recommendations also offer a defined medical endpoint, largely ignoring the post-cure lives of PLHCV²⁴.

Comparison with Existing Literature

The WHO have called for the elimination of viral hepatitis as a major threat to global public health by 2030²⁵. Such an ambitious goal requires substantial increases in the testing and diagnosis of HCV, and innovative reassessment of where, and how, treatment can be provided²⁶. The arrival of DAAs has encouraged creative revision of HCV models of care, aimed at broadening access and simplifying the patient journey by decentralising care into the community, integrating with existing services, and task-shifting to less specialised health-care workers⁹. Of paramount importance, these simplified models of care *work*⁶⁻⁸.

Previous studies have focused on identifying barriers and enablers to GP involvement in decentralised HCV treatment²⁷⁻³¹, but often abstain from identifying how such barriers can be addressed. This study offers a practical way forward. Ingrained barriers such as the pre-treatment assessment of liver fibrosis are removed, supported by a growing evidence base

questioning its necessity³². The frequently cited difficulties of venepuncture are sidestepped by the use of sensitive, specific and accurate dried blood spot testing technologies³³. More problematic is the ability of PCPs to prescribe DAAs. While the WHO recognises much good practice in Scotland coordinating primary and secondary care prescribing, and in medicine reimbursement policies³⁴, current constraints in relation to DAAs prevent PCPs prescribing the drugs. A solution to the convoluted process of DAA reimbursement and prescribing process would allow further simplification of the pathway we propose here.

Scotland aims to exceed WHO targets and achieve HCV elimination by 2024 at the latest³⁵. To date, the primary focus of Scottish decentralised HCV treatment has been the co-location of outreach clinics within existing drug services, with limited task-shifting away from specialist clinicians³⁶. As self-imposed deadlines for HCV elimination loom, general practice and primary care remain peripheral to, or entirely absent from, the vision for community-based HCV treatment in Scotland³⁵. This is echoed throughout the UK, with the latest reports on progress towards HCV elimination from England and Wales locating the GP role clearly within HCV testing and diagnosis alone^{2,37}. The potential significance of omitting the contribution GPs could make to DAA prescribing is illustrated by comparison with other developed nations, notably Australia³⁸.

Globally, Australia is a bellwether for HCV elimination, leading the charge in the decentralisation of HCV treatment from hospital to primary care. Most people now treated for HCV in Australia are prescribed DAAs by non-specialists in community settings³⁸. This fundamental change in disease management was catalysed by the removal of prescribing restrictions for DAAs in March 2016, enabling GPs to evolve their roles from test and refer, to test and treat. This move broadened access to DAAs and helped facilitate a rapid increase in HCV treatment uptake over the first 10 months³⁹. More recent data demonstrates that initial momentum has been sustained, with the number of GPs prescribing HCV treatment continuing to increase¹⁴. While not without its challenges, the inclusion of GPs within models of decentralised HCV treatment expands access to people who use (or crucially *used*) drugs and are living with HCV but who do *not* engage with drug and alcohol services³⁰. While transplanting an Australian model of care into the UK may be tempting, doing so would ignore contextual factors that may limit its success, not least the complex system of drug reimbursement currently in operation for DAAs. A bespoke pathway, sensitive to local contexts, is required.

GP locality is also an important consideration. Scotland is a geographically diverse nation, with 30% of the population living in 'rural' areas⁴⁰. The current focus of HCV treatment provision within drug services ignores the geographic inequity of remote and rural populations experiencing poorer access to health services than their urban counterparts⁴⁰.

Recent drug death figures emphasise that people who use drugs in Scotland are not confined to urban settings⁴¹, and historic data reporting HCV antibody prevalence suggests neither is HCV⁴². For rural and remote communities, the role of the GP can differ from their urban counterparts, providing additional services and offering lifelong care to their communities from 'cradle to grave'^{40,43}. Within this context, a pathway for primary-care initiated HCV treatment may provide the only feasible access to care for some PLHCV.

While we advocate for further inclusion of GPs within the expanding web of community-based HCV treatment provision, we do so with a note of caution. This study sits within an ongoing recruitment crisis in general practice, which has only been exacerbated by the continuing COVID-19 pandemic⁴⁴⁻⁴⁵. While it may be feasible and realisable for GPs to initiate HCV treatment with this pathway, the current strains on primary care provision may preclude its enthusiastic embrace.

Implications for research and/or practice

Current HCV elimination policy in Scotland and the UK restricts the role of GPs to testing, diagnosis and referral. This study provides the first practicable pathway for primary care-initiated HCV treatment in the UK, offering GPs and other PCPs an opportunity to remove barriers to care and improve patient outcomes. Underpinned by theory, the pathway and associated recommendations offer a robust and realisable way to integrate primary care into the growing network of decentralised, community-based HCV treatment hubs.

We identified 57 recommendations to aid pathway implementation within six broad areas of intervention covering 11 pathway steps. These recommendations should not be taken as inflexible dogma, but as a toolkit to help adapt the pathway to different local contexts. For example, not all PCPs will need HCV training, and some GP practices will already have robust relationships with third sector partners and local HCV specialist teams. Our recommendations highlight focal points within the pathway that need to be considered, rather than the prescription of a one-size-fits-all solution. Future research should focus on the operationality and sustainability of the pathway through piloting.

Tables and figures

Figure 1: Sequential depiction of methodology.

Figure 2: The pathway

Supplementary data

Box S1: Overview of barriers and facilitators

Box S2: Summary of recommendations

Box S3: Identification of key barriers and facilitators by pathway step, detailing use of TDF and BCW

Additional information

Funding: This study was funded in full by the Chief Scientist Office of the Scottish Government Health Directorates (HIPS/18/49).

Ethical approval: A favourable ethical opinion was given by the West of Scotland Research Ethics Committee 3 (19/WS/0068). A written record of informed consent was obtained prior to in-person interviews with all participants, and remote interviews with care providers.

Verbal consent was recorded for interviews with PLHCV. GPs were compensated £80 per hour for their time, in line with the service support cost rate agreed by the study funder.

PLHCV received a £20 shopping voucher for their participation in interviews and focus groups. Other participants received no financial recompense.

Competing interests: DW has received speaker's honoraria from Gilead and Janssen and funding for attendance at conferences and educational meetings from Janssen, MSD and BMS. KD has received speaker's honoraria from Abbvie, Gilead, Janssen and MSD and funding for conferences and educational meeting attendance from Abbvie, BMS, Gilead, Janssen and MSD. The remaining authors have nothing to declare.

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References

1. World Health Organisation. *Global progress report on HIV, viral hepatitis and sexually transmitted infections, 2021. Accountability for the global health sector strategies 2016-*

2021: actions for impact. Available from:

<https://www.who.int/publications/i/item/9789240027077> [Accessed 09 Dec 2021]

2. Public Health England. *Hepatitis C in the UK 2020. Working to eliminate hepatitis C as a major public health threat*. Available from:

https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/943154/HCV_in_the_UK_2020.pdf [Accessed 09 Dec 2021]

3. Whiteley D, Elliott L, Cunningham-Burley S, Whittaker A. Health-related quality of life for individuals with hepatitis C: A narrative review. *Int J Drug Policy*. 2015 Oct;26(10):936-949. doi.org/10.1016/j.drugpo.2015.04.019.

4. Zoratti MJ, Siddiqua A, Morassut RE et al. Pangenotypic direct acting antivirals for the treatment of chronic hepatitis C virus infection: A systematic literature review and meta-analysis. *EClinicalMedicine*. 2020 Jan;18:100237. doi.org/10.1016/j.eclinm.2019.12.007.

5. World Health Organisation. *Guidelines for the care and treatment of persons diagnosed with chronic hepatitis C virus infection. July 2018*. Available from:

<https://apps.who.int/iris/bitstream/handle/10665/273174/9789241550345-eng.pdf?ua=1>

[Accessed 09 Dec 2021]

6. Oru E, Trickey A, Shirali R et al. Decentralisation, integration, and task-shifting in hepatitis C virus infection testing and treatment: a global systematic review and meta-analysis. *Lancet Glob Health*. 2021 Apr;9(4): e431-e445. doi.org/10.1016/S2214-109X(20)30505-2

7. Radley A, Robinson E, Aspinall EJ et al. A systematic review and meta-analysis of community and primary-care-based hepatitis C testing and treatment services that employ direct acting antiviral drug treatments. *BMC Health Serv Res*. 2019

Oct;19(1):765. doi.org/10.1186/s12913-019-4635-7

8. Castro R, Perazzo H, de Araujo LAMM et al. Effectiveness of implementing a decentralized delivery of hepatitis C virus treatment with direct-acting antivirals: A systematic review with meta-analysis. *PLoS One*. 2020 Feb;15(2):e0229143. doi.org/10.1371/journal.pone.0229143
9. Cooke GS. Decentralisation, integration, and task shifting: tools to accelerate the elimination of hepatitis C. *Lancet Glob Health*. 2021 Feb;9(4):E375-E376. doi.org/10.1016/S2214-109X(21)00055-3
10. Fox R. Going viral: Why eliminating the burden of hepatitis C requires enhanced cooperation between specialists and primary care providers. *Dig Dis Sci*. 2016 Dec;61(12):3381-3383. doi.org/10.1007/s10620-016-4301-4
11. Karlsen TH, Sheron N, Zelber-Sagi S et al. The EASL-Lancet liver commission: protecting the next generation of Europeans against liver disease complications and premature mortality. *Lancet*. 2021 Dec;S0140-6736(21)01701-3. doi.org/10.1016/S0140-6736(21)01701-3
12. Roberts K, Macleod J, Metcalfe C et al. Cost effectiveness of an intervention to increase uptake of hepatitis C virus testing and treatment (HepCATT): cluster randomised controlled trial in primary care. *BMJ*. 2020 Feb;368:m322. doi.org/10.1136/bmj.m322.
13. Jordan AE, Perlman DC, Reed J et al. Patterns and gaps identified in a systematic review of the hepatitis C virus care continuum in studies among people who use drugs. *Front Public Health*. 2017 Dec;5:348. doi.org/10.3389/fpubh.2017.00348.
14. Stafford F, Dore GJ, Clackett S et al. Prescribing of direct-acting antiviral therapy by general practitioners for people with hepatitis C in an unrestricted treatment program. *Med J Aust*. 2021 Oct;215(7):332-333. doi.org/10.5694/mja2.51204

15. Whiteley D, Speakman E, Elliott L et al. Provider-related barriers and enablers to the provision of hepatitis C treatment by general practitioners in Scotland: A behaviour change analysis. *J Viral Hepat.* 2021 Mar;28(3):528-537. doi.org/10.1111/jvh.13443.
16. Glanz K, Bishop DB. The role of behavioural science theory in development and implementation of public health interventions. *Annu Rev Public Health.* 2010;31:399-418. doi.org/10.1146/annurev.publhealth.012809.103604
17. Michie S, Atkins L, West R. *The Behaviour Change Wheel: A Guide to Designing Interventions.* Bream, Gloucester: Silverback Publishing; 2014.
18. Atkins L, Francis J, Islam R et al. A guide to using the Theoretical Domains Framework of behaviour change to investigate implementation problems. *Implement Sci.* 2017 Jun;12(1):77. doi.org/10.1186/s13012-017-0605-9.
19. Dyson J, Cowdell F. How is the Theoretical Domains Framework applied in designing interventions to support healthcare practitioner behaviour change? A systematic review. *Int J Qual Health Care.* 2021 July;33(3):mzab106. doi.org/10.1093/intqhc/mzab106
20. Michie S, van Stralen MM, West R. The behaviour change wheel: a new method for characterising and designing behaviour change interventions. *Implement Sci.* 2011 Apr;6:42. doi.org/10.1186/1748-5908-6-42.
21. Health Protection Scotland. *Surveillance report: Surveillance of hepatitis C testing, diagnosis and treatment in Scotland, 2019 update.* Available from: https://hpspubsrepo.blob.core.windows.net/hps-website/nss/2834/documents/1_hcv-testing-diagnosis-treatment-scotland-2018.pdf [Accessed 09 Dec 2021]
22. 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 nonrandomised clinical trial. *Ann Intern Med.* 2017 Sep;167(5):311-318. doi.org/10.7326/M17-0118.

23. Wade AJ, McCormack A, Roder C et al. Aiming for elimination: Outcomes of a consultation pathway supporting regional general practitioners to prescribe direct-acting antiviral therapy for hepatitis C. *J Viral Hepat.* 2018 Sep;25(9):1089-1098. doi.org/10.1111/jvh.12910
24. Seear K, Fraser S, Farrugia A, Valentine K. Beyond a 'post-cure' world: Sketches for a new futurology of hepatitis C. *Int J Drug Policy.* 2021 Aug;94:103042. doi.org/10.1016/j.drugpo.2020.103042.
25. World Health Organisation. *Global Health Sector Strategy on Viral Hepatitis 2016-2021: Towards ending viral hepatitis.* Available from: <https://apps.who.int/iris/bitstream/handle/10665/246177/WHO-HIV-2016.06-eng.pdf> [Accessed 09 Dec 2021]
26. Heffernan A, Cooke GS, Nayagam S et al. Scaling up prevention and treatment towards the elimination of hepatitis C: a global mathematical model. *Lancet.* 2019 Mar;393(10178):1319-1329. doi.org/10.1016/S0140-6736(18)32277-3.
27. Doshi RK, Ruben M, Drezner K et al. Knowledge, attitudes, and behaviours related to hepatitis C screening and treatment among health care providers in Washington, DC. *J Community Health.* 2020 Aug;45(4):785-794. doi.org/10.1007/s10900-020-00794-z.
28. Heard E, Massi L, Smirnov A, Selvey LA. Prescribing direct-acting antivirals to treat hepatitis C virus in a general practice setting in Australia: 'so why not do it'? *Intern Med J.* 2020 Sep;50(9):1053-1058. doi.org/10.1111/imj.14648.
29. Johnson S, Aluzaita K, Taar A, Schultz M. Identifying barriers to treatment of HCV in the primary care setting. *Hepatol Int.* 2019 Jan;13(1):58-65. doi.org/10.1007/s12072-018-9902-x.

30. Marshall AD, Grebely J, Dore GJ, Treloar C. Barriers and facilitators to engaging in hepatitis C management and DAA therapy among general practitioners and drug and alcohol specialists – The practitioner experience. *Drug Alcohol Depend.* 2020 Jan;206:107705. doi.org/10.1016/j.drugalcdep.2019.107705.
31. Pourmarzi D, Smirnov A, Hall L et al. Enablers and barriers for the provision of community-based HCV treatment: A case study of a real-world practice. *J Viral Hepat.* 2020 May;27(5):484-496. doi.org/10.1111/jvh.13259
32. Brennan PN, Cartlidge P, Robinson E et al. Cohort study: Apparent redundancy of fibrosis assessment in young persons with HCV; development of realistic approaches to break the paradigm. *Ann Hepatol.* 2022 Jan-Feb;27(1):100550. doi.org/10.1016/j.aohep.2021.100550
33. Vázquez-Morón S, Ardizzone Jiménez B, Jiménez-Sousa MA et al. Evaluation of the diagnostic accuracy of laboratory-based screening for hepatitis C in dried blood spot samples: A systematic review and meta-analysis. *Sci Rep.* 2019 May;9(1):7316. doi.org/10.1038/s41598-019-41139-8
34. World Health Organisation. *Medicines Reimbursement Policies in Europe.* Available from: https://www.euro.who.int/_data/assets/pdf_file/0011/376625/pharmaceutical-reimbursement-eng.pdf [Accessed 09 Dec 2021]
35. Goldberg D, Hutchinson S, Innes H, Dillon J. *Scotland's Hepatitis C Action Plan: Achievements of the first decade and proposals for a Scottish Government Strategy (2019) for the elimination of both infection and disease. Taking advantage of outstanding new therapies.* Available from: https://www.natap.org/2019/HCV/1_HCV-Elimination-Scotland-2019-07-31.pdf [Accessed 09 Dec 2021]

36. Hickman M, Dillon JF, Elliott L et al. Evaluating the population impact of hepatitis C direct acting antiviral treatment as prevention for people who inject drugs (EPIToPe) – a natural experiment (protocol). *BMJ Open*. 2019 Sep;9(9):e029538. doi.org/10.1136/bmjopen-2019-029538.
37. National Assembly for Wales. *Hepatitis C: Progress towards achieving elimination in Wales. June 2019*. Available from: <https://senedd.wales/laid%20documents/cr-ld12601/cr-ld12601-e.pdf> [Accessed 09 Dec 2021].
38. Dore GJ. Elimination of hepatitis C in Australia by 2030: a decade and counting. *Aust Prescr*. 2021 Apr;44(2):36-37. doi.org/10.18773/austprescr.2021.003
39. Hajarizadeh B, Grebely J, Matthews GV et al. Uptake of direct acting antiviral treatment for chronic hepatitis C in Australia. *J Viral Hepat*. 2018 Jun;25(6):640-648. doi.org/10.1111/jvh.12852.
40. Scottish School of Primary Care. *Remote and rural general practice in Scotland: descriptors and challenges. GP clusters briefing paper 20*. Available from: http://www.sspc.ac.uk/media/Media_683800_smxx.pdf [Accessed 09 Dec 2021]
41. National Records of Scotland. *Drug-related deaths in Scotland in 2020*. Available from: <https://www.nrscotland.gov.uk/files//statistics/drug-related-deaths/20/drug-related-deaths-20-pub.pdf> [Accessed 09 Dec 2021]
42. Hutchinson SJ, Roy KM, Wadd S et al. Hepatitis C virus infection in Scotland: Epidemiological review and public health challenges. *Scott Med J*. 2006 May;51(2):8-15. doi.org/10.1258/RSMJM.51.2.8.
43. Pohontsch NJ, Hansen H, Schäfer I, Scherer M. General practitioners' perception of being a doctor in urban vs. rural regions in Germany – A focus group study. *Fam Pract*. 2018 Mar;35(2):209-215. doi.org/10.1093/fampra/cmz083.

44. Carlin E, Alberti H, Davies K. Denigration of general practice as a career choice: The students' perspective. A qualitative study. BJGP Open. 2021 Feb;5(1).

doi.org/10.3399/bjgpopen20X101132

45. Speakman EM, Jarvis H, Whiteley D. Opportunities and risks within the expanding role of general practice. Br J Gen Pract. 2021 Jul;71(709):344-345.

doi.org/10.3399/bjgp21X716489.

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Figure 1: Sequential depiction of methodology, alongside pathway and recommendation development. Stages in green have been described previously¹⁵. BCW: Behaviour Change Wheel; HCV: Hepatitis C Virus; PLHCV: People living with HCV; TDF: Theoretical Domains Framework

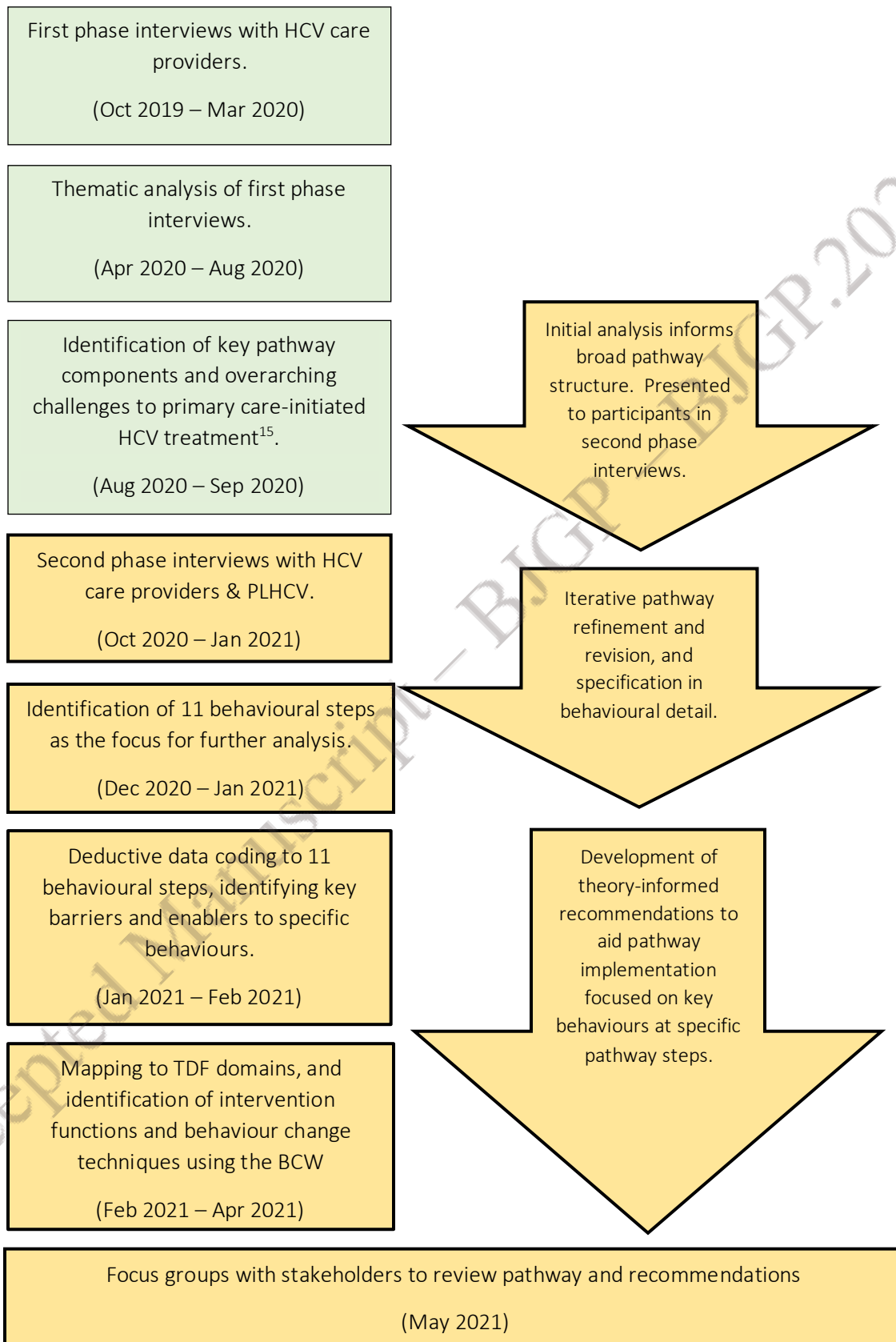
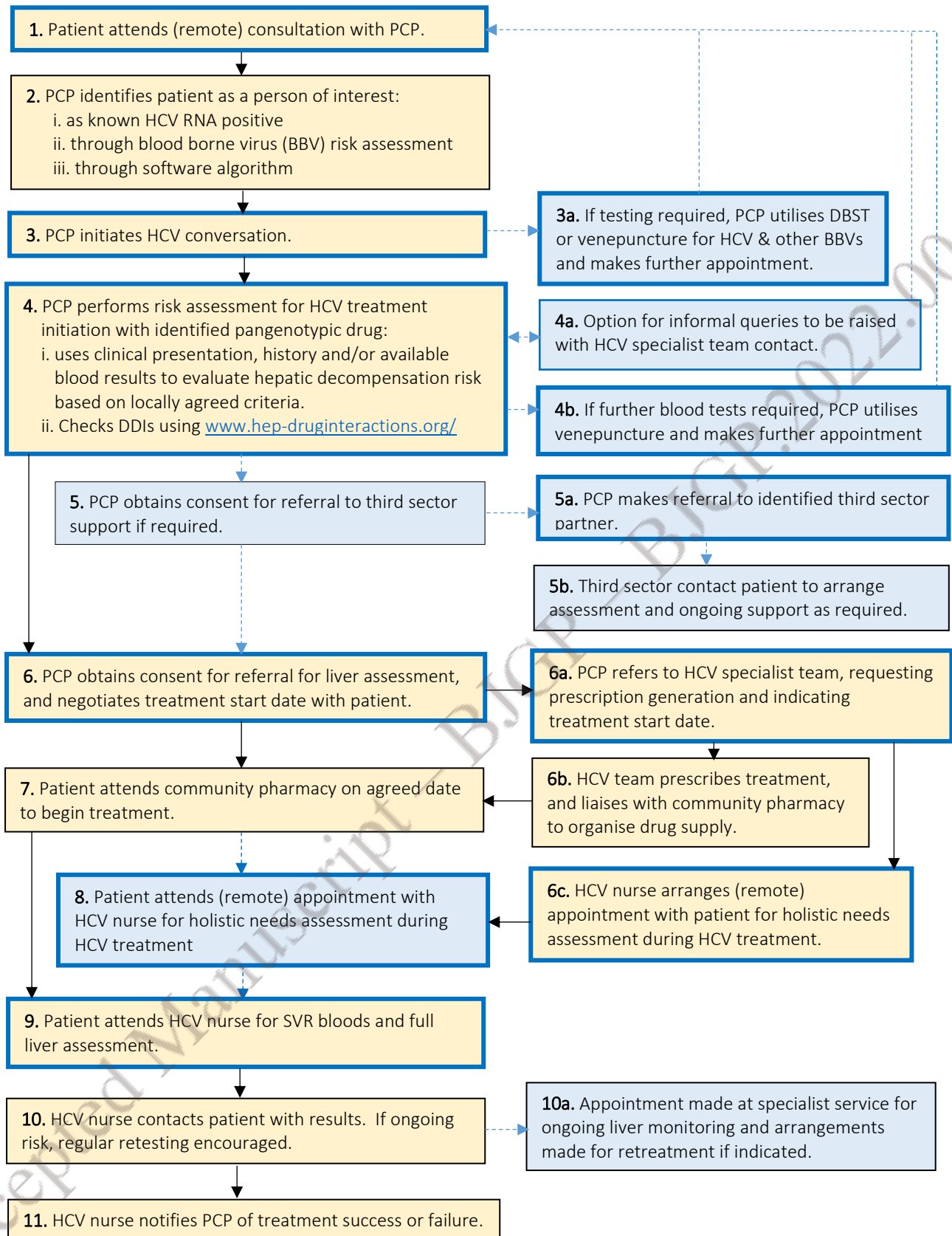


Figure 2: A pragmatic primary care-initiated hepatitis C virus (HCV) treatment pathway.



Key:

DBST (Dried Blood Spot Testing)	 Essential pathway step
DDI (Drug-Drug Interactions)	 Potential pathway step
PCP (Primary Care Provider)	 Pathway 'hotspot'
SVR (Sustained Virological Response)	