

Inferring the effective start dates of non-pharmaceutical interventions during COVID-19 outbreaks

Ilia Kohanovski¹, Uri Obolski^{2,3}, and Yoav Ram^{1,4,a}

¹School of Computer Science, Interdisciplinary Center Herzliya, Herzliya 4610101, Israel

²School of Public Health, Tel Aviv University, Tel Aviv 6997801, Israel

³Porter School of the Environment and Earth Sciences, Tel Aviv University, Tel Aviv 6997801, Israel

⁴School of Zoology, Faculty of Life Sciences, Tel Aviv University, Tel Aviv 6997801, Israel

^aCorresponding author: yoav@yoavram.com, ORCID 0000-0002-9653-4458

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Abstract

During Feb-Apr 2020, many countries implemented non-pharmaceutical interventions, such as school closures and lockdowns, with variable schedules, to control the COVID-19 pandemic caused by the SARS-CoV-2 virus. Overall, these interventions seem to have reduced the spread of the pandemic. We hypothesise that the official and effective start date of such interventions can be noticeably different, for example due to slow adoption by the population, or because the authorities and the public are unprepared. We fit an SEIR model to case data from 12 regions to infer the effective start dates of interventions and contrast them with the official dates. We find mostly late, but also early effects of interventions. For example, Italy implemented a nationwide lockdown on Mar 11, but we infer the effective date on Mar 17 (± 2.99 days 95% CI). In contrast, Germany announced a lockdown on Mar 22, but we infer an effective start date on Mar 19 (± 1.05 days 95% CI). We demonstrate that differences between the official and effective start of NPIS can lead to under-estimating their impact, and discuss potential causes and consequences of our results.

Introduction

26 The COVID-19 pandemic has resulted in implementation of extreme non-pharmaceutical interventions
28 (NPIs) in many affected countries. These interventions, from social distancing to lockdowns, are
30 applied in a rapid and widespread fashion. NPIs are designed and assessed using epidemiological
32 models, which follow the dynamics of infection to forecast the effect of different mitigation and
suppression strategies on the levels of infection, hospitalization, and fatality. These epidemiological
models usually assume that the effect of NPIs on infection dynamics begins at the officially declared
date^{9,11,17}.

34 Adoption of public-health recommendations is often critical for effective response to infectious dis-
eases, and has been studied in the context of HIV¹⁵ and vaccination^{6,22}, for example. However,
36 behavioural and social change does not occur immediately, but rather requires time to diffuse in the
population through media, social networks, and social interactions. Moreover, compliance to NPIs
38 may differ between different interventions and between people with different backgrounds. For ex-
ample, in a survey of 2,108 adults in the UK during Mar 2020, Atchison et al.³ found that those over
70 years old were more likely to adopt social distancing than young adults (18-34 years old), and that
40 those with lower income were less likely to be able to work from home and to self-isolate. Similarly,
42 compliance to NPIs may be impacted by personal experiences. Smith et al.¹⁹ have surveyed 6,149
UK adults in late Apr 2020 and found that people who believe they have already had COVID-19 are
more likely to think they are immune, and less likely to comply with social distancing guidelines.
44 Compliance may also depend on risk perception as perceived by the the number of domestic cases or
even by reported cases in other regions and countries. Interestingly, the perceived risk of COVID-19
46 infection has likely caused a reduction in the number of influenza-like illness cases in the US starting
from mid-Feb 2020²³.

48 Here, we hypothesise that there is a significant difference between the official start of NPIs and their
50 effective adoption by the public and therefore their effect on infection dynamics. We use a *Susceptible-*
Exposed-Infected-Recovered (SEIR) model and a *Markov Chain Monte Carlo* (MCMC) parameter
52 estimation framework to infer the effective start date of NPIs from publicly available COVID-19 case
data in 12 geographical regions. We compare these estimates to the official dates, and find that they

Country	First	Last
Austria	Mar 10 2020	Mar 16 2020
Belgium	Mar 12 2020	Mar 18 2020
Denmark	Mar 12 2020	Mar 18 2020
France	Mar 13 2020	Mar 17 2020
Germany	Mar 12 2020	Mar 22 2020
Italy	Mar 5 2020	Mar 11 2020
Norway	Mar 12 2020	Mar 24 2020
Spain	Mar 9 2020	Mar 14 2020
Sweden	Mar 12 2020	Mar 18 2020
Switzerland	Mar 13 2020	Mar 20 2020
United Kingdom	Mar 16 2020	Mar 24 2020
Wuhan	Jan 23 2020	Jan 23 2020

Table 1: Official start of non-pharmaceutical interventions. The date of the first intervention is for a ban of public events, or encouragement of social distancing, or for school closures. In all countries except Sweden, the date of the last intervention (τ^*) is for a lockdown. In Sweden, where a lockdown was not ordered during the studied dates, the last date is for school closures. Dates for European countries from Flaxman et al.⁹, date for Wuhan, China from Pei and Shaman¹⁸. See Figure S1 for a visual presentation.

include both late and early effects of NPIs on infection dynamics. We conclude by demonstrating
54 how differences between the official and effective start of NPIs can confound assessments of their
impacts.

56 Results

Several studies have described the effects of non-pharmaceutical interventions in different geographical
58 regions^{9,11,17}. Some of these studies have assumed that the parameters of the epidemiological model
change at a specific date (Eq. 6), and set the change date τ to the official NPI date τ^* , usually the
60 lockdown start date (Table 1). They then fit the model once for time $t < \tau^*$ and once for time $t \geq \tau^*$.
For example, Li et al.¹⁷ estimate the infection dynamics in China before and after τ^* , which is set at
62 Jan 23, 2020. Thereby, they effectively estimate the transmission and reporting rates before and after
 τ^* separately.

64 Here, we estimate the joint posterior distribution of the effective start date of NPIs, τ , and the
transmission and reporting rates before and after τ from the entire data, rather than splitting the data
66 at τ . We then estimate $\hat{\tau}$ as the median of the marginal posterior distribution of τ . This is done under
the simplifying assumption that all interventions start at a specific date, despite the reality that the
68 durations between the first and last NPIs were between 4 and 12 days (Table 1, Figure S1).

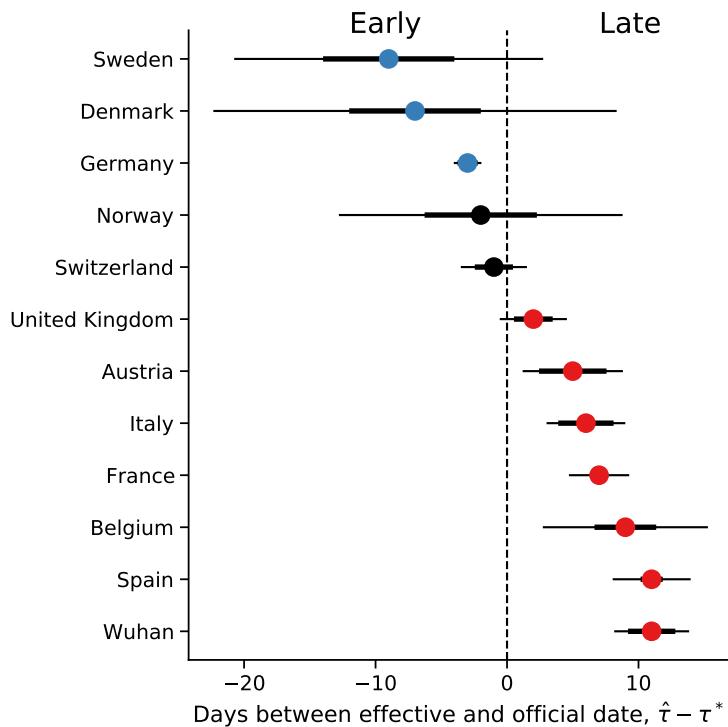


Figure 1: Official vs. effective start of non-pharmaceutical interventions. The difference between τ the effective and τ^* the official start of NPIs is shown for different regions. The effective date is delayed in UK, Austria, Italy, France, Belgium, Spain, and Wuhan, China, compared to the official date (red markers). In contrast, the estimated effective dates in Sweden, Denmark, and Germany are earlier than the official dates (blue markers), although uncertainty is low only for Germany (i.e., zero is not included in 95% CI). The credible intervals for Sweden, Denmark, and Norway are especially wide, see text and Figure 3 for possible explanation. Here, the markers show $\hat{\tau}$, the marginal posterior median, see Table 2. τ^* is the last NPI date (a lockdown except in Sweden, see Table 1). Thin and bold lines show 95% and 75% credible intervals (HDI), respectively. Figure S2 shows a similar summary when estimating $\hat{\tau}$ using case data up to Mar 28, 2020, rather than Apr 11.

We compare the posterior predictive plots of a model with a free τ with those of a model with τ fixed
70 at τ^* and a model without τ (i.e. transmission and reporting rates are constant). All three models
72 were fitted to case data up to Apr 11, used to predict out-of-sample case data up to Apr 24, and these
74 predictions were then compared to the real case data. The model with free τ clearly produces better
76 and less variable predictions (Figure S4a): In all 11 of the European countries, the expected posterior
78 RMSE (root mean squared error) of the out-of-sample predictions is lowest for the model with a free τ
80 (Table S2). When we compare the models using WAIC (Eq. 10), the model with a free τ is strongly
82 preferred in 9 out of 12 countries, the exceptions being Norway (only slight preference), Sweden, and
84 Denmark (Table S1). Notably, the data for Sweden and Denmark do not have a single "peak" during
86 the evaluated dates, possibly leading to wide 95% credible intervals on $\hat{\tau}$ (Figure 1) and poor WAIC in
the model with free τ , whereas the duration between the first and last interventions was especially
long in Norway (Table 1).

We compare the official τ^* and effective τ start of NPIs and find that in most regions the effective
82 start of NPIs differs from the official date: in 10 of 12 countries the 75% credible interval on τ does
not include τ^* (7 of 12 countries when considering a 95% interval; Figure 1). The exceptions are
84 Switzerland and Norway (see below). The latter also has a wide 95% credible interval, perhaps because
it has the longest duration between the first and last NPIs (Table 1). Reported credible intervals, or
86 CI, are calculated as the highest density intervals, or HDI¹⁶.

Late effective start of NPIs. In half of the examined regions, we estimate that the effective start of
88 NPIs τ is later than the official date τ^* .

In Italy, the first case was officially confirmed on Feb 21. School closures were implemented on
90 Mar 5⁹, a lockdown was declared in Northern Italy on Mar 8, with social distancing implemented in
the rest of the country, and the lockdown was extended to the entire nation on Mar 11¹¹. That is, the
92 first and last official NPI dates are Mar 5 and Mar 11. However, we estimate the effective date $\hat{\tau}$ six
days after the lockdown, at Mar 17 (± 2.99 days 95% CI; Figure 2).

94 In Wuhan, China, a lockdown was ordered on Jan 23¹⁷, but we estimate the effective start of NPIs to
be more than a week later, at Feb 3 (± 2.85 days 95% CI; Figure 2).

96 In Spain, social distancing was encouraged starting on Mar 8⁹, but mass gatherings still occurred on
Mar 8, including a march of 120,000 people for the [International Women's Day](#), and a football match
98 between [Real Betis](#) and [Real Madrid](#) (final score 2–1) with a crowd of 50,965 in Seville. A national
lockdown was only announced on Mar 14⁹. Nevertheless, we estimate the effective start of NPI $\hat{\tau}$ ten
100 days later, at Mar 24 (± 2.96 days 95 %CI), rather than Mar 14 (Figure 2).

Similarly, in France we estimate the effective start of NPIs $\hat{\tau}$ at Mar 24 (± 2.29 days 95% CI, Figure 2).
102 This is a week later than the official lockdown, which started at Mar 17, and more than 10 days after
the earliest NPI, banning of public events, which started on Mar 13⁹.

104 **Early effective start of NPIs.** In some regions we estimate an effective start of NPIs τ that is *earlier*
than the official date τ^* (Figure 1). The only conclusive early case is Germany, in which we estimate
106 the effective start of NPIs at Mar 19 (± 1.05 days 95 %CI; Figure 3). This estimate falls between the
first and last official NPI dates, Mar 12 and Mar 22 (Table 1). Therefore, when we refer to this case as
108 "early", we mean that the effective date (Mar 19) occurs *before* the official lockdown date (Mar 22),
not that it occurs before all the NPIs. Interestingly, Germany has the second longest duration between
110 first and last NPIs after Norway (10 and 12 days respectively; see Table 1). However, the credible
interval for the effective date in Germany is narrow (± 1.05 days 95 %CI), whereas it is quite wide in
112 Norway (± 10.79 days; Figure 3). That τ is significantly earlier than τ^* can be suggest that early NPIs

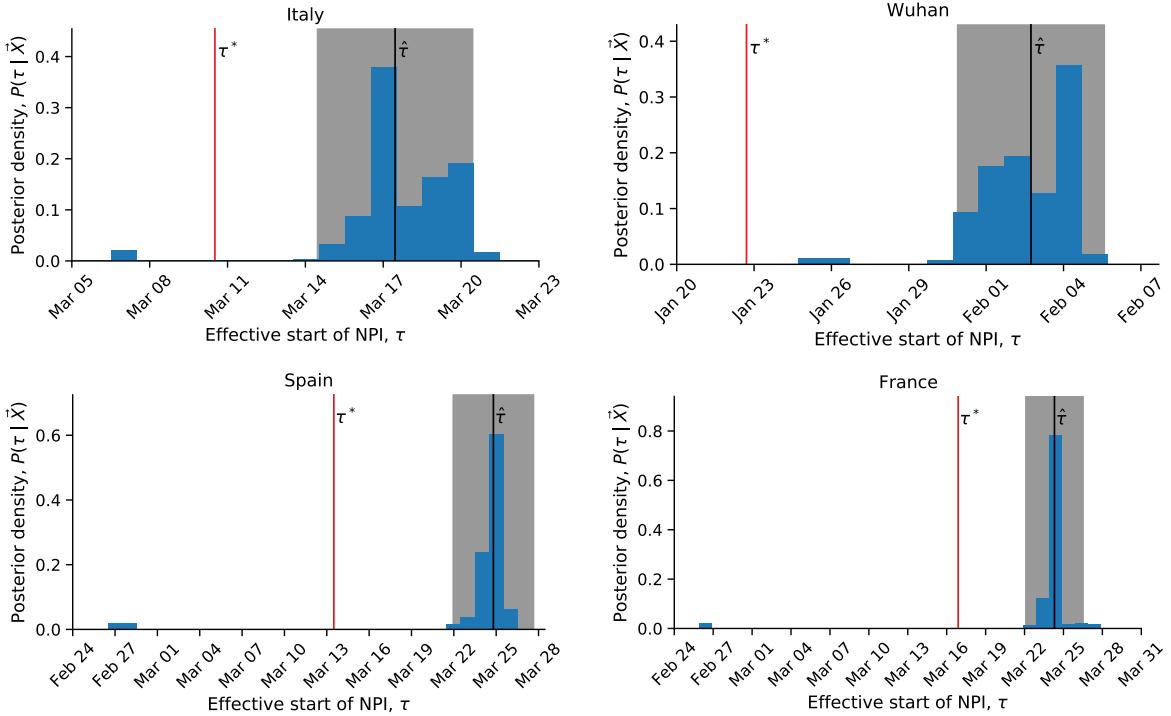


Figure 2: Late effect of non-pharmaceutical interventions. Posterior distribution of τ , the effective start date of NPI, is shown as a histogram of MCMC samples. Red line shows the official last NPI date τ^* . Black line shows the estimate $\hat{\tau}$. Shaded area shows a 95% credible interval.

had been more effective in reducing the transmission rates in Germany compared to other countries.

114 Another possible interpretation is that the German population anticipated the lockdown and began to act in accordance before it started.

116 We also estimate an early effective start of NPIs in Denmark, Norway, and Sweden. However, the
118 credible intervals are quite wide (Figure 1), and in Denmark and Sweden the evidence did not support
the model with free τ over the model with τ fixed at the official date (WAIC values in Table S1).
Indeed, Denmark and Sweden had the highest estimates of λ , that is, the smallest effect of NPIs
120 on transmission rates was the estimated, which probably hinders our ability to estimate τ in these
countries. Moreover, in Sweden, the data do not "peak", i.e., the number of daily cases continues to
122 grow up to Apr 11 (Figure S4a). In Denmark, the opposite occurs: there are seemingly two "peaks",
on Mar 11 and on Apr 8 (Figure S4a); the first "peak" may be a result of stochastic events, for example
124 due to a large cluster of cases or an accumulation of tests. We suspect that these missing and additional
"peaks" increase the uncertainty in our inference.

126 The estimated effective date in Norway is Mar 22, which is two days earlier than the official date of
Mar 24. However, the posterior distribution is very wide (± 10.79 days 95% CI; Figure 3): it covers the
128 range between Mar 9, three days before the first NPI, and Mar 29, five days after the last NPI (Table 1,
Figure S1). The uncertainty in the estimate of the effective start of NPIs might be due to the long
130 duration between the first and last NPIs; however, Germany had the second longest duration between
first and last NPIs, but the corresponding posterior distribution is quite narrow (Figure 3).

132 **Exact effective start of NPIs.** We find one case in which the official and effective dates match and
the credible interval is narrow. Switzerland ordered a national lockdown on Mar 20, after banning
134 public events and closing schools on Mar 13 and 14⁹. Indeed, the posterior median $\hat{\tau}$ is Mar 19 (± 2.51
days 95% CI, see Figure 3). It's also worth mentioning that Switzerland was the first to mandate self
136 isolation of confirmed cases⁹.

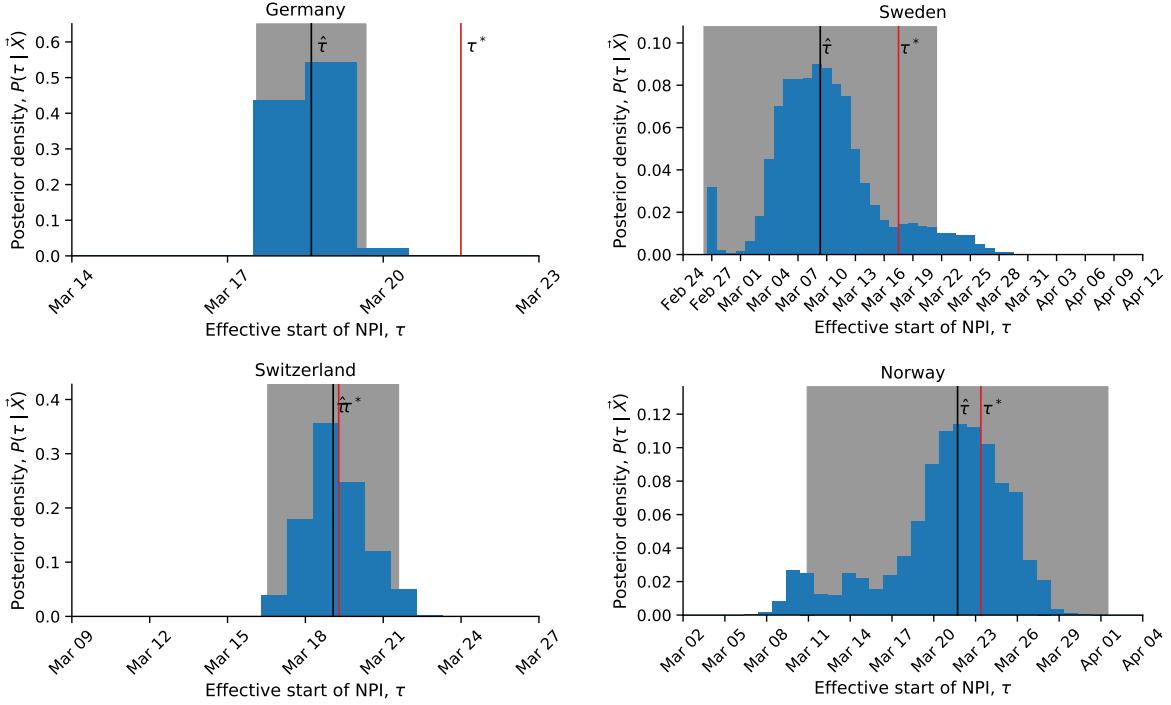


Figure 3: Early and exact effect of non-pharmaceutical interventions. Posterior distribution of τ , the effective start date of NPI, is shown as a histogram of MCMC samples. Red line shows the official last NPI date τ^* . Black line shows the estimated $\hat{\tau}$. Shaded area shows a 95% credible interval.

Assessment of impact of NPIs. The *effective reproduction number* R is the average number of secondary cases caused by an infected individual after an epidemic is already underway⁵. We infer the model-based effective reproduction numbers before and after the implementation of NPIs from model parameters (Eq. 7). We then estimate the impact of NPIs as the relative reduction in the reproduction number⁹. We compare the impacts estimated using the fixed τ model and the free τ model. That is, we compare the impact estimate assuming that NPIs started at the official date τ^* , versus the estimate when inferring the effective start of NPIs from the data. Figure 4 demonstrates that estimates from the fixed τ model are consistently lower than estimates from the free τ model, except in Sweden and Denmark, in which estimation uncertainty is high.

These results suggest that the impact of past NPIs is likely to be under-estimated by health officials and researchers if they assume NPIs effectively start at their official dates. The estimated impact can then be interpreted as ineffectiveness of the NPIs, leading to further escalations.

Health officials might also assess the impact of NPIs by comparing model expectations to actual number of daily cases at a specific number of days after the intervention began. However, a significant difference between the official and effective start of interventions can invalidate such assessments. This is illustrated in Figure 5 using data and parameters from Italy: a lockdown was officially ordered on Mar 10 (τ^*), but its late effect on the infection dynamics starts on Mar 17 ($\hat{\tau}$). If health officials assume that the dynamics change exactly at τ^* , they would expect the number of cases to be within the red lines (posterior predictions assuming $\tau = \tau^*$). This would lead to a significant under-estimation, which might be interpreted as ineffectiveness of the NPI, leading to more aggressive NPIs being applied. However, the number of cases would actually follow the blue lines (posterior predictions using $\tau = \hat{\tau}$), which are a good fit to the real data (stars).

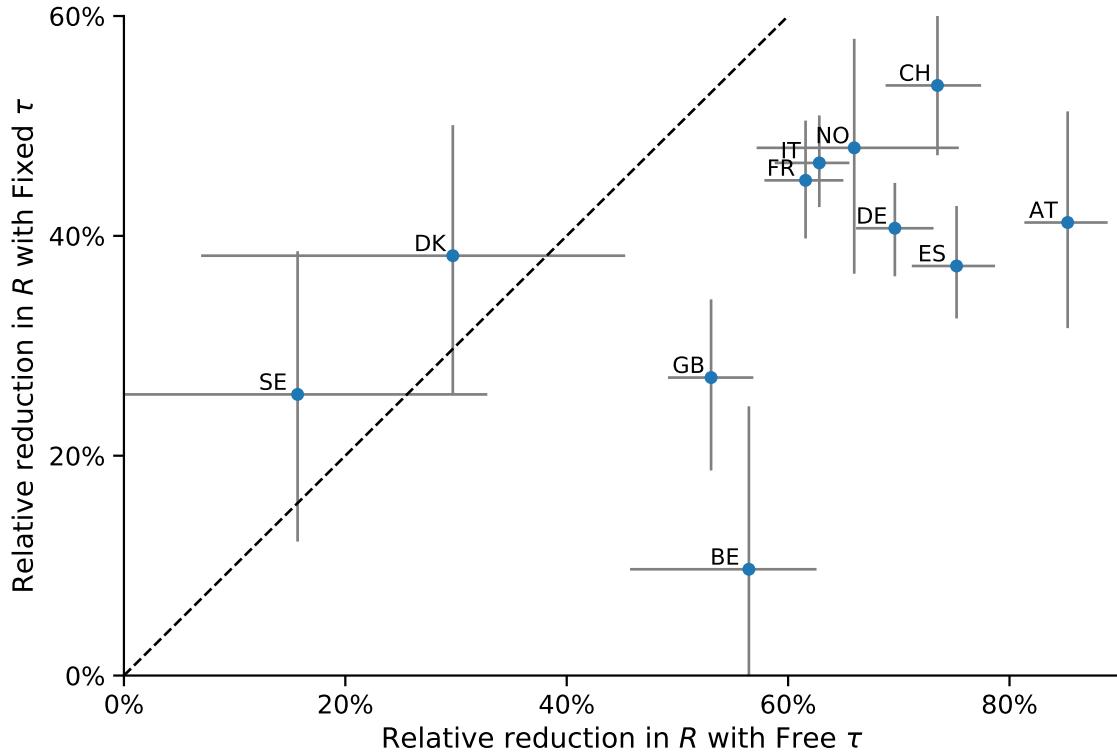


Figure 4: Impact of NPIs is under-estimated when assuming they start at the official date. Shown are estimates of the relative reductions in R (the effective reproduction number), which measures the impact of NPIs on disease transmission. The y-axis shows estimates when assuming the start of NPIs is fixed at the official date (fixed τ); the x-axis shows estimates when inferring the effective start of NPIs from the data (free τ). The dashed line shows a one-to-one correspondence. Markers and bars denote the posterior median and 50% credible intervals. The relative reductions in R are consistently lower for the fixed τ model (below the dashed line), except in Sweden and Denmark in which uncertainty is high.

Discussion

160 We have inferred the effective start date of NPIs in several geographical regions using an SEIR model
 161 under an MCMC parameter estimation framework. We find examples of both late and early effective
 162 start of NPIs relative to the official date (Figure 1).

In most investigated regions we find a late effective start of NPIs. For example, in Italy and in
 163 Wuhan, China, the effective start of the lockdowns seems to have occurred five or more days after
 164 the official date (Figure 2). This difference might be explained, in some cases, by low compliance
 165 or non-adherence to guidelines: In Italy, for example, the government plan to implement a lockdown
 166 in the Northern provinces leaked to the public, resulting in people leaving these provinces before
 167 the lockdown started¹¹. Late effect of NPIs may also be due to the time required by both the
 168 government and the citizens to prepare for a lockdown, and for new guidelines to be adopted by the
 169 population.

In contrast, in some regions we inferred reduced transmission rates even before official lockdowns
 170 were implemented, although this is only conclusive in Germany (Figure 3). An early effective date
 171 might be due to early adoption of social distancing and similar behavioural adaptations in parts of the
 172 population. Adoption of these behaviours may occur via media and social networks, rather than official
 173 government recommendations and instructions, and may be influenced by increased risk perception

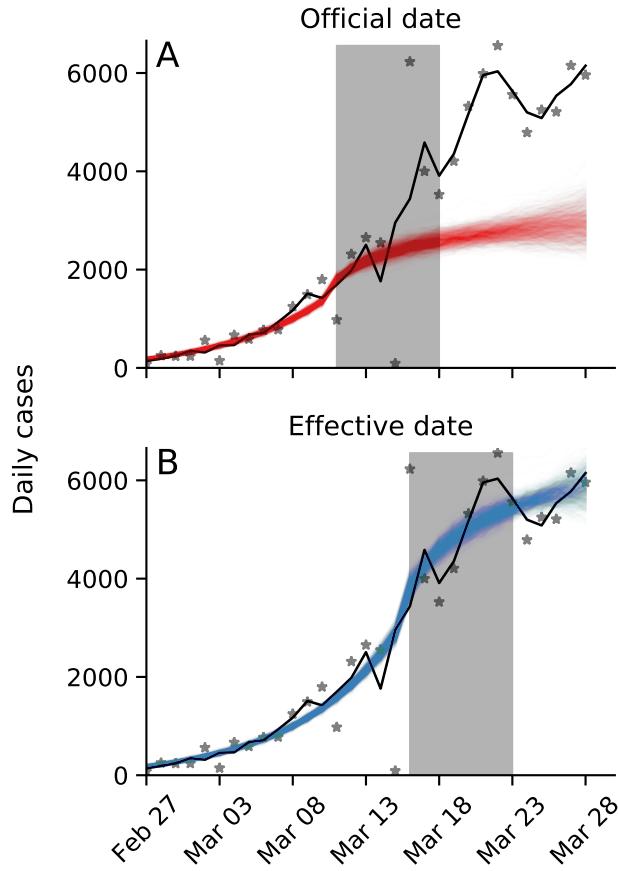


Figure 5: Late effective start of NPIs leads to under-estimation of daily confirmed cases. Real number of daily cases in Italy in black (markers: data; line: smoothed using a Savitzky-Golay filter with window length 3). Model posterior predictions are shown as coloured lines (1,000 draws from the posterior distribution). Shaded box illustrates a serial interval of seven days¹ after the start of NPIs. **(A)** Using the official date τ^* for the effective start of the NPI, Mar 11, the model under-estimates the number of cases seven days after the start of the NPI. **(B)** Using the estimated date $\hat{\tau}$ for the effective start of the NPI, Mar 17, the model precisely estimates the number of cases seven days after the start of the NPI. Here, model parameters are best estimates for Italy (Table 2).

176 due to domestic or international COVID-19-related reports². Indeed, the evidence supports a change
 177 in infection dynamics (i.e. a model with fixed or free τ) even for Sweden (Table S1, Table S2,
 178 Figure S4a), where a lockdown was not implemented*.

As expected, we have found that the evidence supports a model in which the transmission rate changes
 180 at a specific time point over a model with a constant transmission rate (Table S1 and Table S2). It may
 181 be interesting to check if the evidence favors a model with *two* or more change-points, rather than one.
 182 Multiple change-points could reflect escalating NPIs (e.g. school closures followed by lockdowns),
 183 or an intervention followed by a relaxation. However, interpretation of such models will be harder,
 184 as multiple change-points are also likely to result in parameter unidentifiability, for example due to
 simultaneous implementations of NPIs⁹.

186 Interestingly, the effective start of NPIs $\hat{\tau}$ in both France and Spain is estimated to have started on
 187 Mar 24, although the official NPI dates differ significantly: the first NPI in France is only one day
 188 before the last NPI in Spain. The number of daily cases was similar in both countries until Mar 8,
 189 but diverged by Mar 13, reaching much higher numbers in Spain (Figure S3). This may suggest
 correlations between effective starts of NPIs due to global or international events.

*Sweden banned public events on Mar 12, encouraged social distancing on Mar 16, and closed schools on Mar 18⁹.

As different countries experiment with various intervention strategies, we expect similar shifts to occur: in some cases the population will be late to comply with new guidelines, whereas in other cases the population will adopt either restrictions or relaxations even before they are formally announced. Attempts to asses the impact of NPIs^{4,9} generally assume they start at their official date, and that a significant change in the dynamics can be observed roughly seven days after the start of NPIs (due to the characteristic serial interval of COVID-19¹). However, late and early effective start of NPIs, such as we have inferred, can bias these assessments and lead to wrong conclusions about the impact of NPIs (Figure 4, Figure 5).

Our results highlight the complex interaction between personal, regional, and global determinants of behavioral response to an epidemic. Therefore, we emphasize the need to further study heterogeneity in compliance and behavior over both time and space. This can be accomplished both by surveying differences in compliance within and between populations³, and by incorporating specific behavioral models into epidemiological models^{2,7,21}.

Models and Methods

Data. We use daily confirmed case data $\mathbf{X} = (X_1, \dots, X_T)$ from 12 regions during Jan–Apr 2020. These incidence data summarise the number of individuals X_t tested positive for SARS-CoV-2 (using RT-qPCR tests) on each day t . Data for Wuhan, China, Jan 10 to Feb 8, retrieved from Pei and Shaman¹⁸. Data for 11 European countries, Feb 20 to Apr 24, retrieved from Flaxman et al.⁹. Where there were multiple sequences of days with zero confirmed cases (e.g. France), we cropped the data to begin with the last sequence so that our analysis focuses on the first sustained outbreak rather than isolated imported cases. For official NPI dates see Table 1.

SEIR model. We model SARS-CoV-2 infection dynamics by following the number of susceptible S , exposed E , reported infected I_r , unreported infected I_u , and recovered R individuals in a population of size N . This model distinguishes between reported and unreported infected individuals: the reported infected are those that have enough symptoms to eventually be tested and thus appear in daily case reports, to which we fit the model. This model is inspired by Li et al.¹⁷ and Pei and Shaman¹⁸, who used a similar model with multiple regions and constant transmission and reporting rates to study COVID-19 dynamics in China and in the continental US.

Susceptible (S) individuals become exposed due to contact with reported or unreported infected individuals (I_r or I_u) at a rate β_t or $\mu\beta_t$, respectively. The parameter $0 < \mu < 1$ represents the decreased transmission rate from unreported infected individuals, who are often subclinical or even asymptomatic^{8,20}. The transmission rate $\beta_t \geq 0$ may change over time t due to behavioural changes of both susceptible and infected individuals. Exposed individuals, after an average latent period of Z days, become reported infected with probability α_t or unreported infected with probability $(1 - \alpha_t)$. The reporting rate $0 < \alpha_t < 1$ may also change over time due to changes in human behaviour. Infected individuals remain infectious for an average period of D days, after which they either recover, or become ill enough to be quarantined. In either case, they no longer infect other individuals, and therefore effectively become recovered (R). The model is described by the following set of equations,

$$\begin{aligned} \frac{dS}{dt} &= -\beta_t S \left(\frac{I_r}{N} + \mu \frac{I_u}{N} \right) \\ \frac{dE}{dt} &= \beta_t S \left(\frac{I_r}{N} + \mu \frac{I_u}{N} \right) - \frac{E}{Z} \\ \frac{dI_r}{dt} &= \alpha_t \frac{E}{Z} - \frac{I_r}{D} \\ \frac{dI_u}{dt} &= (1 - \alpha_t) \frac{E}{Z} - \frac{I_u}{D} \\ \frac{dR}{dt} &= \frac{I_r}{D} + \frac{I_u}{D}, \end{aligned} \tag{1}$$

230 where N is the population size. The initial numbers of exposed $E(0)$ and unreported infected $I_u(0)$ are free
 231 model parameters (i.e. inferred from the data), whereas the initial number of reported infected and recovered is
 232 assumed to be zero, $I_r(0) = R(0) = 0$, and the number of susceptible is $S(0) = N - E(0) - I_u(0)$.

233 **Likelihood function.** For a given vector θ of model parameters the *expected* cumulative number of reported
 234 infected individuals (I_r) until day t , following Eq. 1, is

$$Y_t(\theta) = \int_0^t \alpha_s \frac{E(s)}{Z} ds, \quad Y_0 = 0. \quad (2)$$

235 We assume that reported infected individuals are confirmed and therefore observed in the daily case report of
 236 day t with probability p_t (note that an individual can only be observed once, and that p_t may change over time,
 237 but t is a specific date rather than the time elapsed since the individual was infected). We denote by X_t the
 238 *observed* number of confirmed cases in day t , and by \tilde{X}_t the cumulative number of confirmed cases until end of
 239 day t ,

$$\tilde{X}_t = \sum_{i=1}^t X_i. \quad (3)$$

240 Therefore, at day t the number of reported infected yet-to-be confirmed individuals is $(Y_t(\theta) - \tilde{X}_{t-1})$. We assume
 241 that X_t conditioned on \tilde{X}_{t-1} is Poisson distributed, such that

$$\begin{aligned} (244) \quad & (X_1 | \theta) \sim Poi(Y_1(\theta) \cdot p_1), \\ & (X_t | \tilde{X}_{t-1}, \theta) \sim Poi((Y_t(\theta) - \tilde{X}_{t-1}) \cdot p_t), \quad t = 2, \dots, T. \end{aligned} \quad (4)$$

242 Hence, the *likelihood function* $\mathcal{L}(\theta | \mathbf{X})$ for a parameter vector θ given the confirmed case data $\mathbf{X} = (X_1, \dots, X_T)$
 243 is defined by the probability to observe \mathbf{X} given θ ,

$$\mathcal{L}(\theta | \mathbf{X}) = P(\mathbf{X} | \theta) = P(X_1 | \theta) \cdot P(X_2 | \tilde{X}_1, \theta) \cdots P(X_T | \tilde{X}_{T-1}, \theta). \quad (5)$$

244 **NPI model.** To model non-pharmaceutical interventions (NPIs), we set the start of the NPIs to day τ and
 245 define

$$(246) \quad \beta_t = \begin{cases} \beta, & t < \tau \\ \beta\lambda, & t \geq \tau \end{cases}, \quad \alpha_t = \begin{cases} \alpha_1, & t < \tau \\ \alpha_2, & t \geq \tau \end{cases}, \quad p_t = \begin{cases} 1/9, & t < \tau \\ 1/6, & t \geq \tau \end{cases}, \quad (6)$$

247 where $0 < \lambda < 1$. The values for p_t follow Li et al.¹⁷, who estimated the average time between infection and
 248 reporting in Wuhan, China, at 9 days before the start of NPIs and 6 days after start of NPIs.

Following Li et al.¹⁷, the effective reproduction numbers before and after the start of NPIs are

$$\begin{aligned} (254) \quad & R_1 = \alpha_1 \beta D + (1 - \alpha_1) \mu \beta D, \\ & R_2 = \alpha_2 \lambda \beta D + (1 - \alpha_2) \mu \lambda \beta D. \end{aligned} \quad (7)$$

The relative reduction in the effective reproduction number due to NPIs is $\frac{R_1 - R_2}{R_1}$.

256 **Parameter estimation.** To estimate the model parameters from the daily case data \mathbf{X} , we apply a Bayesian
 257 inference approach. Model fitting was calibrated for case data up to Mar 28, and then applied to data up to
 258 Apr 11 (for Wuhan, China, model fitting was applied for data up to Feb 8.) We start our model Δt days¹¹ before
 259 the outbreak (defined as consecutive days with increasing confirmed cases) in each country. The model in Eqs. 1
 260 and 6 is parameterised by the vector θ , where

$$\theta = (Z, D, \mu, \beta, \lambda, \alpha_1, \alpha_2, E(0), I_u(0), \Delta t, \tau). \quad (8)$$

262 The likelihood function is defined in Eq. 5. We defined the following prior distributions on the model parameters $P(\theta)$:

$$Z \sim Uniform(2, 5)$$

$$D \sim Uniform(2, 5)$$

$$\mu \sim Uniform(0.2, 1)$$

$$\beta \sim Uniform(0.8, 1.5)$$

$$\lambda \sim Uniform(0, 1)$$

264 $\alpha_1, \alpha_2 \sim Uniform(0.02, 1)$ (9)

$$E(0) \sim Uniform(0, 3000)$$

$$I_u(0) \sim Uniform(0, 3000)$$

$$\Delta t \sim Uniform(1, 5)$$

$$\tau \sim TruncatedNormal\left(\frac{\tau^* + \tau^0}{2}, \frac{\tau^* - \tau^0}{2}, 5, T - 2\right),$$

266 where the prior for τ is a truncated normal distribution shaped so that the date of the first and last NPI, τ^0 and τ^* (Table 1), are at minus and plus one standard deviation, and taking values only between 5 and $T - 2$, where T is the number of days in the data \mathbf{X} . We also tested an uninformative uniform prior, $Uniform(1, T - 2)$.

268 WAIC (Eq. 10) of a model with this uniform prior was either higher, or lower by less than 2, compared to WAIC of a model with the truncated normal prior. The uninformative prior resulted in non-negligible posterior probability for unreasonable τ values, such as Mar 1 in the United Kingdom. We therefore decided to use the more informative truncated normal prior for τ . Other priors follow Li et al.¹⁷, with the following exceptions.

270 λ is used to ensure transmission rates are lower after the start of the NPIs ($\lambda < 1$). We checked values of Δt larger than five days and found they generally produce lower likelihood and unreasonable parameter estimates, 272 and therefore chose $Uniform(1, 5)$ as the prior for Δt . We also tried to estimate the value of p_t before and after 274 τ , instead of keeping it fixed at 1/9 and 1/6. The model with fixed values was supported by the evidence (lower 276 WAIC, see below) in 9 of 12 countries. Moreover, the estimates for Wuhan, China were 1/9 and 1/6, as in Li et al.¹⁷.

278 The posterior distribution of the model parameters $P(\theta | \mathbf{X})$ is estimated using the affine-invariant ensemble sampler for Markov chain Monte Carlo (MCMC)¹³ implemented in the emcee Python package¹⁰. We used 280 the default configuration using the stretch move with stretch scale parameter $a = 2$. For the main analysis we 282 use 50 chains (or walkers) per region, with 7M samples per chain (no thinning was applied). The *integrated* 284 *autocorrelation time*^{10,13} was averaged across parameters and chains for each region (FIGURE). It was found to be between XXX for Norway and XXX for Germany. We examined the trace plots for τ in all regions 286 (FIGURE). All chains seem to converge to the same stationary distribution, in most cases before 2M steps. Thus, we discarded the first 2M samples as burnin samples. The only exception is Spain, in which a single 288 chain converged at around X steps. We considered this chain as part of the burnin and removed in the analysis. We also ran additional chains with 2M samples and a different initialization (i.e. seed). The estimated posterior 290 distributions of τ are very similar to our main analysis, further increasing our confidence in the convergence of our inference. We also examined the joint posterior distributions of τ and λ to check for correlations (FIGURE).

292 **Model comparison.** We perform model selection using two methods. First, we compute WAIC (widely applicable information criterion)¹²,

$$WAIC(\theta, \mathbf{X}) = -2 \log \mathbb{E}[\mathcal{L}(\theta | \mathbf{X})] + 2\mathbb{V}[\log \mathcal{L}(\theta | \mathbf{X})] \quad (10)$$

294 where $\mathbb{E}[\cdot]$ and $\mathbb{V}[\cdot]$ are the expectation and variance operators taken over the posterior distribution $P(\theta | \mathbf{X})$. We compare models by reporting their relative WAIC; lower is better (Table S1).

296 We also plot posterior predictions: we sample 1,000 parameter vectors from the posterior distribution $P(\theta | \mathbf{X})$ fitted to data up to Apr 11, use these parameter vectors to simulate the SEIR model (Eq. 1) up to Apr 24, and plot 298 the predicted dynamics (Figure S4a). Both the accuracy (i.e. overlap of data and prediction) and the precision

(i.e. the tightness of the predictions) are good ways to visually compare models. We also compute the expected
300 posterior RMSE (root mean squared error) of these predictions (Table S2).

Declarations

302 **Ethics approval and consent to participate.** Not applicable.

Consent for publication. Not applicable.

304 **Availability of data and materials.** We use Python 3 with the NumPy, Matplotlib, SciPy, Pandas, Seaborn,
and emcee packages. Source code will be publicly available under a permissive open-source license at
306 github.com/yoavram-lab/EffectiveNPI. We used freely available data, see source code repository and above
Data section for details.

308 **Competing interests.** The authors declare that they have no competing interests.

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312 **Authors' contributions.** UO and YR designed the research. IK and YR performed the research and wrote the
manuscript. All authors read and approved the final manuscript.

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Country	τ^*	τ	$CI_{75\%}$	$CI_{95\%}$	Z	D	μ	β	α_1	λ	α_2	$E(0)$	$I_u(0)$	Δt
Austria	Mar 16	Mar 21	2.5658	3.8148	3.9102	3.6649	0.6407	1.1417	0.1308	0.1387	0.2261	151.1078	131.6623	2.1308
Belgium	Mar 18	Mar 27	2.3456	6.2773	3.9098	3.6152	0.4877	1.0762	0.2721	0.4388	0.2363	309.9114	426.9644	2.1756
Denmark	Mar 18	Mar 11	5.0053	15.3374	4.0957	3.5091	0.4161	1.0747	0.1470	0.6919	0.1867	324.6332	389.5076	2.2071
France	Mar 17	Mar 24	0.5068	2.2928	4.2159	3.1688	0.4863	1.0502	0.3726	0.3765	0.4168	422.6720	1362.5052	1.6717
Germany	Mar 22	Mar 19	0.7657	1.0508	3.3917	3.7166	0.6780	1.1515	0.1753	0.2664	0.4147	529.9397	387.1038	2.1516
Italy	Mar 11	Mar 17	2.1045	2.9945	4.1439	2.5550	0.5371	0.9932	0.5816	0.3860	0.4973	990.1942	1902.7806	1.6367
Norway	Mar 24	Mar 22	4.2700	10.7889	4.0476	3.1061	0.3885	1.0467	0.1536	0.3091	0.2320	478.0678	856.7700	1.9998
Spain	Mar 14	Mar 25	0.8415	2.9656	4.1392	3.3352	0.5974	1.1574	0.3640	0.2551	0.3050	310.0470	957.8280	1.6062
Sweden	Mar 18	Mar 09	4.9930	11.7493	4.0328	3.3962	0.3656	1.0457	0.1427	0.7237	0.3093	394.0202	543.5448	2.3755
Switzerland	Mar 20	Mar 19	1.4513	2.5140	3.9764	3.4425	0.6360	1.1543	0.1283	0.2481	0.2246	336.3842	328.7800	1.9891
United Kingdom	Mar 24	Mar 26	1.4721	2.5543	3.9654	3.6333	0.6414	1.1168	0.1212	0.4504	0.1737	422.8841	486.0467	2.0540
Wuhan, China	Jan 23	Feb 03	1.7984	2.8493	3.7326	3.6320	0.6057	1.1453	0.2754	0.1784	0.3511	597.8676	561.1586	2.4248

Table 2: Parameter estimates for different regions. See Eq. 1 for model parameters. All estimates are posterior medians. 75% and 95% credible intervals given for τ , in days. τ^* is the official last NPI date, see Table 1.

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Supplementary Material

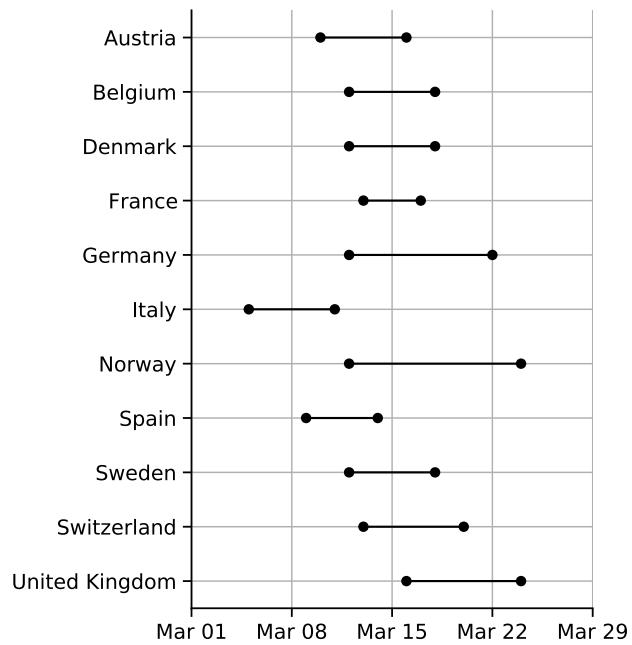


Figure S1: Official start of non-pharmaceutical interventions. See Table 1 for more details. Wuhan, China is not shown.

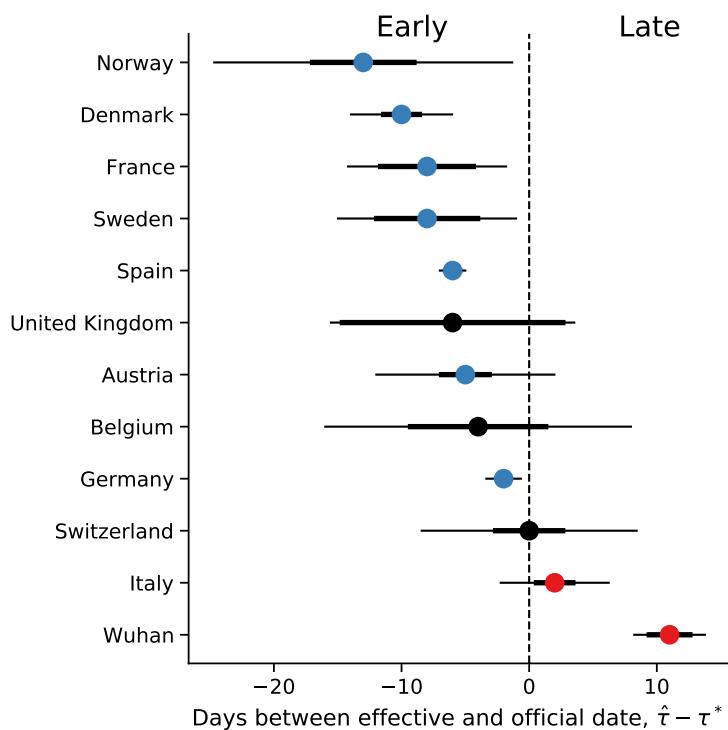


Figure S2: Official vs. effective start of non-pharmaceutical interventions estimated up to Mar 28. The difference between τ the effective and τ^* the official start of NPIs estimated from case data up to Mar 28, 2020, shown for different regions. Here, $\hat{\tau}$ is the marginal posterior median, see Table 2. τ^* is the last NPI date (a lockdown everywhere by Sweden, see Table 1). Thin and bold lines show 95% and 75% credible intervals (HDI), respectively.

WAIC

Country	No	Fixed	Free
Austria	219.49	95.06	35.96
Belgium	148.37	98.41	49.16
Denmark	44.36	40.30	43.11
France	581.59	255.14	172.08
Germany	1029.36	327.50	174.90
Italy	898452.34	5484.56	80.18
Norway	70.03	42.04	39.79
Spain	1476.46	647.34	128.58
Sweden	32.53	30.06	31.10
Switzerland	265.80	83.95	63.89
United Kingdom	258.18	117.54	68.17
Wuhan China	107.31	94.00	73.75

Table S1: WAIC values for the different models. WAIC (widely applicable information criterion; Eq. 10)¹² values for models with: no τ at all, *No*; τ fixed at the official last NPI date τ^* , *Fixed*; and free parameter τ , *Free*. WAIC values are scaled as a deviance measure: lower values imply higher predictive accuracy and a difference of 2 is a popular threshold for model comparison¹⁴. Bold values emphasize cases in which the *Free* model has the lowest WAIC.

RMSE

Country	No	Fixed	Free
Austria	926.7	740.7	58.9
Belgium	3286.0	2541.0	902.9
Denmark	434.8	1058.0	400.0
France	10530.0	8206.0	1816.0
Germany	14240.0	17760.0	1952.0
Italy	14160.0	9536.0	1106.0
Norway	292.2	578.2	63.3
Spain	17840.0	14410.0	1704.0
Sweden	775.9	1205.0	596.4
Switzerland	1839.0	1721.0	214.0
United Kingdom	14700.0	14780.0	2735.0

Table S2: Posterior RMSE of out-of-sample predictions with the different models. Expected posterior predictive RMSE (root mean squared error) for models with: no τ at all, *No*; τ fixed at the official last NPI date τ^* , *Fixed*; and free parameter τ , *Free*. In all cases, the model with free parameter τ has the lowest RMSE. Models were fitted to case data up to Apr 11, 2020, and then used to generate 1,000 predictions up to Apr 24 by sampling model parameters from the posterior distribution. These predictions were then compared to the real data using RMSE, and the mean RMSE value is shown in the table for each country and model. Bold values emphasize cases in which the *Free* model has the lowest RMSE.

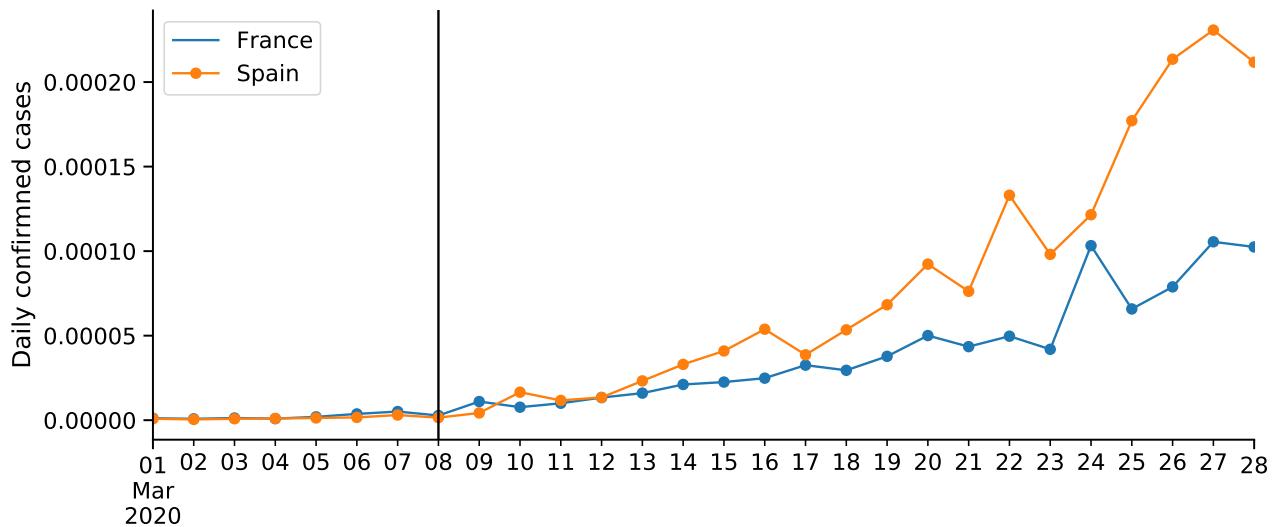
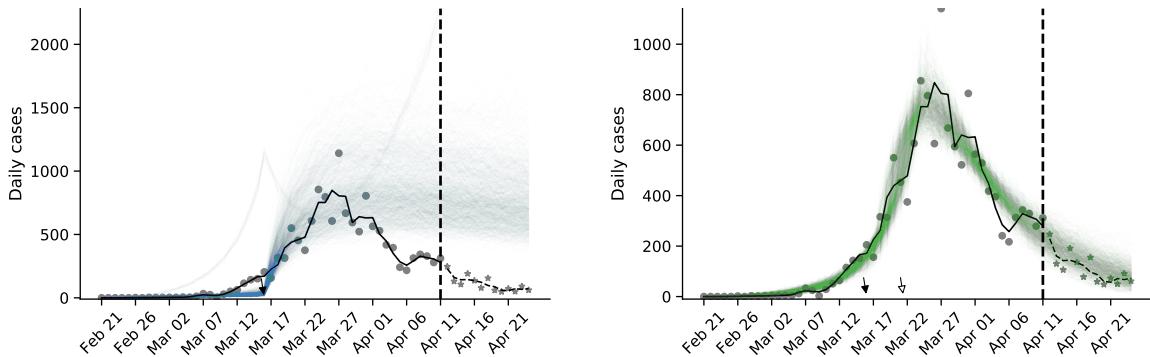
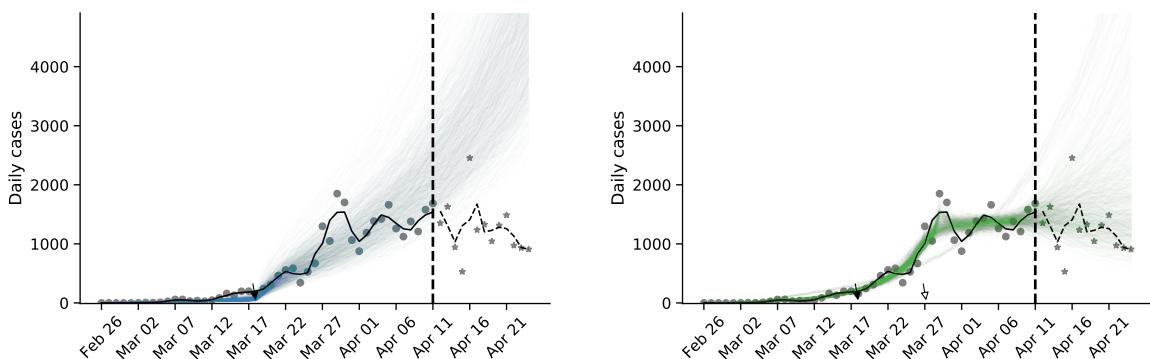


Figure S3: COVID-19 daily confirmed cases in France and Spain. Number of cases proportional to population size (as of 2018). Vertical line shows Mar 8, the effective start of NPIs $\hat{\tau}$ in both countries. Data from Flaxman et al.⁹.

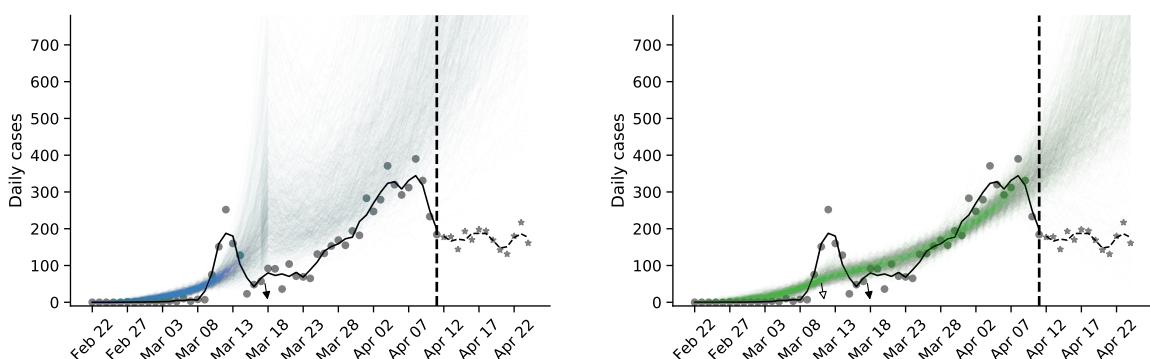
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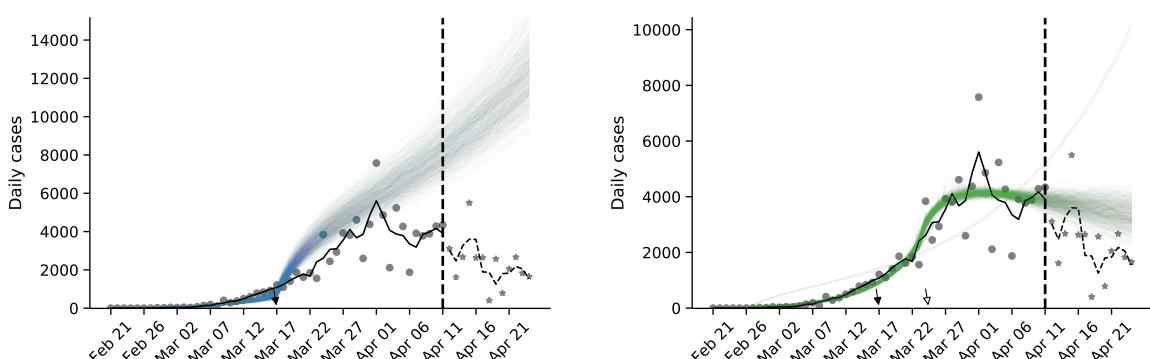
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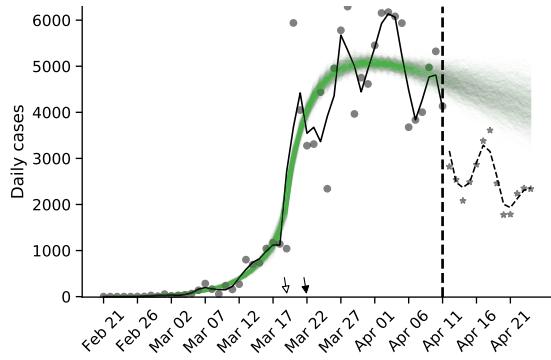
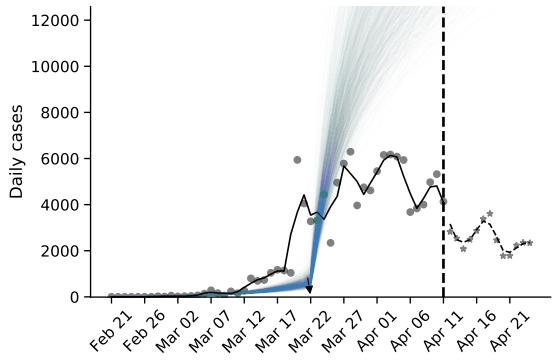
Denmark



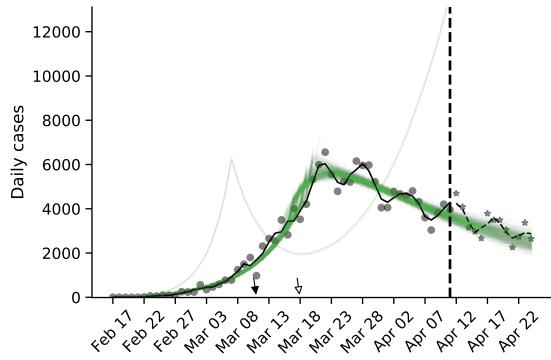
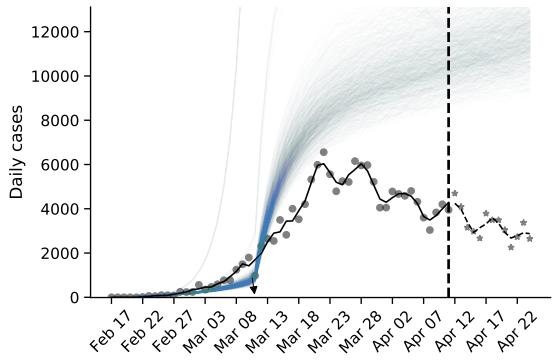
France



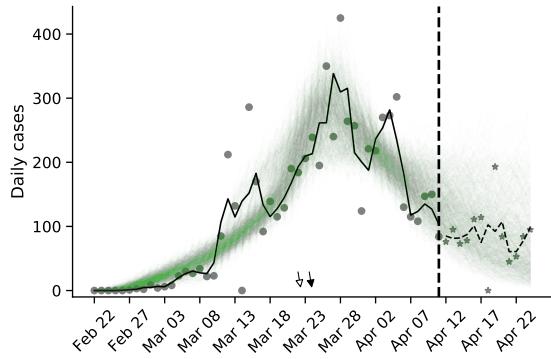
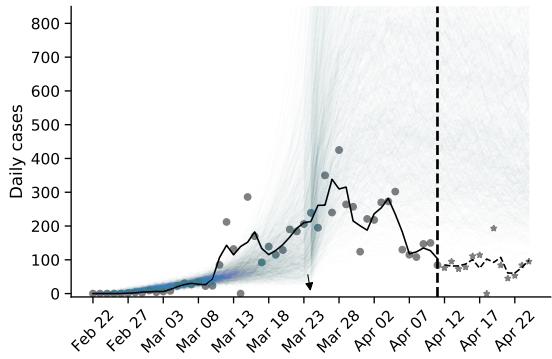
Germany



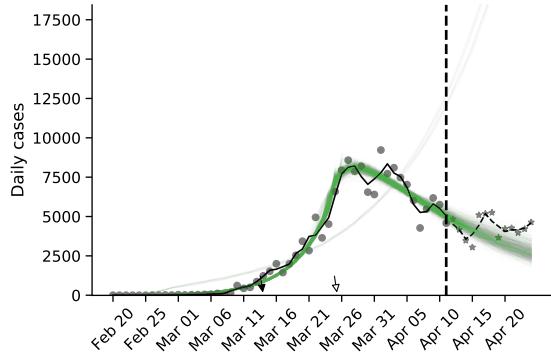
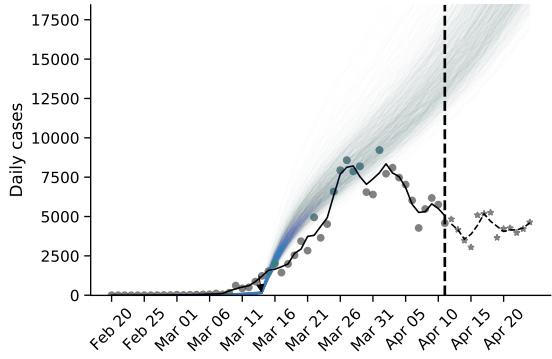
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Norway



Spain



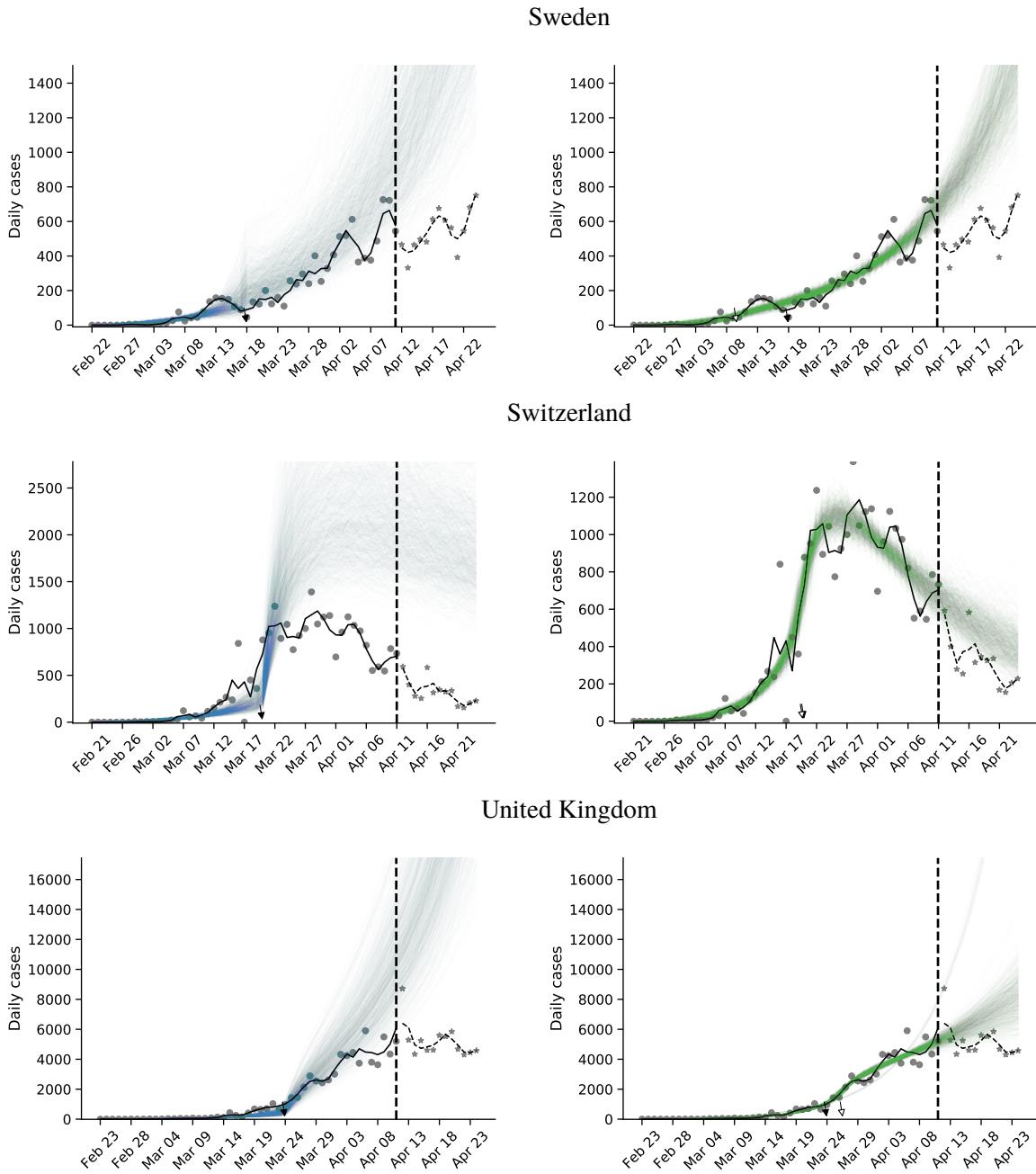


Figure S4. Posterior prediction plots for 11 European countries. The vertical dashed line represents Apr 11, 2020. Circles and stars represent daily case data up to and after Apr 11, respectively. Black and white arrows denote the official τ^* and effective $\hat{\tau}$ start of NPIs, respectively. Black lines represent a smoothing of the data points using a Savitzky-Golay filter with window length 3. Coloured lines represent posterior predictions from a model with fixed τ (blue) and free τ (green). Models were fitted with data up to Apr 11. The predictions are generated by drawing 1,000 parameter sets from the posterior distribution, and then generating a daily case count using the SEIR model up to Apr 24. Note the differences in the y-axis scale. Posterior predictions with the free τ model predict the out-of-sample data well for all countries except Denmark and Sweden, but poorly for the fixed τ model. The predictions of the model without τ (not shown) are even worse.