

1 **Full paper**

2 **Prevalence of hepatitis C virus estimates of undiagnosed individuals in different Italian regions: a**
3 **mathematical modelling approach by route of transmission and fibrosis progression with results**
4 **up to January 2021**

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15 **Short title:** HCV prevalence in Italian regions: an update

16 **SUMMARY**

17 This study provides an update on hepatitis C virus (HCV) estimates across Italy up to January 2021. A
18 mathematical probabilistic modelling approach, including a Markov chain for liver disease progression,
19 was used to estimate current HCV viraemic burden. Prevalence was defined by geographic area using an
20 estimated annual historical HCV incidence by age, treatment, and migration rate from the Italian National
21 database. Viraemic infection was estimated for the main HCV transmission routes by stages F0-F3
22 (patients without liver cirrhosis, i.e., potentially asymptomatic) and F4 (patients with liver cirrhosis, i.e.,
23 potentially symptomatic). By January 2021, we estimated that there were 398,610 individuals in Italy
24 with HCV (prevalence of 0.66%; 95% CI: 0.66-0.67%), of which 287,730 (0.48%; 95% CI: 0.46-0.59%)
25 were stage F0-F3. Prevalence values for all individuals with HCV were: North 0.54% (95% CI: 0.53-
26 0.54%), Central 0.88% (95% CI: 0.87-0.89%), South 0.72% (95% CI: 0.71-0.73%), and the Isles 0.67%
27 (95% CI: 0.66-0.68%). The population at risk for previous/current drug injection accounted for 48.6%
28 of all individuals with HCV. A modelling approach such as this to estimate and update the prevalence of
29 active HCV infection could be a useful methodology for the evaluation of healthcare policies related to
30 HCV elimination plans.

32 *Key words:* HCV, undiagnosed, hepatitis C infection, prevalence, Markov chain

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Ahead of print

36 INTRODUCTION

37 Hepatitis C Virus (HCV) infection is the global leading cause of liver-related morbidity and mortality,
38 and recent estimates report as many as 58 million individuals had chronic HCV in 2020 (Polaris
39 Observatory HCV Collaborators, 2017; *World Health Organization: Hepatitis C* 2021;
40 <https://www.who.int/news-room/fact-sheets/detail/hepatitis-c>). A high proportion of individuals do not
41 show symptoms of the disease for decades, although the rate of disease progression increases with age
42 (Hajarizadeh et al., 2013; Sweeting et al., 2006).

43 Since direct-acting antiviral drugs (DAAs) have become available for the successful treatment of HCV
44 infection (Afdhal et al., 2014; Sulkowski et al., 2014), attention has focussed on identifying infected
45 individuals. Only through the identification of this population will it be possible to achieve 2030 World
46 Health Organization elimination targets (*WHO: Global health sector strategy on viral hepatitis 2016-
47 2021*, pp. 2016–2021; <https://apps.who.int/iris/handle/10665/246177>).

48 Historically, Italy has been considered the country with the highest rate of HCV prevalence in Western
49 Europe (Bellentani et al., 1999; Guadagnino et al., 1997; Kondili et al., 2021a; Maio et al., 2000).
50 However, a prevalence of around 1% (95% uncertainty interval: 0.4-1.4%) has recently been estimated
51 using the “Polaris” model, which is grounded in the natural history of HCV progression and forecasts
52 the HCV burden on the Italian general population (Blach et al., 2022). The reported number of patients
53 treated annually, as tracked in the Italian Medicines Agency (*Agenzia Italiana del Farmaco*; AIFA;
54 <https://www.aifa.gov.it/aggiornamento-epatite-c>) Monitoring Registry for DAAs, was allocated to the
55 age and liver disease stage of the eligible population with HCV by the relative size of the population in
56 each treatment disease stage (Kondili et al., 2018). A free screening program has now been implemented
57 in Italy, with a specific fund approved by law, although decentralized HCV models are still being used
58 without any uniform strategies across different regions (Kondili et al., 2021a, 2020a). The coronavirus
59 disease 2019 (COVID-19) pandemic has increased existing challenges in diagnosis and linkage to care
60 of individuals with HCV (Blach et al., 2021; Kondili et al., 2021c; Mennini et al., 2021). We have
61 recently published national estimates on the number of individuals with HCV up to January 2020 by
62 using a probabilistic approach to estimate the infection rate and a Markov model for liver disease
63 progression in different Italian regions (Kondili et al., 2022a). The present analysis extends these findings
64 further to January 2021, with the goal to update the HCV infection burden by considering the potential
65 role of screening and diagnosis during the pandemic period.

66

67 MATERIALS AND METHODS

68 *Study design*

69 A previously validated mathematical model was used for this study (Kondili et al., 2021b). The model
70 was divided into two distinct computations. It computes the number of infected individuals on a national
71 basis from available literature and the Italian National database (“Statistiche Istat,”
72 <http://dati.istat.it/Index.aspx>) by age, transmission route, and fibrosis stage from 1952 to October 2021.
73 Using the above-described methodology, the model also provides the number of individuals with HCV,
74 subdivided by age, transmission route, and fibrosis status by geographic macroarea from 2002 to January
75 2021 (Kondili et al., 2022a, 2021b).

76 *Study population and literature search*

77 Data on HCV prevalence by route of infection in high risk groups was obtained from a literature search,
78 as previously described elsewhere (Kondili et al., 2021b).

79 *The model*

80 An evolutionary HCV transmission model was developed and implemented using the open-source
81 programming language Python 3.7. The first stage of the model has been previously reported (Kondili et
82 al., 2021b), beginning in 1952 to the end of 2001 calculating the number of infected individuals by age,
83 transmission route, and fibrosis stage. The evolutionary steps considered the insertion of new-borns, new
84 infections (the model accounts for six transmission routes), possible fibrosis evolution (computed using
85 a Markov chain approach), HCV treatment, and liver-related mortality rate. All cause-mortality was also
86 considered (see below). After this stage, the number of individuals with HCV was subdivided for each
87 region considering the distribution of the population and the variability of risk factors among the different
88 regions as explained in the section “Transmission routes and associated risk.” Subsequently, the
89 evolution of the HCV transmission and liver disease progression among the Italian population was
90 performed up to January 2021, with a specific method that included the migration of people (inter-
91 regional and international), referred to as *regional infection burden evaluation*.

92 An overview of these steps is described below.

- 93 1) New-borns: The number of new-borns was added (data were derived from the Italian National
94 database [ISTAT]) (“Statistiche Istat,” <http://dati.istat.it/Index.aspx>).
- 95 2) Internal and external migration (i.e., the movement of people from one Italian region to another or
96 from a foreign country to Italy, respectively). For internal migration, the number of individuals was

97 included independent of age, whereas for external migration a minimum breakdown by age was
98 considered using data derived from ISTAT (“Statistiche Istat,” <http://dati.istat.it/Index.aspx>).

99 3) Individuals with HCV: Individuals recently contracting HCV per region, age, and year were
100 considered (Kondili et al., 2021b). The number of new HCV cases was then calculated following the
101 infection probabilities, depending on age, transmission route (“SEIEVA - Sistema Epidemiologico
102 Integrato dell’Epatite Virale Acuta,” <https://old.iss.it/web/guest/seieva/chi-siamo>), and current year.

103 4) Treated patients: The estimated number of patients treated with anti-viral therapy (i.e., interferon- and
104 DAA-based treatments) was subtracted from the population with HCV based on expert data provided
105 by EpaC onlus Patient Association up to 2014 (<https://www.epac.it/>) and AIFA Registry data for DAA
106 monitoring since 2015 (“Aggiornamento dati Registri AIFA DAAs - Epatite C cronica,”
107 <https://www.aifa.gov.it/aggiornamento-epatite-c>). Estimates from the Italian Platform for the Studies
108 of Viral Hepatitis Therapies (PITER) cohort, for unclassified fibrosis stage by AIFA Registry, were
109 also used (Kondili et al., 2021b, 2017). The average number of individuals successfully treated and
110 subtracted were 2,000 every year from 1993 to 2000 and 7,000 every year from 2001 to 2014. The
111 number of treated patients was based on AIFA Monitoring Registry for DAA
112 (<https://www.aifa.gov.it/aggiornamento-epatite-c>) data as follows: from 2015 to 2017 (mainly
113 prioritized treatment criteria). an average of 36,296 patients each year was subtracted at a 95% rate of
114 sustained virologic response from the number of total infected patients. Following universal treatment
115 since 2017, a total of 56,499 treated patients for 2018, 36,348 treated patients for 2019, and 15,664
116 treated patients for 2020 were subtracted each year at a 98% rate of sustained virologic response from
117 the total of infected patients.

118 5) Fibrosis evolution: Individuals with HCV who were considered to undergo a possible fibrosis
119 progression according to a Markov chain probability evolution process (Kondili et al., 2021b).

120 6) Mortality: The number of annual deaths per region and age provided by ISTAT (“Statistiche Istat,”
121 <http://dati.istat.it/Index.aspx>) was subtracted. In our model, we accounted for HCV-related mortality
122 by considering the average values of transition probabilities following the F4 fibrosis stage reported
123 by Linthicum et al., 2016 (Linthicum et al., 2016) and Kondili et al., 2017 (Dienstag et al., 2011;
124 Kondili et al., 2017; Townsend et al., 2011; Wright et al., 2006). All transition probabilities were
125 adjusted for competing probabilities of death from other causes according to official data
126 [ISTAT](“Statistiche Istat,” <http://dati.istat.it/Index.aspx>).

127 *Transmission routes and associated risk*

128 The model independently tracked each transmission route, distributing the weight of the effect of each
129 route over time, as previously described (Kondili et al., 2021b). Data for the following high-risk routes
130 of HCV transmission were considered: previous or current drug injection, tattoos or body piercing, sexual
131 transmission, glass syringe, blood transfusion, and vertical transmission. Key criteria for input of data in
132 the model regarding age and year of infection were defined for each risk group. Based on HCV
133 prevalence and the time-series of the Italian population, we reconstructed the probability of infection for
134 ages 0-100, for years 1952 to 2021, and for the six different infection routes.

135 The total burden of infection at the beginning of 2002 was obtained using the results of the national phase
136 (Kondili et al., 2021b), divided for each macroregion. The criteria used to distribute the burden of
137 infection across different macroregions are described below for each high-risk route of infection.

138 *a) History of previous or current drug use*

139 The distribution of previous or current drug use was estimated according to the 2002 national report on
140 drug addiction (*Relazione Annuale Al Parlamento Sullo Stato Delle Tossicodipendenze in Italia 2002*,
141 2002; <http://www.edscuola.it/archivio/handicap/tossicodipendenze02.pdf>). Distribution of this risk
142 factor was also based on the number of individuals with HCV in proportion to the number of individuals
143 under treatment in a drug addiction service (SERD; *Servizi per le Tossicodipendenze*) according to
144 fractions. Peak age of infection was 27 years, and HCV infection through injection of drugs was assumed
145 to begin in 1970.

146 *b) History of tattoo or body piercing*

147 Nationally, ages of infection from tattoo or body piercing were between 15 and 70 years, with a peak age
148 of 35 years (Kondili et al., 2021b). However, data were not available to evaluate the prevalence of this
149 route of infection across different regions; therefore, the number of individuals with HCV was subdivided
150 proportionally to avoid errors in estimates. A region yields the number of those with HCV for a given
151 age group proportionally to the number of those of that age in the region.

152 *c) Sexual transmission*

153 Due to the lack of available data for the route of sexual transmission, we applied the same formula as
154 that for the route of tattoo and body piercing (Kondili et al., 2021b). Transmission of HCV in this
155 population occurred from 15 to 65 years of age (peak age of 35 years).

156 *d) Previous glass syringe use*

157 A higher prevalence was assumed to have occurred at an early age (0-8 years) due to glass syringe
158 vaccination since 1950. However, neither infection by glass syringe (in 1975, the single use of disposable

159 plastic syringes became law in Italy, substituting glass syringes nationwide) nor blood transfusion routes
160 were considered in any new HCV cases since 2002; therefore, these routes can be referred to as “extinct”
161 for the regional phase.

162 *e) Risk of infection by blood transfusion in the past*

163 Since blood transfusion was more likely to occur at older ages due to surgical intervention, the peak risk
164 was centred at around 60 years of age. The age distribution profile started when the model began and
165 peaked in the nineties (when the virus was finally isolated), and decreased gradually in subsequent years.

166 *f) Vertical transmission*

167 The risk of vertical transmission was calculated from the number of mothers with HCV; these females
168 transfer the virus to new-borns at a given percentage (values retrieved from ISTAT]). The risk was
169 estimated to be around 5.8% (Hofstraat et al., 2017; “Statistiche Istat,” <http://dati.istat.it/Index.aspx>).
170 Reduction of the risk has been modelled with a linear decrease up to 0.015% (Kondili et al., 2021b).

171 *Markov chain progression for fibrosis evolution*

172 The evolution from one stage to the next was modelled according to a Markov chain approach, previously
173 described elsewhere (Kondili et al., 2021b). Briefly, the following annual transition probabilities were
174 used in the reference case: $F0 \rightarrow F1 = 7.6\%$, $F1 \rightarrow F2 = 9.5\%$, $F2 \rightarrow F3 = 10.8\%$, and $F3 \rightarrow F4 = 13.4\%$
175 (Linthicum et al., 2016). It was assumed that not more than one transition per year could occur per patient,
176 and potential spontaneous liver fibrosis regression was not considered.

177 *Sensitivity analysis*

178 Sensitivity analysis was performed using a Monte Carlo approach to estimate the effect of each input on
179 the number of estimated individuals with HCV and their annual distribution among F0-F4 stages as
180 described previously in detail (Kondili et al., 2022a, 2021b). In brief, we repeatedly ran the model for
181 1,000 simulations, each time randomly picking the value of some input parameters (still around the
182 values), and we assessed the effect of this randomization on the results of the model. Our Monte Carlo
183 scheme intercepts two different types of randomization: risk distribution and transition probabilities. This
184 approach considered the uncertainty in annual distribution among the F0-F4 stages, in the fibrosis annual
185 transition probabilities, and the probability of self-curing (recover spontaneously without treatment). In
186 each simulation, the self-curing and transition probabilities were derived (independently) from their
187 predefined random distributions. We also conducted a sensitivity analysis of our model by transmission
188 route for the different macroareas for individuals with stage F0-F3 and F4 (Kondili et al., 2022a).
189 Sensitivity analysis provides information on the margins for which prevalence estimates are valid, so as

190 to understand the degree of variability in the data. The output of the Monte Carlo consisted of mean
191 values of the different prevalence values computed using our model, together with a 95% confidence
192 interval (CI), from 2.5 to 97.5 percent for each.

193 **RESULTS**

194 *HCV prevalence estimates by Italian macroarea*

195 The number of individuals with HCV in Italy in January 2021 was estimated at 398,610 (prevalence of
196 0.66%; 95% CI: 0.66-0.67%). Higher prevalence estimates were seen in regions of Central Italy (0.88%;
197 95% CI: 0.87-0.89%), such as Umbria (1%) and Marche (1%), followed by the South (0.72%; 95% CI:
198 0.71-0.73%), such as Basilicata (0.83%) and Calabria (0.81%), and the Isles (0.67%; 95% CI: 0.66-
199 0.68%), in regions such as Sicily (0.66%) and Sardinia (0.71%); the lowest values were seen in regions
200 in the North (0.54%; 95% CI: 0.53-0.54%), such as Piedmont (0.67%) and Veneto (0.6%) (Table 1 and
201 Figure 1). The distributions of HCV prevalence in potentially asymptomatic individuals (stage F0-F3,
202 i.e., patients without liver cirrhosis; estimated to be 287,730) and those with F4 stage disease (patients
203 with liver cirrhosis, thus potentially symptomatic; estimated to be 110,880) for the 4 macroareas are
204 summarised in Table 1. While the prevalence values for individuals with stage F0-F3 across the four
205 macroareas were similar, varying from 0.43% to 0.59%, and similar to the overall prevalence for this
206 stage (0.48%; 95% CI: 0.46-0.59%) (Table 1), prevalence values for individuals with stage F4 disease
207 were generally two-fold lower than those with stage F0-F3, apart from the North, where it was
208 approximately four-fold less (0.11% of F4 vs. 0.43% of those with F0-F3 fibrosis stage) (Table 1).

209 *HCV prevalence by high-risk transmission route*

210 Marked differences in overall prevalence estimates across the six high-risk infection routes were
211 observed (Table 2). The highest prevalence was observed by previous or current drug use (0.32%; 95%
212 CI: 0.32-0.33%) followed by previous or current tattoo use (0.17%; 95% CI: 0.16-0.17%). Stratifying by
213 macroarea showed little variation in prevalence estimates for each infection route (Table 3).

214 *Sensitivity analysis*

215 Ranges in variability of estimates by fibrosis stage and age of cohort by sensitivity analysis were obtained
216 as final results of the computations (up to January 2021). Variations around the mean values and CI
217 ranges were computed by 1,000 Monte Carlo simulations. Specifically, the overall percentage variations
218 around prevalence values were 0.42%-0.83%, for variations within 95% confidence intervals. The

219 variations in 95% confidence intervals were 2.7-5.4% for F0-F3 and 7-14% for F4, indicating the
220 robustness of our model's findings.

221 **DISCUSSION**

222 The present study confirms findings from our recent regional analysis of the prevalence of HCV across
223 Italy up to January 2020 (Kondili et al., 2022a, 2021b) and extends them to provide more recent estimates
224 up to January 2021. Compared with our present findings, after 12 months (up to January 2021) and by
225 using the same modelling approach, we have observed a slight decrease of 10,389 in the overall absolute
226 number of individuals with HCV compared with estimates for the previous year (409,183 in January
227 2020 vs. 398,610 in January 2021). While the number of individuals with stage F0-F3 decreased from
228 300,171 to 287,981 (a decrease of 4.1%), the number of those with stage F4 (symptomatic) increased
229 slightly, from 109,012 to 110, 813 (1,800 individuals and an increase of 1.6%).

230 The estimated number (and distribution across Italy) of patients with F4 cirrhosis is concerning and was
231 not expected, considering their potential symptomatic disease and the high importance of viral
232 eradication in these patients, who have been prioritized for treatment since 2015 when DAAs became
233 available. The lack of early diagnosis in people with severe liver damage has also been documented by
234 the AIFA DAA monitoring registry, which reported 20% of patients treated having a diagnosis of liver
235 cirrhosis from 2019 to date ("Aggiornamento dati Registri AIFA DAAs - Epatite C cronica,"
236 <https://www.aifa.gov.it/aggiornamento-epatite-c>), patients that would have been diagnosed from 2015 in
237 Italy. In addition, many people with viral hepatitis are still diagnosed too late (Lazarus et al., 2019), often
238 resulting in increased risk of severe liver complications and death (Picchio et al., 2021). Furthermore,
239 several studies conducted in the opportunistic screening setting still suggest a high prevalence of
240 individuals with active infection in severe stages of liver disease who have not yet been treated (Piazzolla
241 et al., 2021). These data imply that intervention focusing on case finding needs to be improved to ensure
242 earlier diagnosis of populations at risk of disease progression. In terms of disease control, it is also
243 important to increase our understanding of medical barriers that limit access to healthcare in different
244 settings, starting from the general practitioner's awareness and clinical behaviour to patients with high
245 risk of HCV infection and signs of chronic severe liver damage. Our findings indicate an estimated
246 110,880 individuals in need of urgent diagnosis, immediate linkage to care, and treatment to eradicate
247 HCV infection to stop liver disease progression.

248 Considering the different factors for high risk of infection, people 46-55 years of age who previously
249 injected or currently inject drugs and those who had undergone aesthetic procedures such as tattoos or

250 body piercing were identified as the main populations in which higher prevalence of undiagnosed
251 asymptomatic individuals (stage F0-F3 disease) is expected, with similar distribution across regions
252 (Kondili et al., 2021b). In these populations of potentially asymptomatic individuals, active HCV
253 screening should be implemented across Italy to identify the submerged population of asymptomatic
254 people with chronic HCV infection, as the required step toward achieving HCV elimination. This focused
255 screening approach, if appropriately undertaken, would identify a high rate of individuals with HCV, due
256 to the higher rate of infection related to risk factors in the younger population (drug use and aesthetic
257 procedures at risk), and simultaneously reduce onward viral transmission by clearing HCV from those
258 who are also at higher risk of spreading the infection (Kondili et al., 2022b).

259 Although a higher estimated HCV prevalence of undiagnosed individuals was seen in some specific
260 regions in Central Italy and in the Basilicata and Calabria regions in the South, similar but slightly lower
261 prevalence estimates of undiagnosed individuals were generally observed across the four macroareas.
262 This would suggest a decrease in the overall level of prevalence in HCV infection compared with the
263 past in Italy, where higher prevalence values and a gradient from North to South (3.9% in Veneto to
264 16.2% in Campania) was related mainly to the nosocomial transmission of infection (A.I.S.F,
265 ASSOCIAZIONE ITALIANA PER LO STUDIO DEL FEGATO, [https://www.webaisf.org/per-il-](https://www.webaisf.org/per-il-paziente/le-malattie-del-fegato/)
266 [paziente/le-malattie-del-fegato/](https://www.webaisf.org/per-il-paziente/le-malattie-del-fegato/); Guadagnino et al., 1997). Despite using a different approach compared
267 with the modelling used to estimate the global HCV burden, the results of this modelling study report an
268 estimation within the range reported for overall active infection in Italy up to 2020. Specifically, an
269 overall prevalence of 1% has been estimated up to 2020, with ranges varying from 0.4-1.4%, by the
270 Polaris Observatory (Blach et al., 2022), whereas this differential modelling of the HCV infection burden
271 by transmission route granularity in Italy up to January 2021 shows a prevalence of active infection of
272 0.66% (0.66-0.67). These two independent estimates are very similar, confirming the robustness of this
273 modelling approach, and suggest that Italy is currently not a country with high HCV endemic levels.
274 However, in terms of achieving the elimination target, screening for active HCV infection is the only
275 currently available tool for reaching asymptomatic individuals and ensuring their linkage to care and
276 viral eradication. In this regard, despite these relatively low prevalence values compared with past values,
277 the screening tools for the birth 1948-1988 cohort have been shown to be highly cost-effective in Italy
278 (Kondili et al., 2020a).

279 Analyses for the present study were performed up to January 2021, therefore including data up to 1 year
280 after the severe acute respiratory syndrome coronavirus 2 virus pandemic (WHO,
281 <https://covid19.who.int/table>). It is well documented that healthcare facilities across all Italian regions

282 had to be re-organized in order to cope with the increased number of patients with COVID-19 admitted
283 to emergency departments, resulting in the postponement of medical services and procedures considered
284 “non-essential” or “deferrable.” The reduction in the treatment rate seen in 2019 and the nearly complete
285 interruption of DAA treatment during the COVID-19 pandemic are likely to impact directly on the
286 management of HCV and are of serious concern (Blach et al., 2021). A recent nationwide survey
287 undertaken by the Italian Association for the Study of the Liver highlighted a significant decrease and
288 suspension in the number of outpatient visits and prescriptions of antiviral treatment, which partially
289 recovered 1 year later. However, the increase in outpatient visits was not followed by an increase in DAA
290 prescriptions (“Aggiornamento dati Registri AIFA DAAs - Epatite C cronica,”
291 <https://www.aifa.gov.it/aggiornamento-epatite-c>; Aghemo et al., 2020; Mennini et al., 2021; Ponziani et
292 al., 2021). The potential impact of this deferral has been specifically assessed in a separate analysis,
293 where it was estimated that deferring DAA treatment for a further 6 months could increase the number
294 of deaths from liver-related disease in HCV patients in Italy to more than 500 patients after 5 years
295 (Kondili et al., 2020b). Regions that have experienced a high number of cases of patients hospitalized
296 for COVID-19, such as Lombardy, may be more susceptible to deferral in DAA treatment (Buoro et al.,
297 2020).

298 *Limitations*

299 The impact of other less-frequent transmission routes was not considered in the present analysis, and this
300 limitation may have resulted in an underestimation of the observed prevalence values. Non-liver related
301 mortality (i.e., natural mortality due to other causes) was based on ISTAT data (Italian Mortality
302 Registry), which does not specify the cause of death and may have led to a potential overestimation of
303 the population living with HCV infection (i.e., F4 stage disease). Another limitation is the high number
304 of unregistered immigrants (potentially asymptomatic and undiagnosed for HCV) (Coppola et al., 2019;
305 Fedeli et al., 2019; Massimo et al., 2018) in Italy who were not considered in our estimations. Further
306 studies are needed to explore the impact of the immigrant population on HCV prevalence.

307 **CONCLUSION**

308 Findings from the present analysis showed that the number of individuals with HCV in Italy was
309 estimated at 398,610 in January 2021 (prevalence of 0.66%; 95% CI: 0.66-0.67%). Based on data derived
310 from this modelling study, it is necessary to stress the importance of promoting attentive case finding
311 and linkage to care in individuals with symptomatic liver disease (estimated to be 110,880) in addition

312 to the active screening of asymptomatic people (estimated to be 287,730) in order to reduce prevalence,
313 limit incidence, and potentially achieve the elimination of HCV in Italy.

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321 **Conflicts of interest**

322 LAK received teaching grants from AbbVie and Gilead. MA received funding for membership on
323 Advisory Boards, for the preparation of educational materials, for research and educational grants, for
324 membership on speaker panels and for support for travel to conferences from the following companies:
325 Gilead Sciences, Janssen-Cilag, ViiV Healthcare, Merck Sharp and Dohme, Abbvie, Angelini, Pfizer,
326 GSK, Menarini. AA serves on the advisory boards for AbbVie, Gilead, MSD, Intercept, Sobi, and Mylan
327 and has received speaker fees from AbbVie, Gilead, Sobi, and Alfasigma. CMM received an
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329 and VG are AbbVie employees and may own AbbVie stocks and options.

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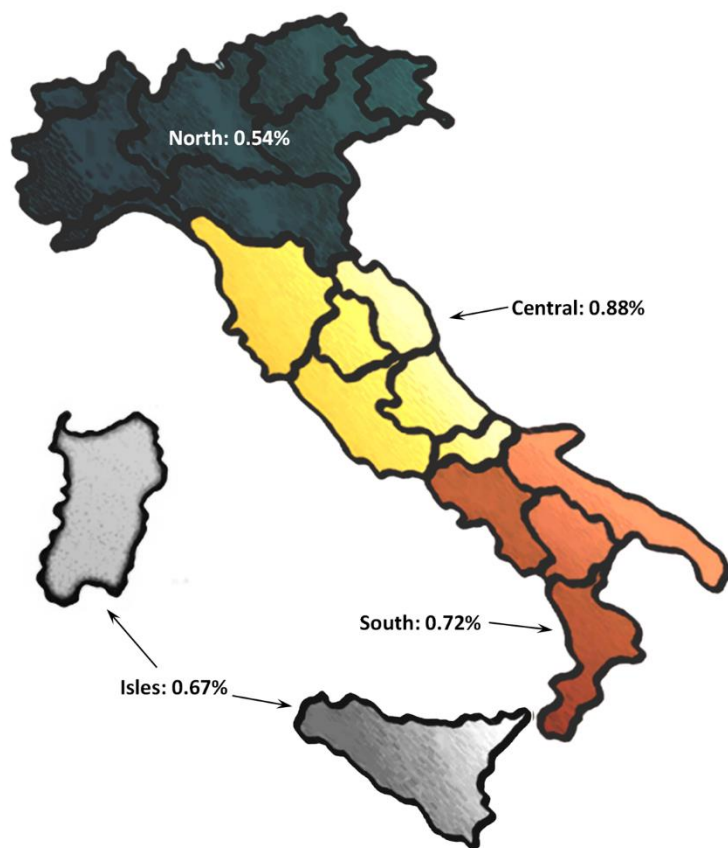
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464 **Figure 1.** Map showing the estimated prevalence of hepatitis C virus in the 4 macroareas of Italy up to
465 January 2021.



466
467

468 **Table 1** - Estimates of the absolute number and percentage of viraemic HCV individuals in Italy
 469 according to fibrosis stage and macroarea up to January 2021.

Macroarea/ fibrosis stage	Absolute number and 95% CI	Prevalence (%) and 95% CI (%)	Percentage
North			
F0-F3	121,533 (118,036-125,030)	0.43 (0.42-0.45)	42.2
F4	29,450 (26,073-32,827)	0.11 (0.09-0.12)	26.6
Total	150,983 (149,557-152,410)	0.54 (0.53-0.54)	37.9
Central			
F0-F3	70,028 (67,839-72,217)	0.59 (0.57-0.60)	24.3
F4	35,323 (33,388-37,257)	0.30 (0.28-0.31)	31.9
Total	105,351 (104,188-106,513)	0.88 (0.87-0.89)	26.4
South			
F0-F3	65,506 (63,448-67,564)	0.48 (0.46-0.49)	22.8
F4	33,422 (31,325-35,519)	0.24 (0.23-0.26)	30.1
Total	98,928 (87,623-100,233)	0.72 (0.71-0.73)	24.8
Isles			
F0-F3	30,663 (29,461-31,864)	0.47 (0.46-0.49)	10.7
F4	12,686 (11,473-13,898)	0.20 (0.18-0.21)	11.4
Total	43,348 (42,513-44,183)	0.67 (0.66-0.68)	10.9
Italy			
F0-F3	287,730 (279,911-295,549)	0.48 (0.46-0.59)	100
F4	110,880 (103,130-118,630)	0.18 (0.17-0.20)	100
Total	398,610 (396,960-400,260)	0.66 (0.66-0.67)	100

470 CI = confidence interval; HCV = hepatitis C virus.

471 Fibrosis stages: F0-F3 = asymptomatic; undiagnosed/unlinked to care and F4 = symptomatic;
 472 potentially linked to care and cure. Prevalence estimates were calculated according to each region's
 473 population by age group annually from 1952 up to January 1, 2021, from the Italian National Database
 474 (ISTAT).

475

476 **Table 2 - Estimates of the absolute number and percentage of viraemic HCV individuals in Italy**
 477 *according to fibrosis stage and risk factor for route of transmission up to January 2021.*

Transmission route/ fibrosis stage	Absolute number and 95% CI	Percent of attributable risk and 95% CI (%)	Percentage
Glass syringe			
F0-F3	9,516 (8,617-10,415)	0.016 (0.14-0.17)	3.3
F4	37,159 (35,511-38,808)	0.06 (0.06-0.06)	33.5
Total	46,675 (45,552- 47,799)	0.08 (0.08-0.08)	11.7
Transfusion			
F0-F3	4,037 (3,532-4,541)	0.007 (0.006-0.008)	1.4
F4	5,914 (5,347-6,482)	0.01 (0.009-0.011)	5.3
Total	9,951 (9,374-10,528)	0.02 (0.02-0.02)	2.5
PWID*			
F0-F3	144,307 (139,845-148,768)	0.24 (0.23-0.25)	50.2
F4	49,404 (45,371-53,437)	0.082 (0.075-0.089)	44.6
Total	193,710 (192,186-195,235)	0.32 (0.32-0.33)	48.6
Sex			
F0-F3	42,141 (41,117-43,165)	0.07 (0.068-0.072)	14.6
F4	3,496 (2,915-4,078)	0.006 (0.005-0.007)	3.2
Total	45,637 (44,837-46,438)	0.08 (0.07-0.08)	11.4
Tattoo			
F0-F3	86,491 (84,161-88,822)	0.14 (0.14-0.15)	30.1
F4	13,681 (11,895-15,466)	0.023 (0.02-0.026)	12.3
Total	100,172 (98,922-101,423)	0.17 (0.16-0.17)	25.1
Vertical			
F0-F3	1,237 (1,034-1,440)	0.002 (0.002-0.002)	0.4
F4	1,227 (997-1,458)	0.002 (0.002-0.002)	1.1
Total	2,464 (2,233-2,695)	0.004 (0.004-0.005)	0.6
All routes			
F0-F3	287,730 (279,911-295,549)	0.48 (0.46-0.59)	100

F4	110,880 (103,130-118,630)	0.18 (0.17-0.20)	100
Total	398,610 (396,960-400,260)	0.66 (0.66-0.67)	100

478 CI = confidence interval; HCV = hepatitis C virus; PWID = people who inject drugs.
479 Fibrosis stages: F0-F3 = asymptomatic; undiagnosed/unlinked to care and F4 = symptomatic;
480 potentially linked to care and cure. *Refers to previous or current risk of infection by drug use.
481

Ahead of print

482 **Table 3** - Estimates of the absolute number and percentage of viraemic HCV individuals in Italy
 483 according to fibrosis stage, risk factor for route of transmission, and macroarea up to January 2021.

Region	Transmission route	Fibrosis stage	Absolute number			Percent of attributable risk		
			Mean number	Lower 95% CI	Upper 95% CI	Mean (%)	Lower 95% CI	Upper 95% CI
North	Glass syringe	F0-F3	2,069	1724	2414	0.007	0.006	0.009
		F4	5,772	5051	6494	0.02	0.02	0.02
		Total	7,841	7176	8507	0.03	0.03	0.03
	Transfusion	F0-F3	936	742	1130	0.003	0.003	0.004
		F4	962	762	1163	0.003	0.003	0.004
		Total	1,898	1622	2174	0.007	0.006	0.008
	PWID	F0-F3	61,834	59693	63975	0.22	0.21	0.23
		F4	16,244	14395	18094	0.06	0.05	0.06
		Total	78,078	76951	79206	0.28	0.27	0.28
	Sex	F0-F3	18,567	17915	19218	0.07	0.06	0.07
		F4	1,278	943	1613	0.005	0.003	0.006
		Total	19,844	19255	20434	0.07	0.07	0.07
	Tattoo	F0-F3	37,600	36356	38843	0.13	0.13	0.14
		F4	4,798	3962	5634	0.02	0.01	0.02
		Total	42,397	41520	43275	0.15	0.15	0.15
	Vertical	F0-F3	529	368	689	0.002	0.001	0.003
		F4	396	256	535	0.001	0.001	0.002
		Total	924	733	1115	0.003	0.003	0.004
Central	Glass syringe	F0-F3	2,980	2593	3,368	0.03	0.02	0.03
		F4	12,494	11,932	13,055	0.11	0.10	0.11
		Total	15,474	14,962	15,986	0.13	0.13	0.13
	Transfusion	F0-F3	1,201	958	1,443	0.010	0.008	0.01
		F4	1,973	1,693	2,254	0.02	0.01	0.02
		Total	3,174	2,860	3,488	0.03	0.02	0.03
	PWID	F0-F3	35,458	34,145	36,770	0.30	0.29	0.31
		F4	15,502	14,393	16,611	0.13	0.12	0.14

		Total	50,959	50,208	51,711	0.43	0.42	0.43
	Sex	F0-F3	9,647	9,258	10,036	0.08	0.08	0.08
		F4	974	728	1,220	0.01	0.01	0.01
		Total	10,621	10,265	10,976	0.09	0.09	0.09
	Tattoo	F0-F3	20,465	19,697	21,232	0.17	0.17	0.18
		F4	4,002	3,406	4,598	0.03	0.03	0.04
		Total	24,467	23,911	25,023	0.21	0.20	0.21
	Vertical	F0-F3	278	186	370	0.002	0.002	0.003
		F4	377	286	469	0.003	0.002	0.004
		Total	655	559	752	0.01	0.00	0.01
South	Glass syringe	F0-F3	3,220	2,828	3,611	0.02	0.02	0.03
		F4	14,048	13,304	14,792	0.10	0.10	0.11
		Total	17,268	16,601	17,934	0.13	0.12	0.13
	Transfusion	F0-F3	1,353	1,078	1,629	0.010	0.008	0.01
		F4	2,175	1,854	2,496	0.02	0.01	0.02
		Total	3,528	3,150	3,907	0.03	0.02	0.03
	PWID	F0-F3	32,384	31,077	33,691	0.24	0.23	0.25
		F4	12,772	11,689	13,854	0.09	0.09	0.10
		Total	45,156	44,183	46,128	0.33	0.32	0.34
	Sex	F0-F3	9,266	8,824	9,707	0.07	0.06	0.07
		F4	837	633	1,040	0.006	0.005	0.008
		Total	10,102	9,682	10,522	0.07	0.07	0.08
	Tattoo	F0-F3	19,005	18,363	19,646	0.14	0.13	0.14
		F4	3,287	2,767	3,807	0.02	0.02	0.03
		Total	22,292	21,659	22,924	0.16	0.16	0.17
	Vertical	F0-F3	279	172	385	0.002	0.001	0.003
		F4	304	200	407	0.002	0.002	0.003
		Total	582	456	709	0.004	0.003	0.005
Isles	Glass syringe	F0-F3	1,247	979	1,515	0.02	0.02	0.02
		F4	4,845	4,415	5,275	0.08	0.07	0.08
		Total	6,093	5,653	6,532	0.09	0.09	0.10

Transfusion	F0-F3	547	388	706	0.009	0.006	0.01
	F4	804	602	1,006	0.013	0.009	0.02
	Total	1,351	1,132	1,569	0.02	0.02	0.02
PWID	F0-F3	14,631	13,896	15,367	0.23	0.22	0.24
	F4	4,886	4,236	5,536	0.08	0.07	0.09
	Total	19,517	18,850	20,184	0.30	0.29	0.31
Sex	F0-F3	4,662	4,327	4,998	0.07	0.07	0.08
	F4	408	250	565	0.01	0.004	0.01
	Total	5,070	4,738	5,402	0.08	0.07	0.08
Tattoo	F0-F3	9,422	8,822	10,023	0.15	0.14	0.16
	F4	1,593	1,289	1,898	0.02	0.02	0.03
	Total	11,016	10,494	11,538	0.17	0.16	0.18
Vertical	F0-F3	153	82	223	0.002	0.001	0.004
	F4	150	74	225	0.002	0.001	0.004
	Total	302	214	391	0.005	0.003	0.006

484 CI = confidence interval; HCV = hepatitis C virus; PWID = (past or current) people who inject drugs.

485 Fibrosis stages: F0-F3 = asymptomatic; undiagnosed/unlinked to care and F4 = symptomatic;

486 potentially linked to care and cure.

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