

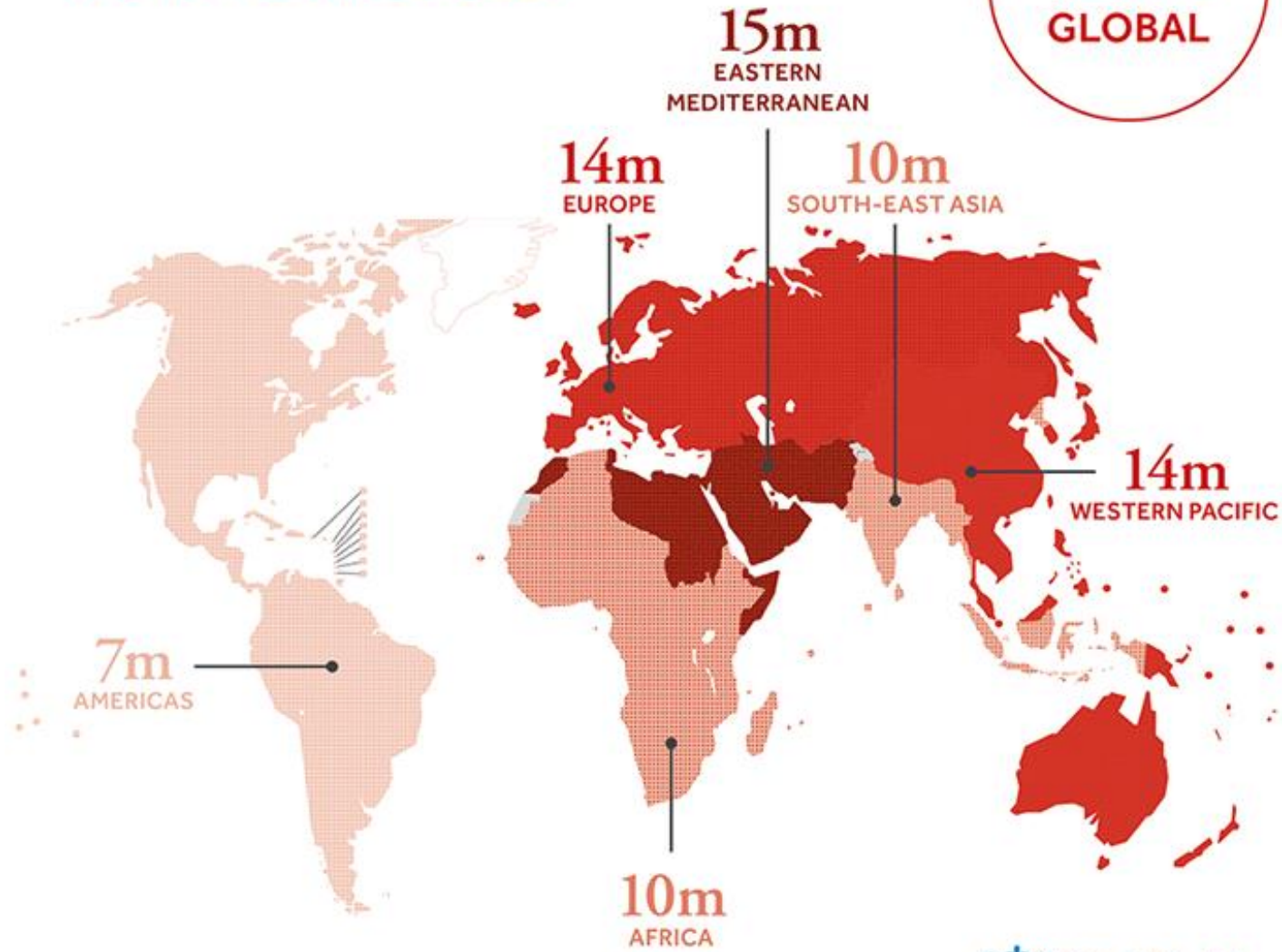
HCV: A UK view on treatment coverage

Dr Emma E Page



VIRAL HEPATITIS C IN THE WORLD

71m
GLOBAL



COMBATING HEPATITIS B AND C TO REACH ELIMINATION BY 2030

GHSS on viral hepatitis

MAY 2016



World Health
Organization

| Target areas | | Baseline 2015 | 2020 target | 2030 target |
|-------------------------------|---|---------------|------------------------------------|----------------------|
| | 4 Harm reduction (sterile syringe/needle set distributed per person per year for people who inject drugs [PWID]) | 20 | 200 | 300 |
| | 5 Treatment | | | |
| | 5a. Diagnosis of HBV and HCV (coverage %) | <5% | 30% | 90% |
| | 5b. Treatment of HBV and HCV (coverage %) | <1% | 5 million (HBV) 3 million (HCV) | 80% eligible treated |
| Impact leading to elimination | Incidence of chronic HBV and HCV infections | 6–10 million | 30% reduction | 90% reduction |
| | Mortality from chronic HBV and HCV infections | 1.46 million | 10% reduction | 65% reduction |

COMBATING HEPATITIS B AND C TO REACH ELIMINATION BY 2030

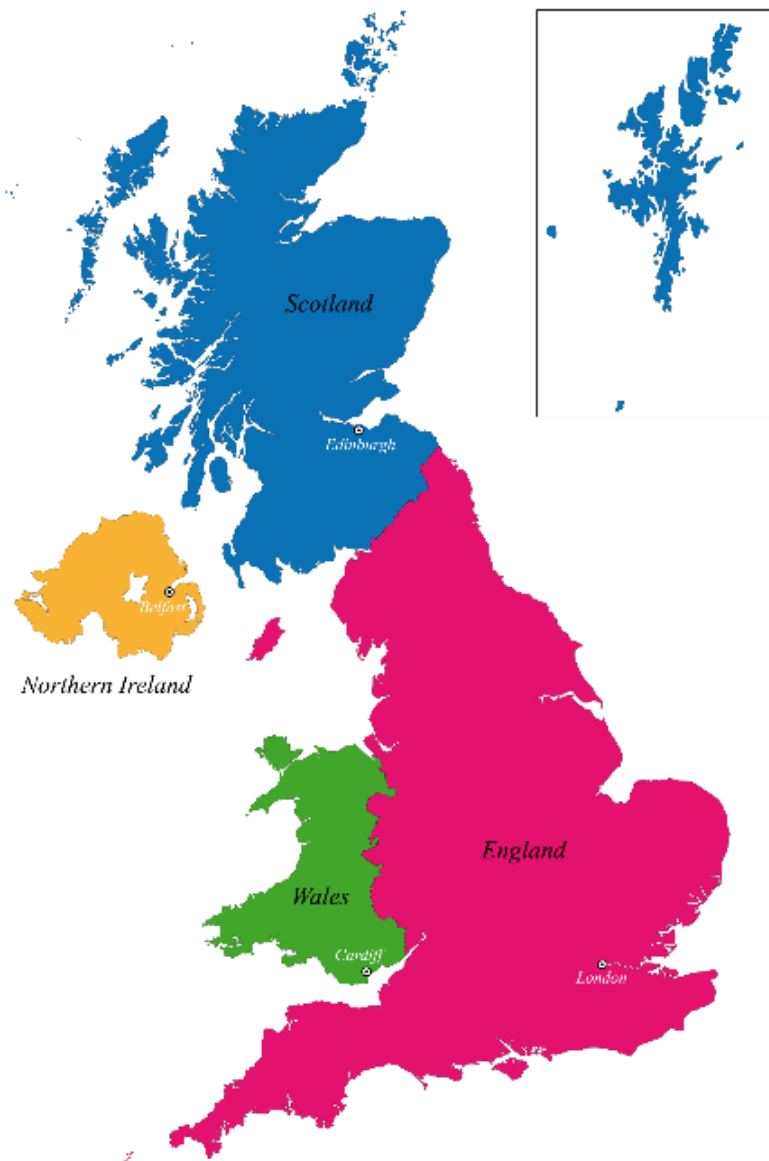
GHSS on viral hepatitis
MAY 2016



World Health
Organization

Updated for
WHO
European
Region

| Target areas | | Baseline 2015 | 2020 target | 2030 target |
|-------------------------------|---|---------------|---------------|----------------------|
| | 4 Harm reduction (sterile syringe/needle set distributed per person per year for people who inject drugs [PWID]) | 20 | 200 | 300 |
| | 5 Treatment | | | |
| | 5a. Diagnosis of HBV and HCV (coverage %) | <5% | 75% | 90% |
| | 5b. Treatment of HBV and HCV (coverage %) | <1% | 50% | 80% eligible treated |
| Impact leading to elimination | Incidence of chronic HBV and HCV infections | 6–10 million | 30% reduction | 90% reduction |
| | Mortality from chronic HBV and HCV infections | 1.46 million | 10% reduction | 65% reduction |



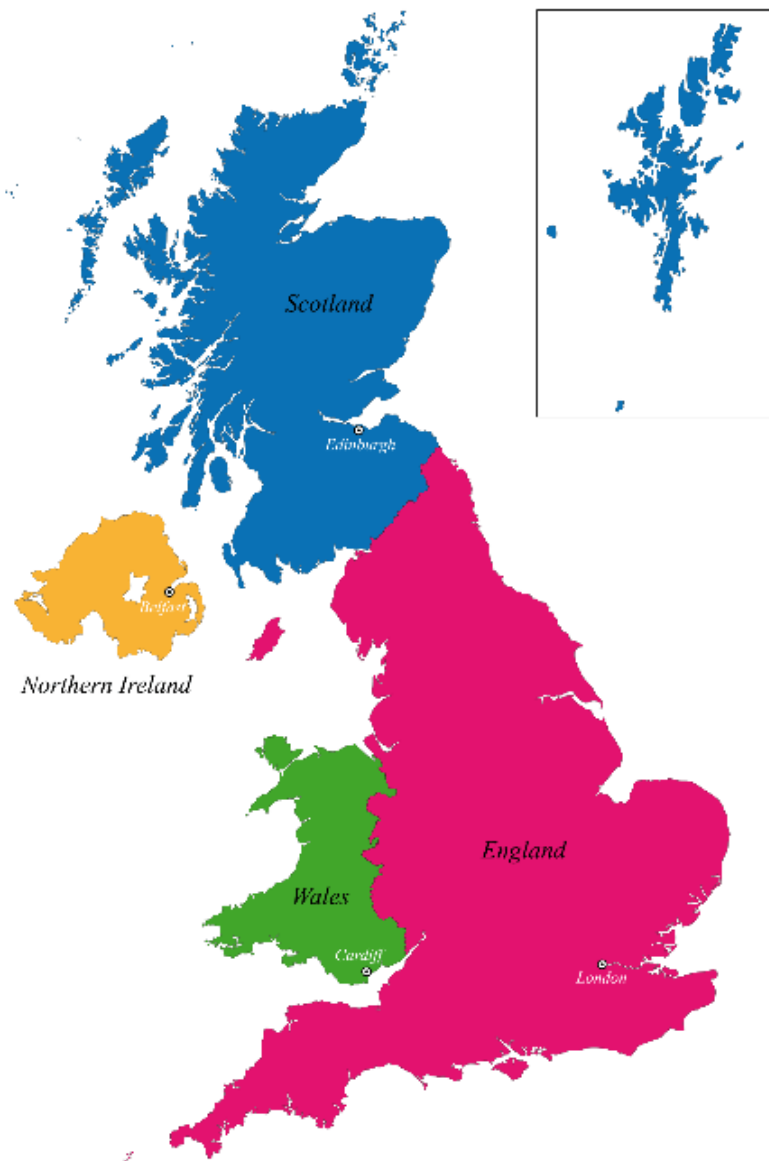
HCV in the UK – 2020 targets

- 200,000 people have chronic HCV



HCV in the UK – 2020 targets

- 200,000 people have chronic HCV
 - 60% PWID have ‘adequate’ syringe/needle provision

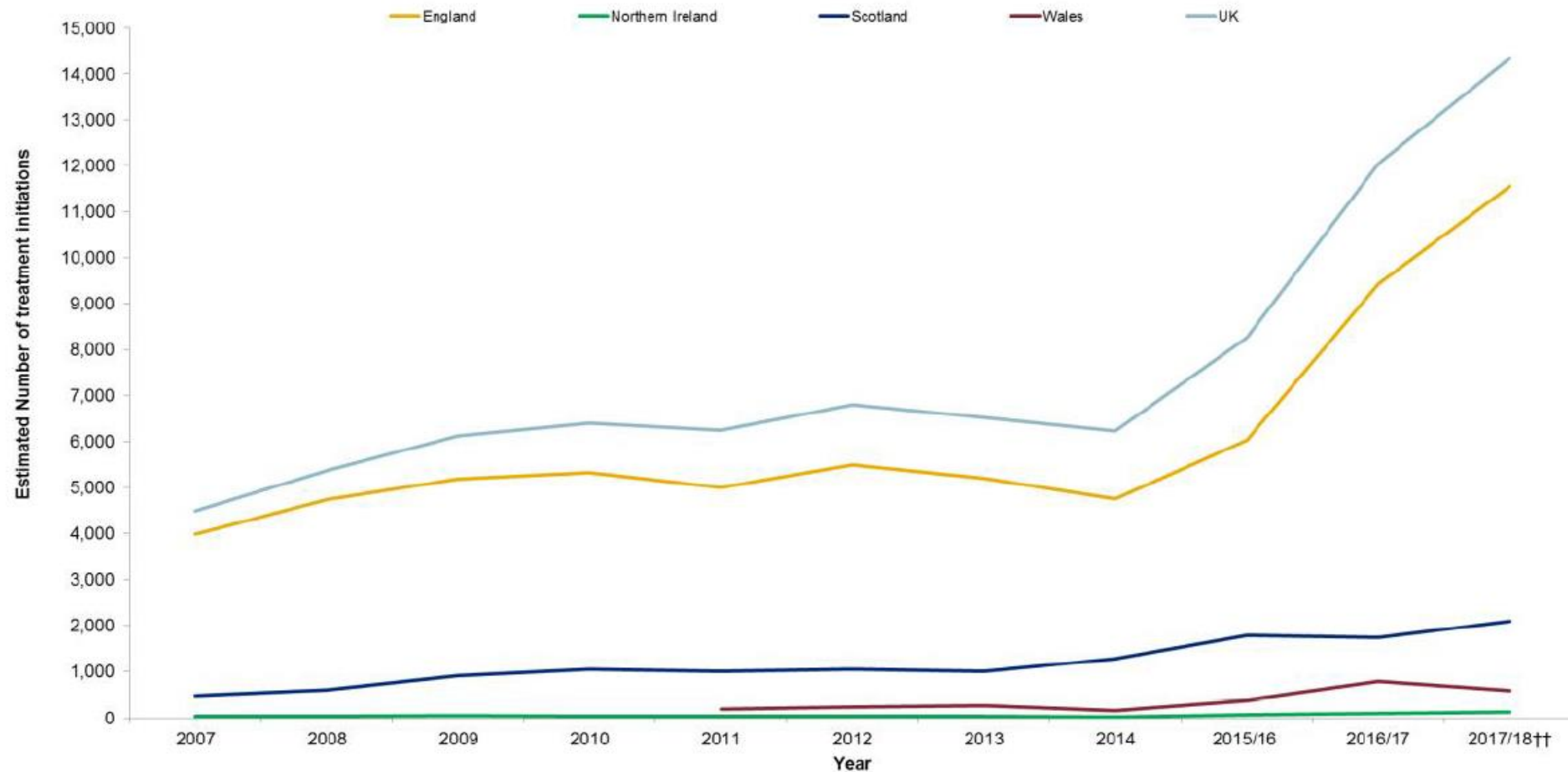


HCV in the UK – 2020 targets

- 200,000 people have chronic HCV
 - 60% PWID have ‘adequate’ syringe/needle provision
 - Estimated 66% aware of status (75%)



Figure 3. UK-wide estimates of numbers initiating HCV treatment, calendar years 2007 to 2014 and financial years 2015 to 2016 – 2017 to 2018^{*,}**



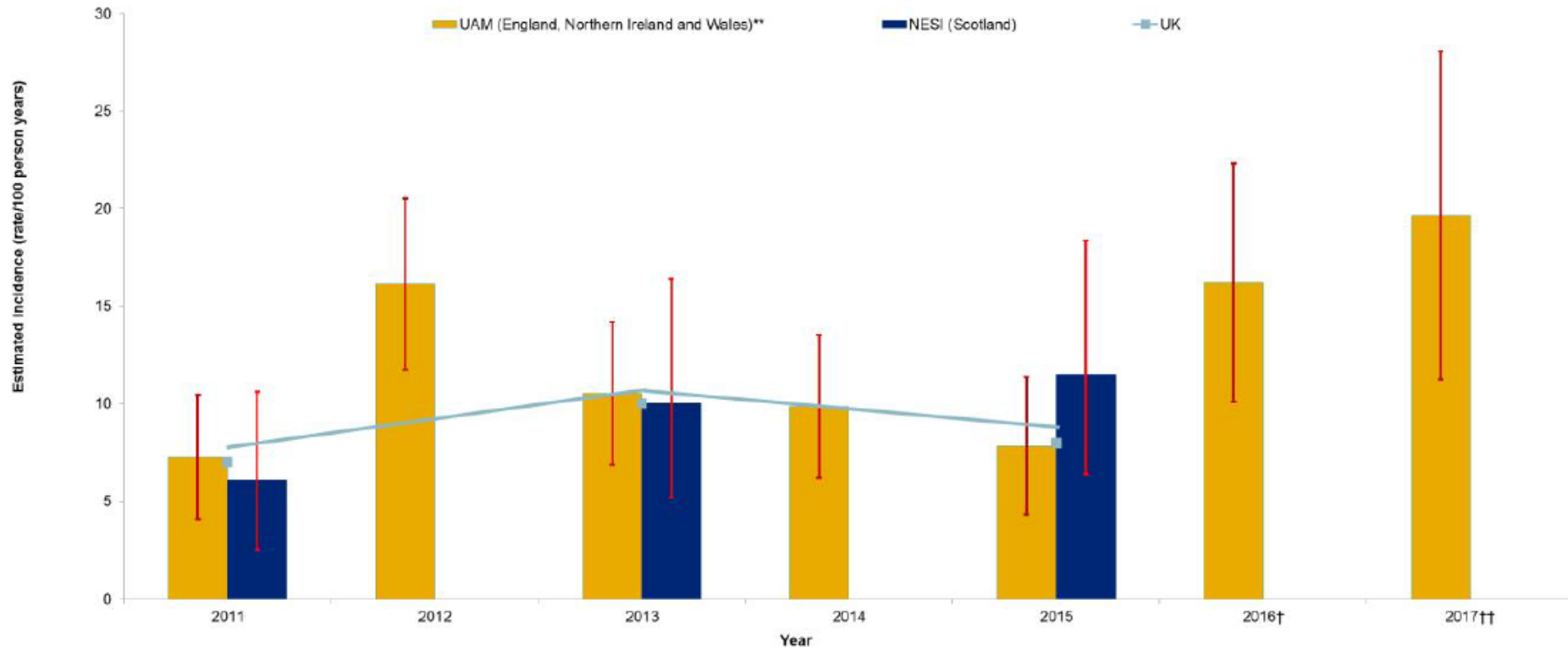


HCV in the UK – 2020 targets

- 200,000 people have chronic HCV
 - 60% PWID have ‘adequate’ syringe/needle provision
 - Estimated 66% aware of status (75%)
 - Currently around 14,000/yr (50%)
 - 125% increase from pre-2015
 - 19% increase from 2016/2017



Figure 8. Estimated UK-wide incidence of HCV among PWID, 2011-2017*,**



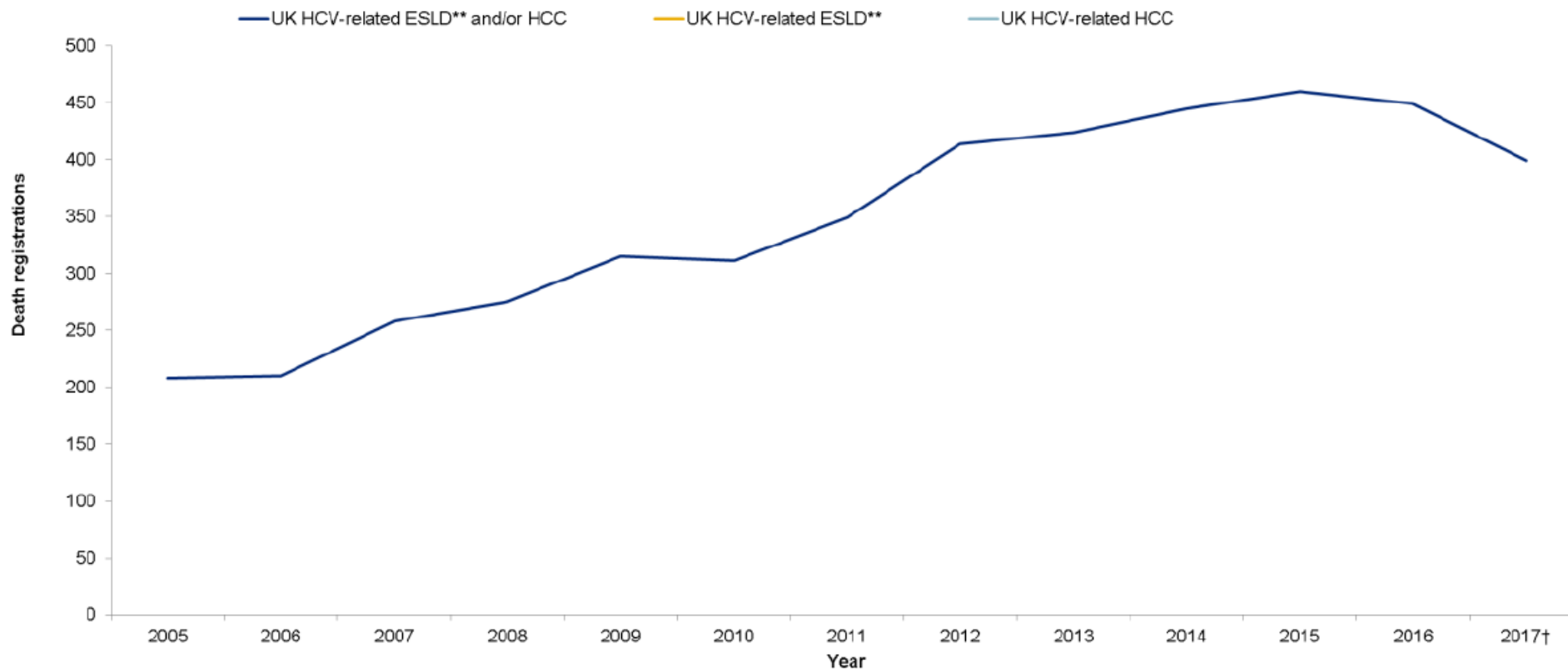


HCV in the UK – 2020 targets

- 200,000 people have chronic HCV
 - **60% PWID have 'adequate' syringe/needle provision**
 - **Estimated 66% aware of status (75%)**
 - **Currently around 14,000/yr**
 - 125% increase from pre-2015
 - 19% increase from 2016/2017
 - **Incidence in PWIDs appears to be increasing (50% reduction)**



Figure 6. Death registrations* for HCV-related ESLD and HCC in UK***
Countries: 2005 to 2017**





HCV in the UK – 2020 targets

- 200,000 people have chronic HCV
 - 60% PWID have ‘adequate’ needle/syringe provision
 - Estimated 66% aware of status (75%)
 - Currently around 14,000/yr
 - 125% increase from pre-2015
 - 19% increase from 2016/2017
 - Incidence in PWIDs appears to be increasing (50% reduction)
 - Early estimates suggest on target for 10% reduction in deaths by 2020



Public Health
England

Hepatitis C in the UK

2018 report

In the UK, eliminating hepatitis C as a major public health threat by driving down HCV-related mortality and preventing new infections from occurring is potentially feasible with the tools currently available. Investment in 3 core intervention areas is needed: (i) ensuring adequate harm reduction for PWID, (ii) increasing the proportion of infected individuals who are diagnosed, and (iii) increasing the proportion of infected individuals who access and complete treatment, achieving a sustained virological response (SVR).

HCV in the UK



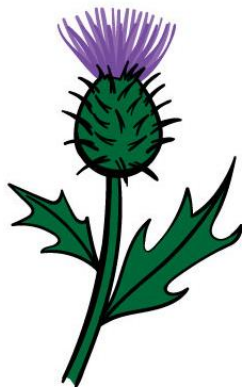
England

- 0.4 % prevalence
- 113,000 chronic HCV
- 33% undiagnosed



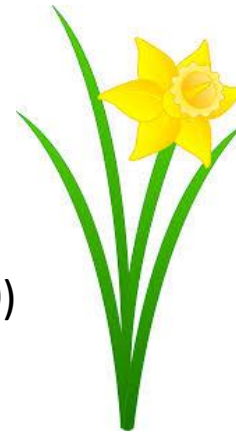
N.Ireland

- 0.4% prevalence
- 13,000 chronic HCV
- 50% undiagnosed



Scotland

- 0.7% prevalence
- 34,500 chronic HCV
(Recent estimates HPS: 29,000)
- 40% undiagnosed

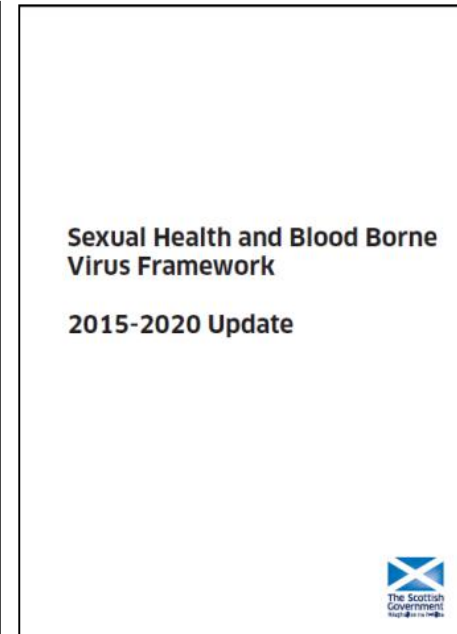
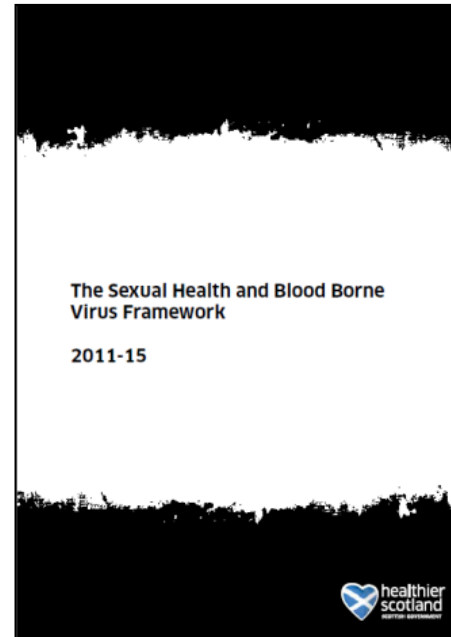
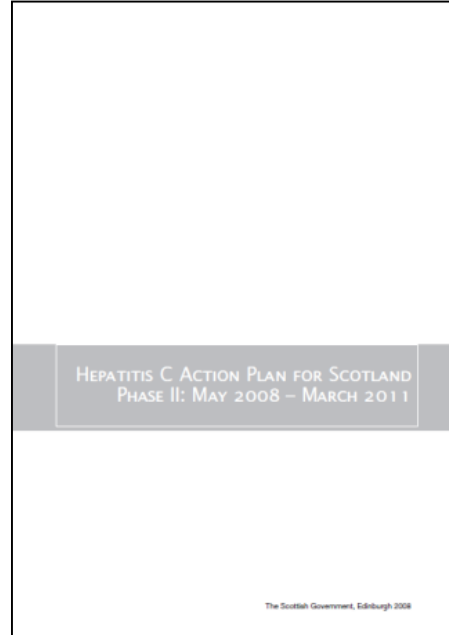
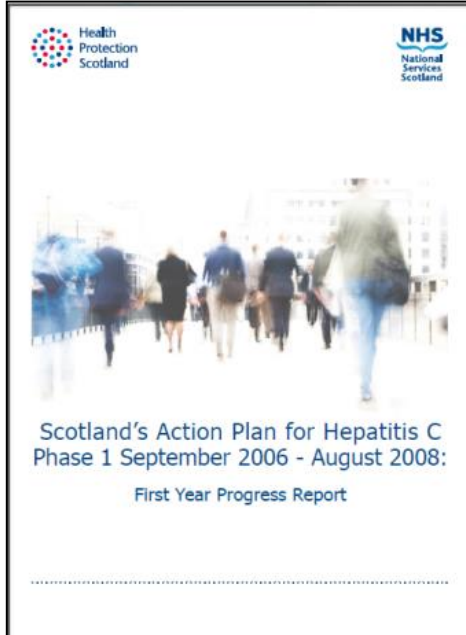


Wales

- 0.4% prevalence
- 13,000 chronic HCV
- 50% undiagnosed

Treatment coverage: Scotland

Past



Treatment coverage: Scotland

Current



Scottish
Medicines
Consortium

- DAAs approved by SMC with few restrictions
 - National guidelines guide treatment
 - Budget constraints
 - Priority to most advanced disease (until 2018)
 - Ultimate goal: offer all treatment TasP
- Move treatment to prisons, drug services & pharmacies.....

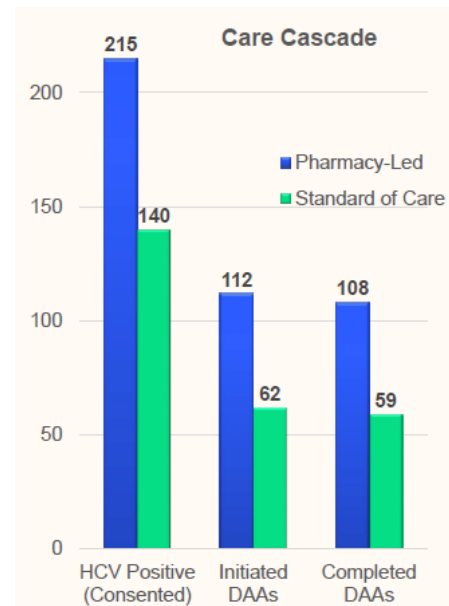
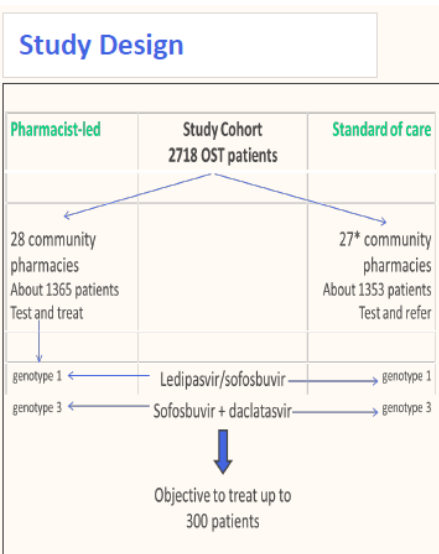
Treatment coverage: Scotland

Current



PRELIMINARY ANALYSIS OF THE SUPERDOT-C STUDY: A CLUSTER RANDOMISED CONTROLLED TRIAL OF PHARMACY LED VERSUS CONVENTIONAL TREATMENT FOR HCV POSITIVE PATIENTS RECEIVING DAILY OPIOD SUBSTITUTION THERAPY WITHIN NHS SCOTLAND

Radley AS¹, de Bruin M², Inglis SK³, Donnan PT³, Beer LJ³, Barclay S⁴, Fraser A⁵, Dillon JF¹
University of Dundee¹, University of Aberdeen², Tayside Clinical Trials Unit³, NHS Greater Glasgow & Clyde⁴, NHS Grampian⁵



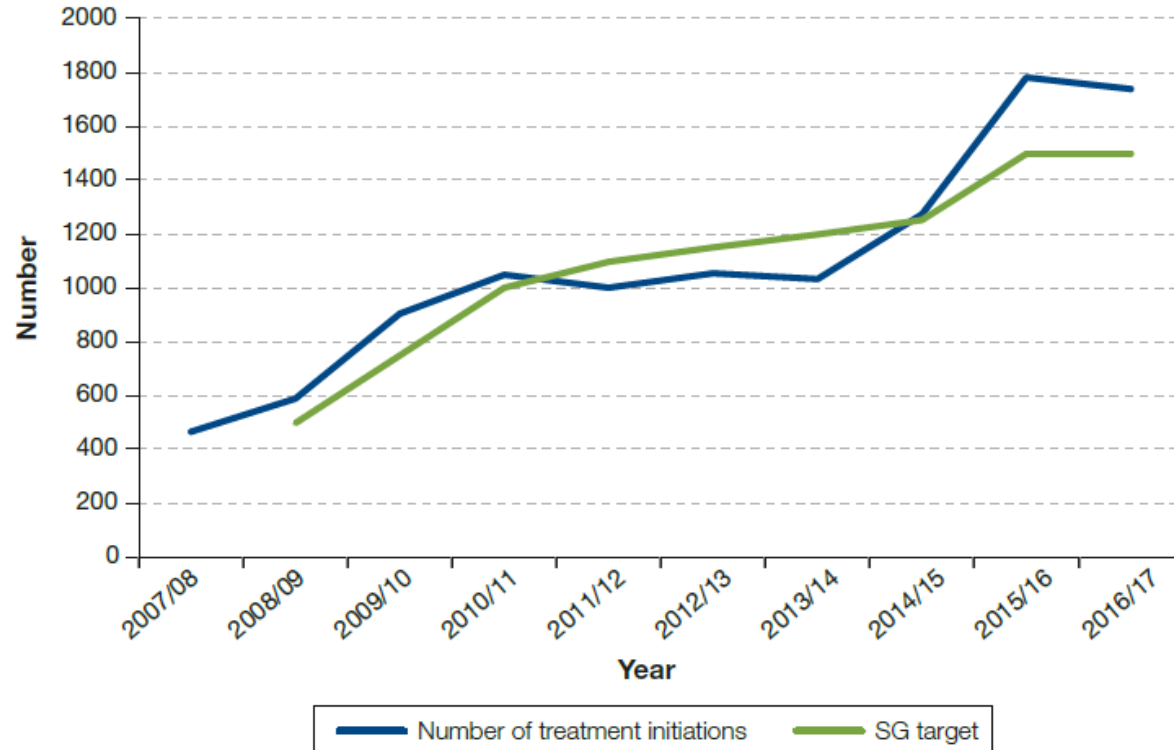
- 55 Pharmacies
- Pharmacy treatment sites increased:
 - numbers identified
 - numbers treated
- SVR12 74% vs 58% (incomplete data)
- Tayside: 'diagnosed 80% and treated 70%'

Treatment coverage: Scotland

Current



Figure 14: Estimated numbers initiating HCV treatment in Scotland vs. national treatment targets, 2007-2016.



Scale up in treatment numbers:

- 450 a year 2007
- 1,739 a year 2016/17

Treatment rates still rising

- 2,000 2018/19
- Target of 2,500 by 2020

Treatment coverage: Scotland Future



Eliminating Hepatitis C in Scotland: A Call to Action


A summary of evidence from the Hepatitis C Elimination Inquiry held by the cross-party Scottish Hepatitis C Parliamentary Champions group and The Hepatitis C Trust

- Committed to elimination HCV by 2030
- Not on track to meet GHSS 2030 targets
- Highlighted number of issues around access to treatment:
 - Budget constraints = prioritisation
 - Some areas not treating current PWID
 - LTC improvement needed
 - Delivering treatment in the community
 - Reported Increased incidence in PWID
- Outcome: Scottish Government to develop Hepatitis C elimination strategy
.....currently awaiting publication

Treatment coverage: England



Current

|  | |
|---|--|
| SCHEDULE 2 – THE SERVICES | |
| A. Service Specifications | |
| Service Specification No. | F04 S f |
| Service | Operational Delivery Networks for Hepatitis C Care in Adults |
| Commissioner Lead | NHS England |
| Provider Lead | |
| Period | 01.04.2015 to 31.03.2016 |
| Date of Review | 31.03.2017 |

- Ensure equitable access to treatments
- And act as gate keepers to DAAS:
 - All patients treated must be discussed at ODN
 - Access to DAA controlled by NHSE

HEP C ODNs AND CLINICAL LEADS



Treatment coverage: England

Current

Interim Clinical Commissioning
Policy Statement:
Sofosbuvir + Daclatasvir/Ledipasvir
+/- Ribivirin for defined patients with
Hepatitis C
April 2014
Reference: NHS ENGLAND A02/PS/b



Treatment coverage: England

Current

Interim Clinical Commissioning
Policy Statement:
Sofosbuvir + Daclatasvir/Ledipasvir
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April 2014
Reference: NHS ENGLAND A02/PS/b



June 2015

Clinical Commissioning Policy
Statement:
Treatment of chronic Hepatitis C in
patients with cirrhosis

Treatment coverage: England

Current

Interim Clinical Commissioning Policy Statement:
Sofosbuvir + Daclatasvir/Ledipasvir +/- Ribivirin for defined patients with Hepatitis C
April 2014
Reference: NHS ENGLAND A02/PS/b



Dec 2015 'rate cards' = treatment for all*

- Drug choice: genotype, cirrhosis, past treatment
- Each ODN associated run rate with penalties and cap
- Updated 6 mnthly after NICE approval and tenders bt NHS E & pharma



June 2015

Clinical Commissioning Policy Statement:
Treatment of chronic Hepatitis C in patients with cirrhosis

West Yorkshire ODN Network - Summary of treatment guidelines for chronic HCV infection
Revised September 2017

| Genotype | Cirrhosis | Treatment experience | Drug regime |
|----------|-------------------------|-------------------------------------|------------------------------|
| G1 | No cirrhosis | Naive | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| | | G1a (or G1 without sub-genotype) | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| G1 | Compensated cirrhosis | Naive | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| | | G1a (or G1 without sub-genotype) | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| G1 | Decompensated cirrhosis | Naive | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| | | G1a (or G1 without sub-genotype) | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| G2 | No cirrhosis | Naive or Peg/NS5B treatment failure | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| | | Compensated cirrhosis | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| G2 | Compensated cirrhosis | Naive or Peg/NS5B treatment failure | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| | | Decompensated cirrhosis | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| G3 | No cirrhosis | Naive or Peg/NS5B treatment failure | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| | | Compensated cirrhosis | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| G3 | Compensated cirrhosis | Naive or Peg/NS5B treatment failure | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| | | Decompensated cirrhosis | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |

| Genotype | Cirrhosis | Treatment experience | Drug regime |
|----------|-----------------------|-------------------------|------------------------------|
| G4 | No cirrhosis | Naive | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| | | Compensated cirrhosis | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| G4 | Compensated cirrhosis | Naive | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |
| | | Decompensated cirrhosis | Sofo/led ± ribi* 12 weeks |
| | | | Sofo/led ± ribi* 8 weeks |
| | | | Sofo/led ± ribi* 12 weeks |

3. Response to consider 2* one treatment include: non-sustained ODN with 2* or 2* one treatment, absolute contraindication or previous severe intolerance to 2*
4. Cirrhosis is defined as liver biopsy evidence of cirrhosis, e.g. ICG R15 score > 14% (or ICG R15 score > 14% and MELD score > 10) or histological evidence (Schussler, CT or MRI) of liver cirrhosis or evidence of portal hypertension (e.g. varices, ascites) without other obvious cause.
5. Decompensated cirrhosis is defined as: previous or ongoing episode of ascites, variceal bleeding or encephalopathy and/or Child Pugh score > 7 (B or C) and/or bilirubin > 0.5 mg/dL at treatment baseline.
6. For G3 infected compensated cirrhotic patients being treated with sofosbuvir/ledipasvir, add weight-based RBV if treatment experienced, ideally add RBV to all G3 compensated cirrhotic patients, regardless of treatment response. For significant RBV use:
7. All G1a treatment-naïve patients offered 12 weeks sofosbuvir/ledipasvir/sofosbuvir + daclatasvir rather than considering certain patients potentially suitable for 8 weeks.
8. **Genotype 1 HCV RNA > 100,000 IU/mL** - add EDC4 sample to genotype for NS5A resistance associated substitution (RAS) testing - 4 significant NS5A RAS at baseline (significant NS5A RAS = L159V, Y155H, H58R, Y93H, Y93H/H) - treat with sofosbuvir/ledipasvir + weight-based RBV for 16 weeks. If no significant NS5A RAS at baseline, can treat with sofosbuvir/ledipasvir without RBV for 12 weeks. If significant NS5A RAS present for genotype 1, no RBV tolerance, consider referral to HCV for 2* one RBV-free regimen. **Genotype 1 HCV RNA > 100,000 IU/mL** - treat with sofosbuvir/ledipasvir (no RBV) for 12 weeks.
9. Consider consent weight-based RBV for G1 patients with cirrhosis who are either treatment-experienced and/or who have genotype 1a (1*).
10. For G1a patients with compensated cirrhosis being treated with sofosbuvir/ledipasvir + daclatasvir with weight-based RBV. If either sofosbuvir/ledipasvir/sofosbuvir non-responder and/or if 1* Peg/NS5B, add-on 2* and/or 2* RBV, then treat with 24 weeks sofosbuvir/ledipasvir/sofosbuvir + daclatasvir with RBV, otherwise treat for 12 weeks.

Treatment coverage: England

Current



Interim Clinical Commissioning Policy Statement:
Sofosbuvir + Daclatasvir/Ledipasvir +/- Ribivirin for defined patients with Hepatitis C
April 2014
Reference: NHS ENGLAND A02/PS/b



Dec 2015 'rate cards' = treatment for all*

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June 2015

Clinical Commissioning Policy Statement:
Treatment of chronic Hepatitis C in patients with cirrhosis

West Yorkshire ODN Network - Summary of treatment guidelines for chronic HCV infection
Revised September 2017

| Genotype | Cirrhosis | Treatment experience | Drug regime |
|-------------------------|-----------------------|---|---|
| G1 | No cirrhosis | Naive | <p>WFOA Ombitasvir/paritaprevir + dasabuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Esofenavir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Sofosbuvir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> |
| | | <p>or</p> <p>or</p> | |
| | | <p>or</p> <p>or</p> | |
| | Compensated cirrhosis | Naive | <p>WFOA Ombitasvir/paritaprevir + dasabuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Esofenavir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Sofosbuvir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> |
| | | <p>or</p> <p>or</p> | |
| | | <p>or</p> <p>or</p> | |
| Decompensated cirrhosis | Naive | <p>WFOA Sofosbuvir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> | |
| | <p>or</p> <p>or</p> | | |
| | <p>or</p> <p>or</p> | | |
| G2 | No cirrhosis | Naive | <p>WFOA Ombitasvir/paritaprevir + dasabuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Esofenavir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Sofosbuvir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> |
| | | <p>or</p> <p>or</p> | |
| | | <p>or</p> <p>or</p> | |
| | Compensated cirrhosis | Naive | <p>WFOA Ombitasvir/paritaprevir + dasabuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Esofenavir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Sofosbuvir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> |
| | | <p>or</p> <p>or</p> | |
| | | <p>or</p> <p>or</p> | |
| Decompensated cirrhosis | Naive | <p>WFOA Sofosbuvir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> | |
| | <p>or</p> <p>or</p> | | |
| | <p>or</p> <p>or</p> | | |
| G3 | No cirrhosis | Naive | <p>WFOA Ombitasvir/paritaprevir + dasabuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Esofenavir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Sofosbuvir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> |
| | | <p>or</p> <p>or</p> | |
| | | <p>or</p> <p>or</p> | |
| | Compensated cirrhosis | Naive | <p>WFOA Ombitasvir/paritaprevir + dasabuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Esofenavir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Sofosbuvir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> |
| | | <p>or</p> <p>or</p> | |
| | | <p>or</p> <p>or</p> | |
| Decompensated cirrhosis | Naive | <p>WFOA Sofosbuvir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> | |
| | <p>or</p> <p>or</p> | | |
| | <p>or</p> <p>or</p> | | |

| Genotype | Cirrhosis | Treatment experience | Drug regime |
|----------|--------------|----------------------|---|
| G4 | No cirrhosis | Naive | <p>WFOA Ombitasvir/paritaprevir + dasabuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Esofenavir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Sofosbuvir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> |
| | | <p>or</p> <p>or</p> | |
| | | <p>or</p> <p>or</p> | |
| G4a | No cirrhosis | Naive | <p>WFOA Ombitasvir/paritaprevir + dasabuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Esofenavir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> <p>WFOA Sofosbuvir/sofosbuvir + sofosbuvir + weight based RBV + ribavirin 12 weeks</p> |
| | | <p>or</p> <p>or</p> | |
| | | <p>or</p> <p>or</p> | |

3. Response to consider **2** the treatments include: re-treatable ODNs with **1** or **2** the treatments, absolute contraindication or previous severe intolerance to sofosbuvir.

4. Cirrhosis is defined as liver biopsy evidence of cirrhosis, e.g. ICG/APRI score 5 or 6 (METAVIR score F3-4 or F4 or liver biopsy fibrosis (e.g. Fibroscan™) evidence of cirrhosis, e.g. Fibroscan™ median 12.111 kPa or HCV fibrosis score 3 or equivalent on other disease scales or BSA/TAT 1.6 and APRI score 10.0 or radiological evidence (ultrasound, CT or MRI) of liver cirrhosis or evidence of portal hypertension (e.g. varices, ascites) without other obvious cause.

5. Decompensated cirrhosis is defined as: previous or ongoing episode of ascites, variceal bleeding or encephalopathy and/or Child Pugh score 7 (B or C) and/or bilirubin >0.5µmol/L at treatment baseline.

6. For G3 fibrotic compensated cirrhotic patients being treated with sofosbuvir/sofosbuvir, add weight-based RBV if treatment experienced, ideally add RBV to all G3 compensated cirrhotic patients, regardless of treatment response. For significant RBV use.

7. All G3 treatment-naïve patients offered 12 weeks ombitasvir/paritaprevir + dasabuvir rather than considering certain patients potentially suitable for 8 weeks.

8. **Excluded HCV RNA 100 IU/ml** and **COA** patients to therapy for NS5A resistance associated substitution (RAS) testing - 4 significant NS5A RAS at baseline (significant NS5A RAS = L159V, Y93H, A156V, S122R, Y93H, H159V, H159Y, H159L or Y93H/N) treat with sofosbuvir/sofosbuvir + weight based RBV for 16 weeks. If no significant NS5A RAS at baseline, can treat with sofosbuvir/sofosbuvir + weight based RBV for 12 weeks. If significant NS5A RAS present for 16 weeks, no RBV tolerance, consider referral to HCV for **2** the RBV-free regimen. **Excluded HCV RNA 100 IU/ml** treat with sofosbuvir/sofosbuvir (no RBV) for 12 weeks.

9. Consider interim weight-based sofosbuvir for G1 patients with cirrhosis who are either treatment experienced and/or who have passed 75kPa.

10. For G1a patients with compensated cirrhosis being treated with ombitasvir/paritaprevir + dasabuvir with weight based RBV. If either sofosbuvir/sofosbuvir, sofosbuvir/sofosbuvir or sofosbuvir + weight based RBV, then treat with 24 weeks ombitasvir/paritaprevir + dasabuvir with RBV, otherwise treat for 12 weeks.

Urgent Clinical Commissioning Policy Statement: Retreatment of Chronic Hepatitis C Infection in Adults with Advanced or Decompensated Cirrhosis

NHS England Reference: 170020/PS

A clinical commissioning policy statement is an interim commissioning position pending the formation of a Clinical Policy.

Sept 2017

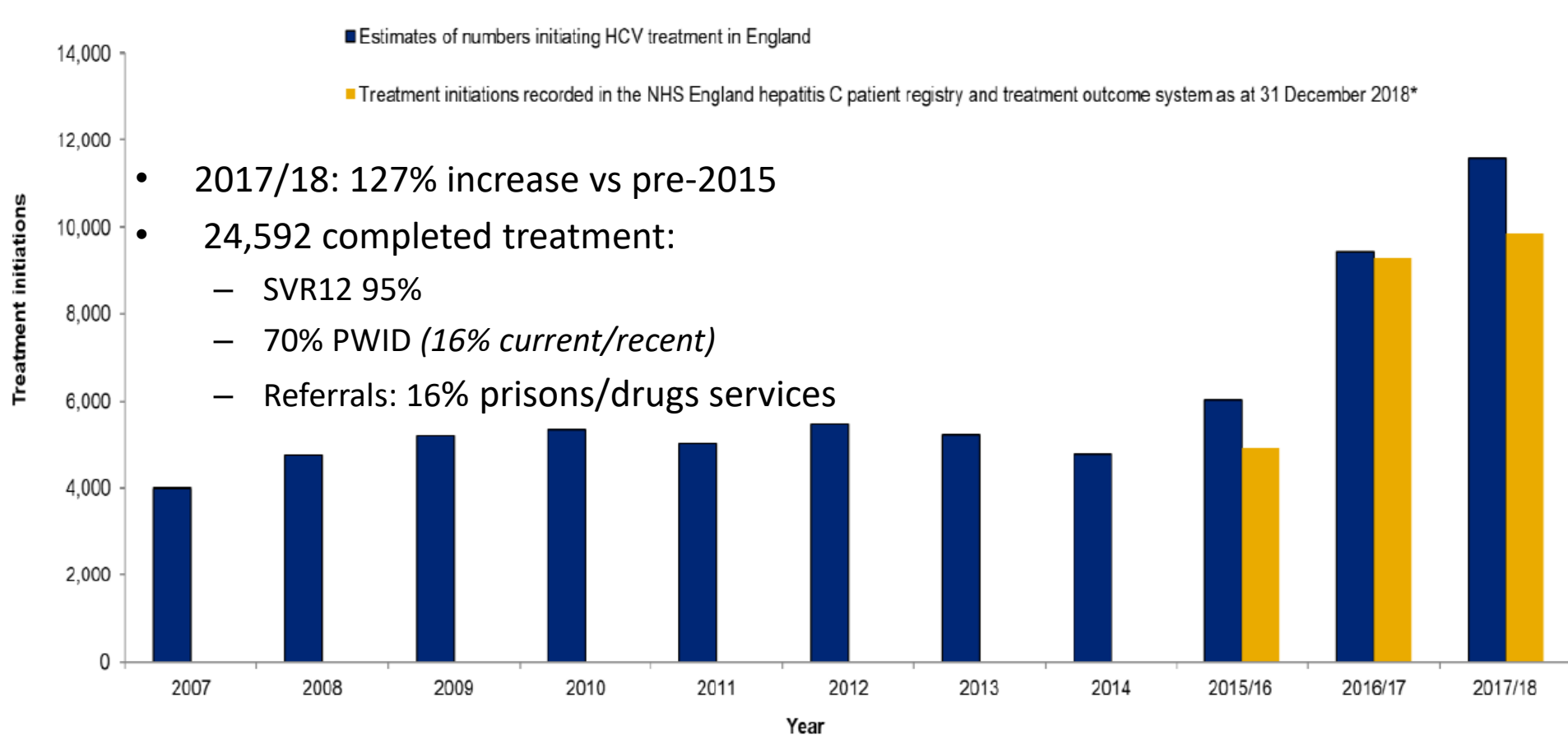




Treatment coverage: England

Current

Figure 26. Provisional estimates of numbers initiating HCV treatment in England, 2007-2017/2018



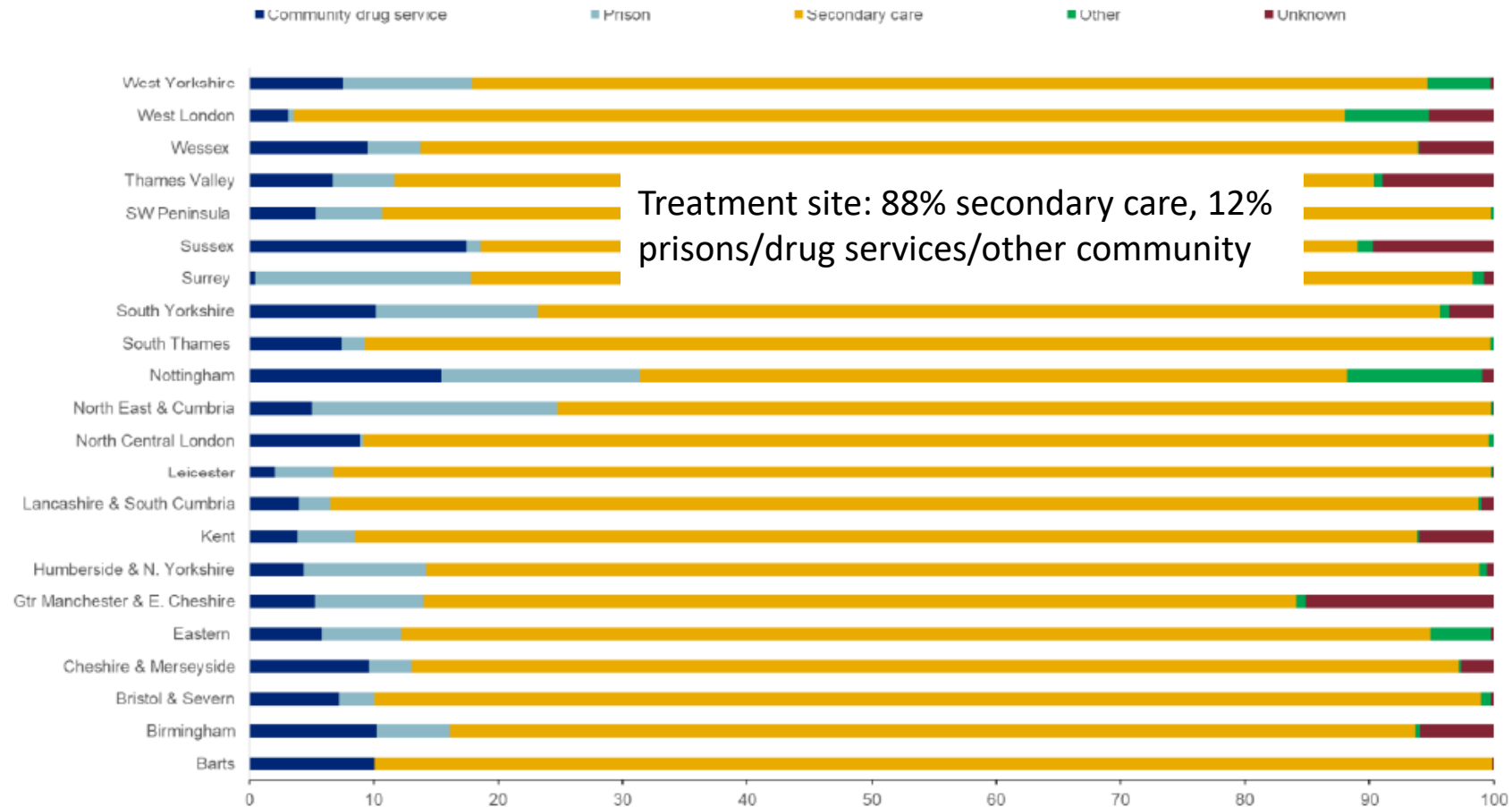


Treatment coverage: England

Current




Figure 32: Distribution of treatment setting (%) for patients with a treatment episode in the Hepatitis C Patient Registry and Treatment Outcome System, by ODN (n=30,870)



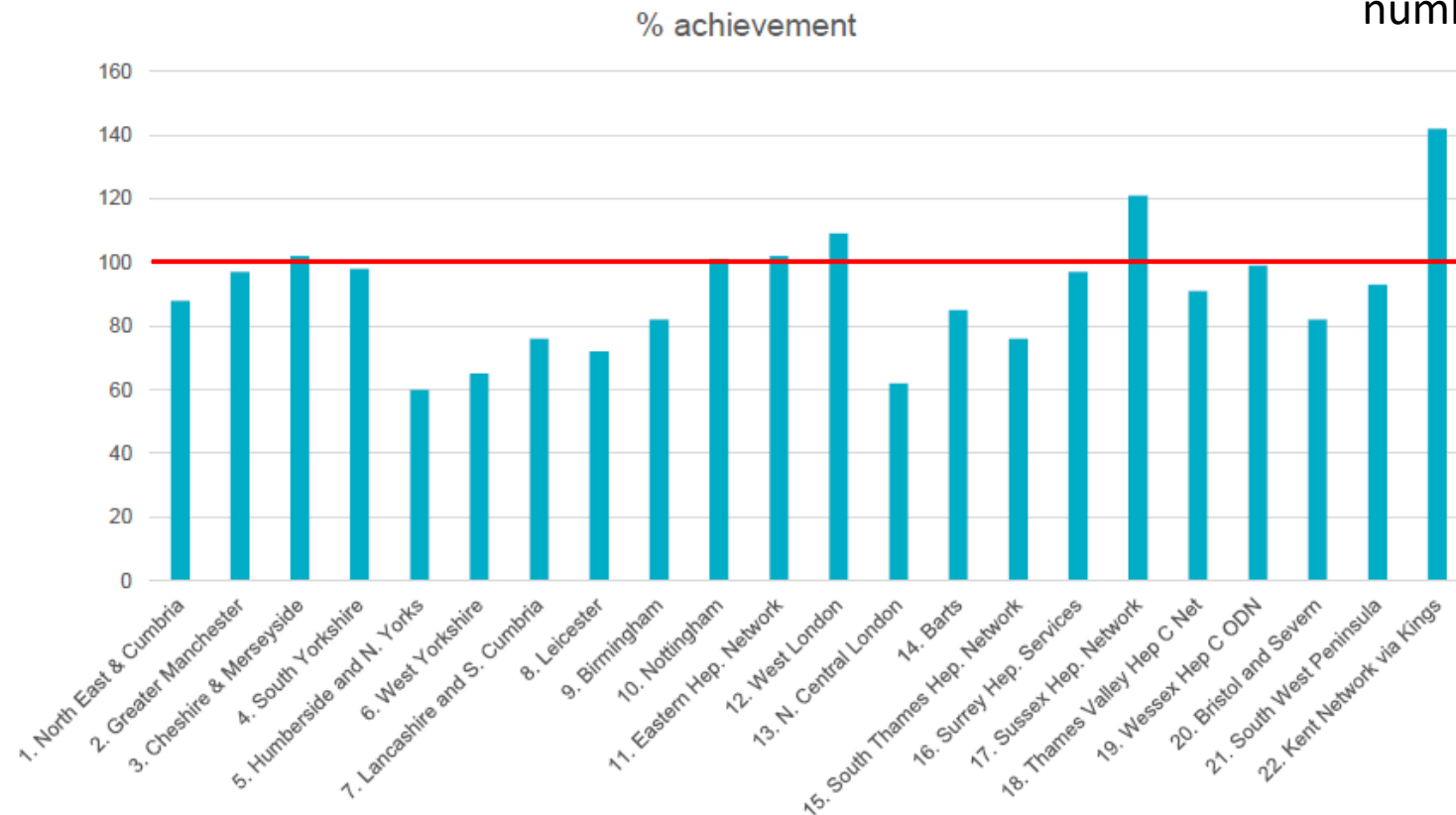
Treatment coverage: England

Current



2018/2019 ODN run rate % achievement

- 2018/29 run rate not achieved:
 - Over estimation of number of patients





Treatment coverage: England

Current

Hepatitis C treatment monitoring in England

Content, completeness and preliminary findings from the Hepatitis C patient registry and treatment outcome system

- 2018/29 run rate not achieved:
 - Over estimation of number of patients
 - ‘easy’ to engage patients treated
- Move treatment to the community

Table 9. Infection details for patients in the Hepatitis C patient registry and treatment outcome system yet to be treated compared to those with a treatment episode in the Register.

| Variable | Not yet treated (n= 7,816) | Treated (n=24,592) | Significance (P-value) |
|---|----------------------------|--------------------|------------------------|
| Injecting route of transmission (%) | | | <0.001 |
| Current/recent PWID (injected in past 3 years.) | 30.4 | 16.2 | |
| Past PWID | 38.7 | 46.3 | |

Treatment coverage: England

Current




Polaris Observatory *'without improved diagnosis rates number treated could fall to 5000/yr'*

- 2018/29 run rate not achieved:
 - Over estimation of number of patients
 - 'easy' to engage patients treated
 - Move treatment to the community
 - Still significant numbers undiagnosed:
 - Increase community testing

Treatment coverage: England Future



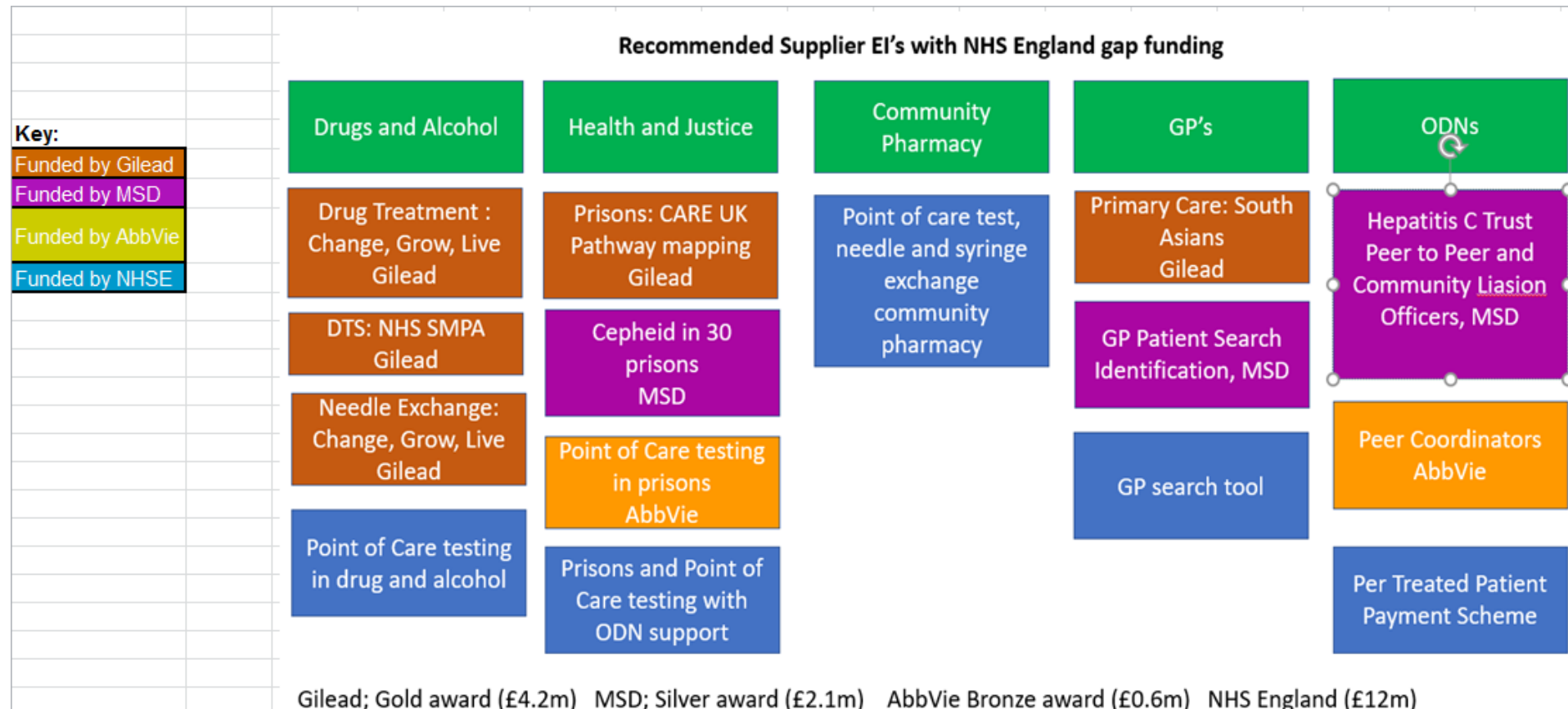
May 2019

- 3+2 year contract
- 3 bidders given medal status
- Rate card continues:
 - Gilead 60% 
 - MSD 24% 
 - AbbVie 17% 

NHS England has unveiled its plan to completely eliminate hepatitis C, signing up three pharma companies to the £1bn (\$1.29bn) programme which involves identifying and curing patients with drug treatments.

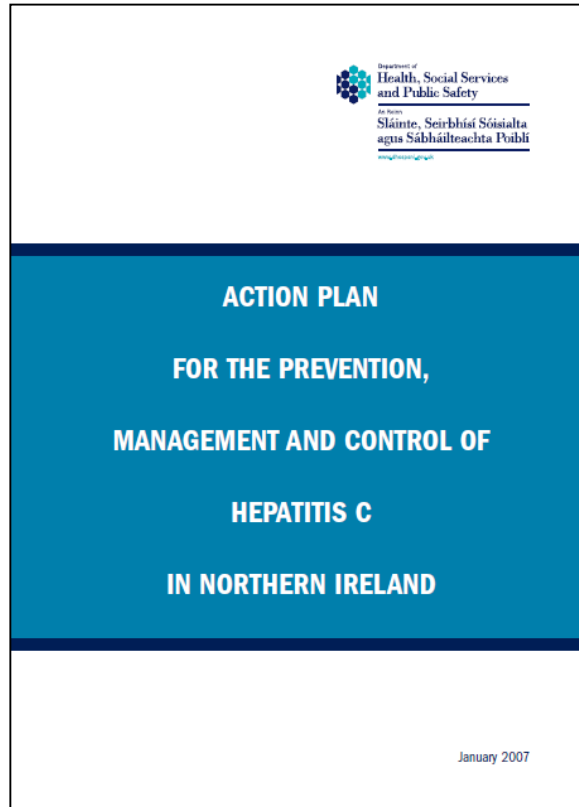
Treatment coverage: England

Future



Treatment coverage: N Ireland

Past



- Action plan since 2007
 - Prevention:
 - PWID
 - Healthcare settings
 - Prisons
 - Clinical services
 - Testing
 - Treatment
 - Establishment of Managed Clinical Network

Treatment coverage: N Ireland

Current



Northern Ireland Hepatitis B & C Managed Clinical Network

To reduce the transmission of Hep B & C (HBV & HCV) infection among injecting drug users by promoting routine and low threshold testing for Hep B & C infection among injecting drug users, and exploring best practice models for engaging and retaining injecting drug users in HBV & HCV treatment.

To promote timely diagnosis of HBV & HCV infection in NI (particularly in high risk groups), and to facilitate equitable access to high quality treatment of diagnosed HBV & HCV infections through the regional hepatology unit in RVH.

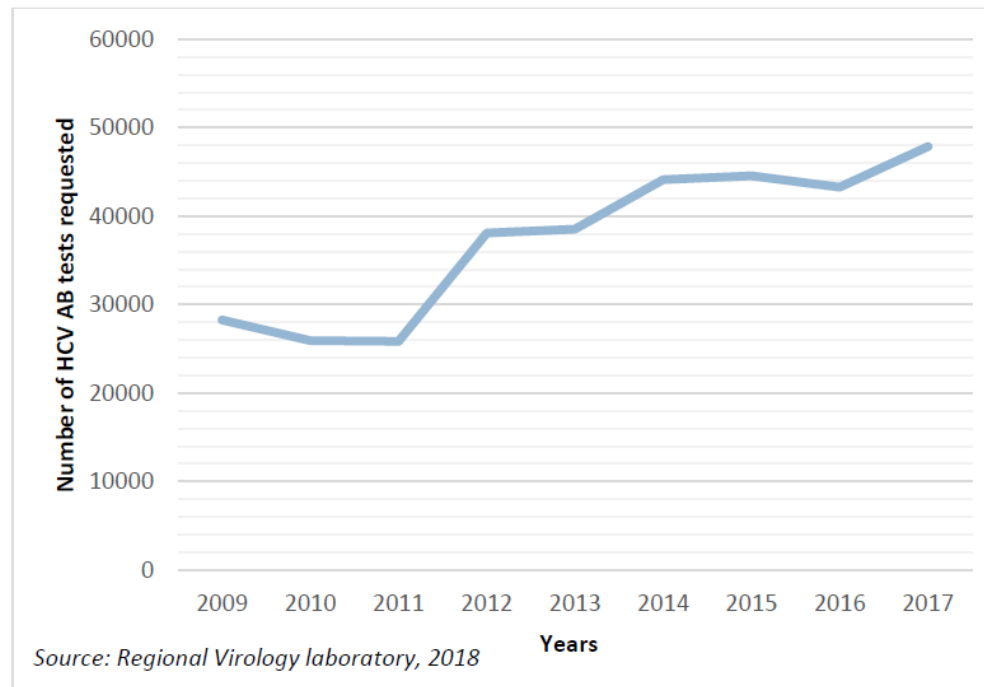
To deliver all treatment of HBV and HCV infection in NI through the regional hepatology unit, using treatment guidelines based on NICE recommendations;

Treatment coverage: N Ireland

Current

NI Regional Hepatitis B&C
Managed Clinical Network
Annual Report 2018

FIGURE 6: NUMBER THE HEPATITIS C TEST REQUESTS IN NORTHERN IRELAND, 2009 - 2017



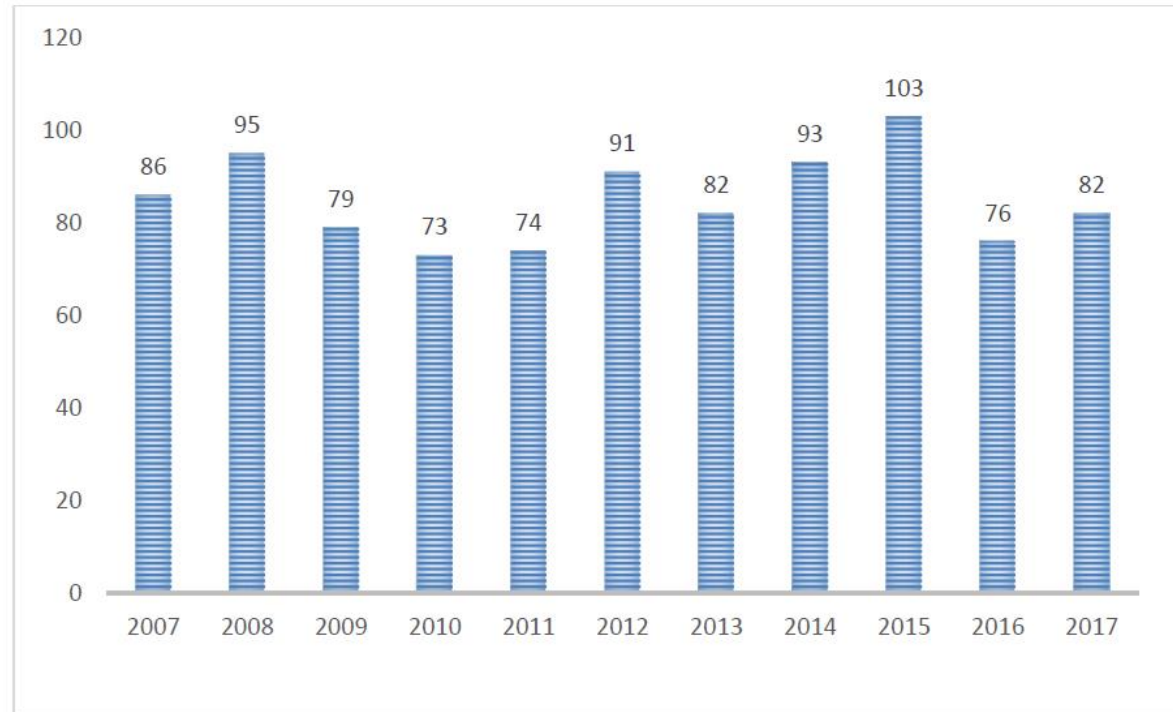
- Significant scaling up in testing numbers:
 - 2009: 28,256
 - 2017: 47,864

Treatment coverage: N Ireland

Current

NI Regional Hepatitis B&C
Managed Clinical Network
Annual Report 2018

FIGURE 7: LABORATORY CONFIRMED HCV PCR POSITIVE CASES, NORTHERN IRELAND, 2007-2017

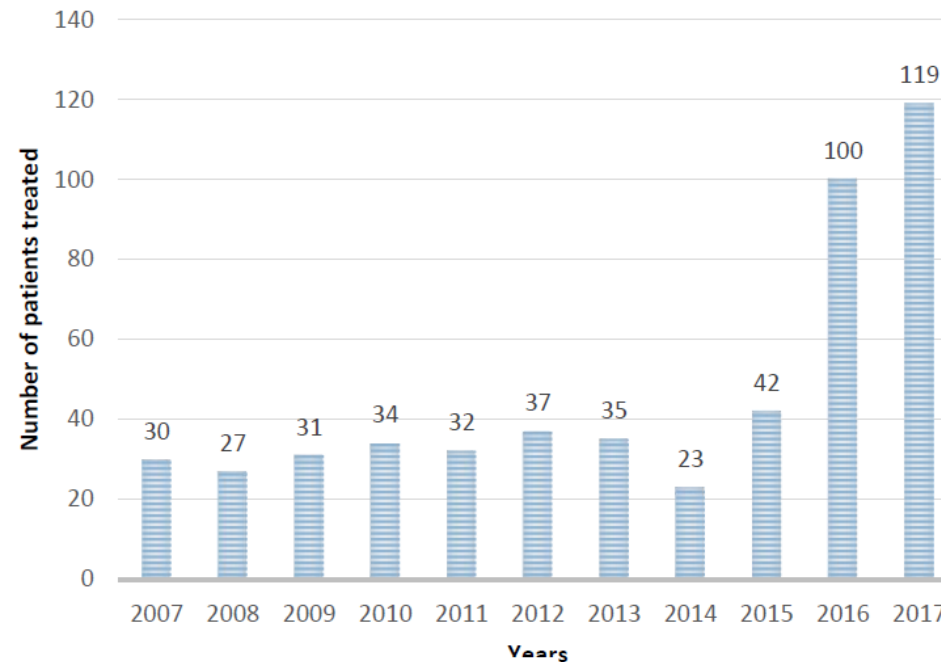


Treatment coverage: N Ireland

Current

NI Regional Hepatitis B&C
Managed Clinical Network
Annual Report 2018

FIGURE 9: HEPATITIS C TREATMENT INITIATIONS IN NORTHERN IRELAND, 2007-2017



- 2016 SVR 97%
- 2017: 2/3 referrals from primary and secondary care

Source: Local RVH database/Regional Hepatology clinic 2018

Treatment coverage: N Ireland

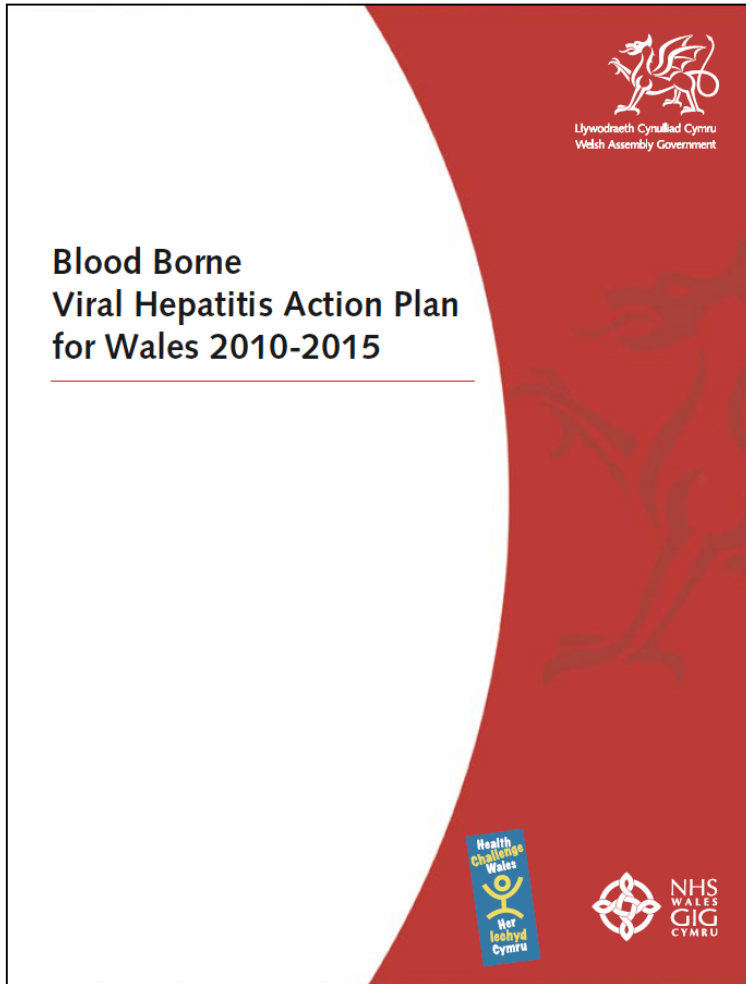
Future

NI Regional Hepatitis B&C
Managed Clinical Network
Annual Report 2018

- 2016/17 increase in cases of HCV among PWID:
 - increase in syringe/needle provision
 - increase in DBT
- Plans to produce an Action Plan towards WHO elimination goals

Treatment coverage: Wales

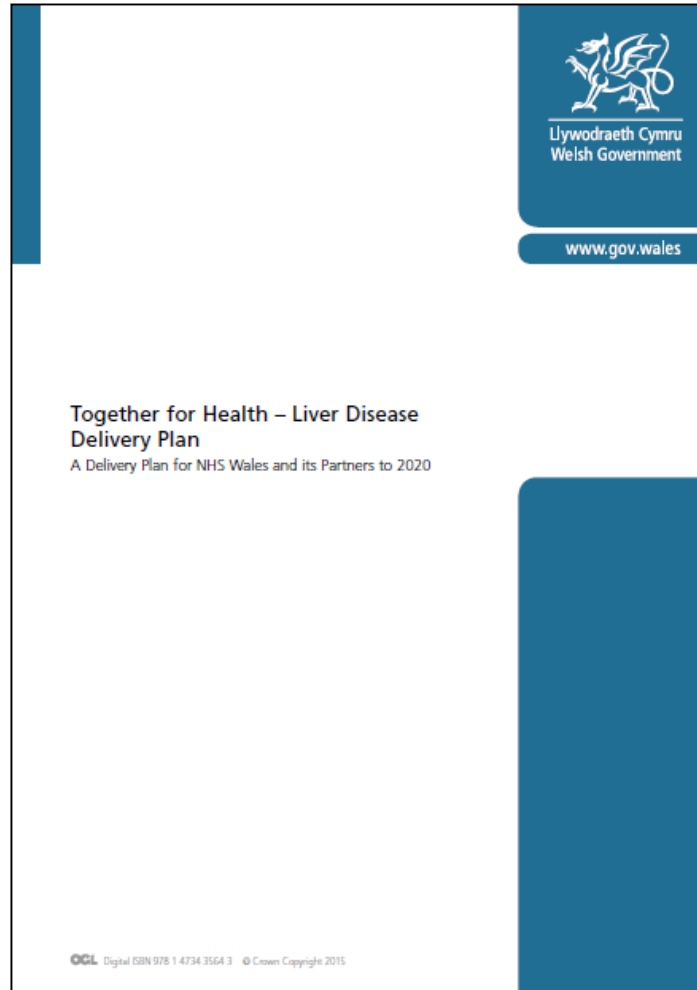
Past



- Action Plan 2010-2015
 - Focus on: prevention, testing and treating
 - DBS drug services & prisons
 - BBV prisons nurse specialist
 - Move to ‘opt-out’ testing in prisons
 - 10% men HCV Ab positive
 - Database developed

Treatment coverage: Wales

Current



- Liver Disease Delivery Plan 2015-2020
 - Roll out of DAAs from 2014
 - No restrictions on numbers
 - Minimum targets set
 - BUT... since 2015 targets have been missed
 - 2017/18 600 treated (target 900)

 - 5,000 diagnosed not yet treated

 - High numbers of undiagnosed

Treatment coverage: Wales

Future



Cynulliad
Cenedlaethol
Cymru

National
Assembly for
Wales

- Wales has signed up to the WHO elimination strategy
- Jan 2019 NAW held inquiry to look at what was needed to achieve elimination
 - If treatment targets met elimination delayed by 1-2 years
 - On current rates elimination could be delayed to 2040.
 - The development of an Elimination Action Plan was advised

HCV: A UK view on treatment coverage



- All 4 countries at risk of not meeting WHO elimination target for 2030
- All facing similar issues:
 - Reported increasing incidence in PWID
 - Significant numbers still undiagnosed
 - ‘Easy’ to engage patients treated LTC problems
 - Treatment numbers need to be sustained/increased

HCV: A UK view on treatment coverage



Public Health
England

Hepatitis C in the UK

2018 report

Working to eliminate hepatitis C as a major public health threat

To achieve WHO goals need to radically change our response to tackling HCV in PWID

- Increase NSP access
- Find and inform large numbers of undiagnosed
- Increase linkage to care & access to treatment:
 - significantly expand community & prison based diagnosis and treatment
 - simplify requirements for treatment.... Pangenotypics / FibroScans
 - Be flexible'one size will not fit all'
- Monitor and ensure equity of access to treatment to all

HCV: A UK view on treatment coverage

Dr Emma E Page

Disclosures:

Clinical Lead for Get Tested LeEDs which is a joint working project between Leeds Teaching Hospitals Trust and Gilead Sciences Ltd.