

A Bayesian Machine Learning Approach for Spatio-Temporal Prediction of COVID-19 Cases

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**1 A Bayesian Machine Learning Approach for
2 Spatio-Temporal Prediction of COVID-19 Cases**

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7 Abstract Modeling the behavior and spread of infectious diseases on space and
8 time is key in devising public policies for preventive measures. This behavior is so
9 complex that there are lots of uncertainties in both the data and in the process
10 itself. We argue here that these uncertainties should be taken into account in the
11 modeling strategy. Machine learning methods, and neural networks, in particular,
12 are useful in modeling this sort of complex problems, although they generally lack
13 of probabilistic interpretations. We thus present here a neural network method
14 embedded in a Bayesian framework for modeling and predicting the number of
15 cases of infectious diseases in areal units. A key feature is that our combined
16 model considers the impact of human movement on the spread of the infectious
17 disease, as an additional random factor to the also considered spatial neighborhood
18 and temporal correlation components.

19 Our model is evaluated over a COVID-19 dataset for 245 health zones of
20 Castilla-Leon (Spain). The results show that a Bayesian model informed by a neu-
21 ral network method is generally able to predict the number of cases of COVID-19
22 in both space and time, with the human mobility factor having a strong influence
23 on the model.

24 Keywords Bayesian inference, COVID-19, Neural network, Poisson regression,
25 Public mobility

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26 1 Introduction

27 Infectious diseases are the main cause of health hazards in the world and are re-
28 sponsible for deaths of millions of people around the world (WHO, 2019). Various
29 outbreaks of infectious diseases have occurred throughout human history, and in-
30 deed there is currently a global health pandemic caused by the novel Coronavirus
31 disease (COVID-19). More than 90 million people have been infected and more
32 than 2 million people have lost their lives (Worldometer, 2020; Wu et al., 2020)
33 as of January 2021 due to COVID-19. To contain the spread of this virus, vari-
34 ous regulations such as social distancing measures, travel restrictions, and city or
35 nation-wide lockdowns have been put in place by policy makers around the world.
36 These regulations, although effective in containing the spread of the disease, have
37 also impacted the daily lives of people, social behavior and the global supply chain
38 (Jones et al., 2008). The transmission of general infectious diseases (e.g. COVID-
39 19) exhibits spatio-temporal patterns and can be predicted based on ecological,
40 environmental and socio-economic factors (Anno et al., 2019; Yang et al., 2020).
41 Prediction of these infections is important for government and health workers to
42 plan for effective mitigation by prioritizing the actions of prevention and control
43 measures (Remuzzi & Remuzzi, 2020).

44 Human movement typically stimulates the introduction of infectious diseases
45 into a new region. There are various evidences that due to human movement,
46 a region-specific disease is introduced to a new region (Stoddard et al., 2009;
47 Nunes et al., 2014) and spreads locally (Stoddard et al., 2013; Gross et al., 2020).
48 Indeed, a number of recent studies have incorporated human movement factors
49 into the modeling strategy (M. U. G. Kraemer et al., 2019; Massaro et al., 2019;
50 Mukhtar et al., 2020). For example, the increased human mobility in western Africa
51 had a high impact in making the Ebola virus catastrophic (Farrar & Piot, 2014).
52 Bogoch et al. (2015) studied the air transport data of flights going out of the
53 Ebola virus affected countries, finding air transport also one of the reasons for the
54 transmission. In the case of COVID-19, it is also seen that the measures related
55 to human movements, such as travel restrictions and social distancing, have been
56 effective in containing the diseases (Fang et al., 2020; M. Kraemer et al., 2020).
57 It is a fact that the introduction of human mobility in epidemiological studies has
58 been more accessible due to technological advancements in locational services and
59 availability of movement data (Guinness, 2016; Sedlar et al., 2019). In this context,
60 availability of technologies such as WiFi or cell phone tower positioning systems
61 and global navigation satellite systems have made the analysis of mobility much
62 easier (Gonzalez et al., 2008; Toch et al., 2019).

63 The spread of infectious diseases in space and their outbreak in time consti-
64 tute a complex spatio-temporal problem, which is an effect of complex dynamics
65 of human behavior, environment, and their interactions. Furthermore, as reported
66 in Pan et al. (2020), during pandemics the human mobility pattern changes com-
67 pared to that of other times which makes the problem more complex and difficult
68 to analyze. Deep learning methods have proven to be suitable for modeling such
69 complex problems (Mosavi et al., 2020). Indeed, (Ak et al., 2018; Titus Muurlink
70 et al., 2018; Akhtar et al., 2019; Anno et al., 2019; Kapoor et al., 2020; Wiec-
71 zorek et al., 2020) have used neural networks (some with human mobility data)

72 to model the spread of infectious diseases. Neural network-based methods rely on
73 a hidden stage to learn from the data and are unable to explicitly account for
74 the spatial and spatio-temporal random effects. However, although these methods
75 have performed well, they are unable to provide uncertainties in the predictions,
76 which we believe are essential in statistical inference and probabilistic forecasting.
77 We argue that predictions accompanied with uncertainties provide further
78 confidence on the results (Beale & Lennon, 2012). To incorporate uncertainties
79 in neural networks, Bayesian neural networks have been developed (Kononenko,
80 1989; Dhamodharavadhani et al., 2020) and applied over various spatio-temporal
81 problems (McDermott & Wikle, 2019). However, in the field of modeling and un-
82 derstanding the dynamics of COVID-19, the use of neural networks in combination
83 with Bayesian inference is limited. Cabras (2020) presented a method of combining
84 neural networks with Bayesian inference having a focus on COVID-19 infections
85 in Spain. However, mobility and its influences were not considered. As spatio-
86 temporal predictions help in understanding the spread of the disease to further
87 identify the regions of high risk, a large number of papers can be found in the field
88 of spatio-temporal modeling of diseases. Among them, generalized linear models
89 (GLM) with the addition of spatial effects of nearby places and/or temporal effects
90 from past events are found to be often used and proven to be useful in prediction
91 (Guo et al., 2017; Cabrera & Taylor, 2019; Giuliani et al., 2020). For example,
92 Giuliani et al. (2020) have used GLM to predict COVID-19 infections in regions
93 of Italy, and found the spatial interactions of nearby places to have a high influ-
94 ence on modeling; this shows the importance of accounting for the spatial effects
95 explicitly. In a parallel vein, Bayesian modeling methods have also been used in
96 this epidemiological context (Gelman et al., 2013; Aswi et al., 2019; Song et al.,
97 2019; Torres-Signes et al., 2020).

98 In this context, our approach focuses on the use of deep learning methods (us-
99 ing a Long Short Term Memory-LSTM) informing a Poisson regression model in
100 a Bayesian framework to model and predict the spread and outbreak of COVID-19
101 with uncertainties. In particular, human mobility data along with socio-demographic
102 variables are incorporated in the combined model to predict the dynamics of
103 COVID-19. In doing so, we highlight the importance of human mobility in mod-
104 eling the dynamics of infectious diseases.

105 The plan of the paper is as follows. Section 2 presents the data along with
106 all covariates considered in the model to motivate the proposed statistical model.
107 We also consider some spatial weights built from the movement data. Section 3
108 presents the statistical model, and the results come in Section 4. The paper ends
109 with some conclusions and a discussion in Section 5.

110 **2 Study area and data**

111 Daily COVID-19 infections aggregated per 245¹ health zones in the community
112 of Castilla-Leon (Spain) were used in this paper. The temporal range goes from
113 March 1, 2020 to February 5, 2021. Castilla-Leon is the largest community in Spain

¹ Here, the health zones SORIA NORTE, SORIA SUR and SORIA RURAL are aggregated to a single unit.

114 by area located in the northwest part of Spain. This region has a population of
 115 around 2.5 million and is ranked third among the communities in offering social
 116 services to the citizens. Figure 1 shows the location map of the study area and also
 117 cumulative COVID-19 cases per 10000 inhabitants per health zone until February
 118 5, 2021. We note that COVID-19 has spread throughout the study area with
 119 clusters around major urban areas.

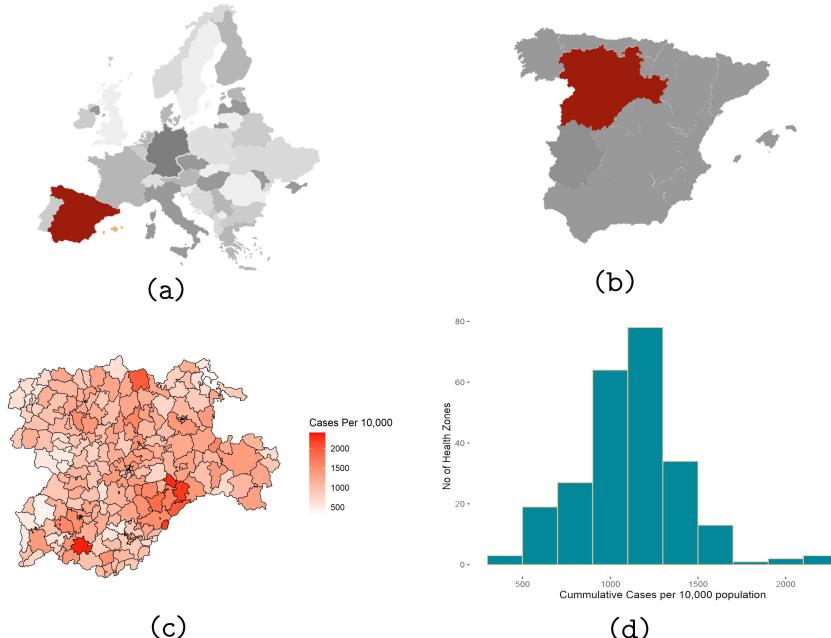


Fig. 1: (a) Location of Spain; (b) Location of Castilla-Leon in Spain; (c) Cumulative numbers of COVID-19 cases per 10000 inhabitants and health zones; (d) Histogram of cumulative cases per health zones of Castilla-Leon

120 COVID-19 cases data were retrieved from the open data portal of Castilla-
 121 Leon ². Similarly, the socio-demographic datasets and the health zone boundary, in
 122 shapefile form, were downloaded from the open data platform of Instituto Nacional
 123 de Estadística ³. The human mobility data for the study area was acquired from
 124 Barcelona Supercomputing Center flowmap dashboard ⁴. A brief description and
 125 source of the datasets used in the current paper are reported in Table 1.

126 Figure 2 shows the daily number of COVID-19 cases per 10000 inhabitants.
 127 The highlighted red line represents the daily mean number of cases per 10000
 128 inhabitants. The cases increased in March and April 2020 (defining the first wave),
 129 and then started to decrease until August 2020 due to the imposed lockdown

² <https://datosabiertos.jcyl.es/web/es/datos-abiertos-castilla-leon.html>

³ <https://www.ine.es/en/index.htm>

⁴ <https://flowmaps.life.bsc.es/flowboard/>

Table 1: Summary of data used and their sources

Data	Data Sources	Description of data
COVID-19	Open data portal of Castilla-Leon	Daily infected cases at health zone level
Mobility data	Barcelona Supercomputing Center	Daily human mobility matrices at municipality level
Socio-demographic	Open data portal of Castilla-Leon	Individual health zone total population, unemployment level and number of urban offices
Geometry	Open data portal of Castilla-Leon	Boundary shapefiles of 245 health zones

measures. However, due to a certain relaxation towards the summer period, the cases started to increase late August to end up with a second wave in October and November 2020. A third wave of infection is noted in January and February 2021, and started to decrease again due to some partial restrictions and the onset of the vaccination program. Similarly, weekly trends in the number of cases is visible with a drop of cases on weekends, due to the reduced number of tests done over the weekends.

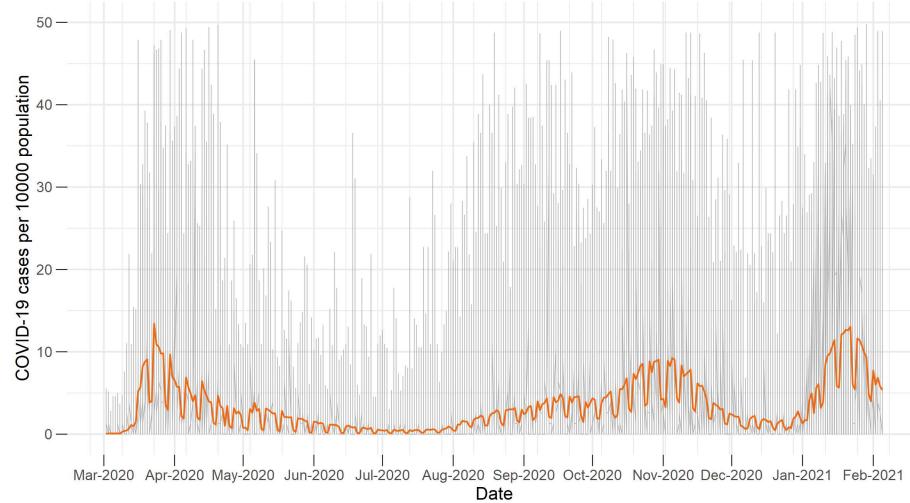


Fig. 2: Temporal trend of COVID-19 cases in the study area. The red line represents the daily mean number of cases per 10000 inhabitants.

The mobility data acquired from the data portal of Barcelona Supercomputing Center was prepared by the Ministry of Transport, Mobility, and Urban Agenda. The data was preprocessed to guarantee anonymized records from mobile phones.

140 These recorded events contain both active events also known as Call Detail Records
 141 (CDR) and passive events with a periodic update of device position, change of
 142 coverage area, etc. The location information is at the level of the coverage area of
 143 each antenna, which is merged to create origin-destination matrices at municipali-
 144 ty, districts and provinces level. Along with these records from the cell phones,
 145 landuse data, population data, transport network data such as train lines, and
 146 location of airports have been used to create the merged matrices (Ministry of
 147 Transport & Agenda, 2020). The available daily mobility data was at the munici-
 148 pality level; those municipalities with population less than 1000 were combined to
 149 form aggregated zones. As all other available data were at the health zones level,
 150 these aggregations were converted to the health zone level by applying spatial
 151 overlay functions and dividing the movement data in proportion to the area. The
 152 socio-demographic covariates considered in this paper were the following: total
 153 population per health zone, number of people demanding for employment, num-
 154 ber of unemployed people, number of commercial units, office units, and industrial
 155 units in the urban areas of each health zone (see a description in Table 2). Addi-
 156 tionally, we also considered some built-in variables (see Table 3). In particular, we
 157 computed the average number of cases and average number of deaths in the direct
 158 neighborhood. The cumulative cases of COVID-19 for the last 14 days were also
 159 computed to consider the aggregated impact for a short time frame.

Table 2: Summary of socio-demographic variables

Variable Name	Description
<code>total.pop</code>	total population of the health zone
<code>demanding_total_employment</code>	Number of people demanding for employment
<code>registered_unemployed_total</code>	Number of people registered as unemployed
<code>number_of_urban_commercial_units</code>	Number of commercial offices in the urban areas
<code>number_of_urban_industrial_units</code>	Number of industrial units in the urban areas
<code>number_of_urban_office_units</code>	Number of offices units in the urban areas

Table 3: Summary of built-in variables

Variable Name	Description
Day of the week	Computed from the date
Cumulative cases	Cumulative number of cases for last 14 days
Average number of cases in neighboring health zones	Average of number of cases in health zones that share a common border
Average number of deaths in neighboring health zones	Average number of deaths in health zones that share a common border

160 Last, but not least, we introduce new spatial weights based on the movement
 161 data that represent the associated movement-based risk. These weights are com-

162 puted per health zone and day. We add a temporal lag to handle past-term move-
163 ment data and the daily data are weighted depending on the temporal distance.

164 These spatial weights take into account the mobility from all other regions j
165 into region i , and the weights are interpreted as the chance of a moving person to
166 import the infection of the disease into region i from all the other regions. This
167 spatial weight for a region i and day t , $W_{i,t}$, can be computed as

$$168 \quad W_{i,t} = \sum_{j=1}^n \left[\sum_{t'=t-1}^{t-\Delta t} m_{ji,t'} * w'_{t'} \right] * \frac{I_{j,t}}{P_j} \quad (1)$$

169 where n is total number of regions, $m_{ji,t}$ is the mobility from all regions j to i on
170 day t , $I_{j,t}$ is the number of infected cases at region j at time t , P_j is the total
171 population of the region j and $w'_{t'}$ is the weight given to the mobility data on day
172 t .

173 A time lag Δt is added to the computation of the spatial weights as the spread
174 of a disease on the region is dependent on the mobility and infections on past days
175 in all other regions of the study area. We used a 7-day lag as infection is assumed
176 to act a week before first symptoms. We assigned the following weights: given t ,
177 we give $t - 1$ and $t - 2$ only a weight of 5%, this weight increases up to 10% for
178 $t - 3$ and $t - 4$, then goes up to 20% for $t - 5$ and $t - 6$, and finally the weight is
179 30% for $t - 7$.

180 Figure 3 shows the temporal series of the spatial weights for 4 selected health
181 zones along with the daily number of COVID-19 cases for the study period. It
182 is evident that increasing weights correspond to increased COVID-19 cases. Simi-
183 larly, Figure 4 shows the flowmap of the median mobility for the week 2021-01-30
184 till 2021-02-05, prepared with the flowmapblue R package ⁵, and the spatial dis-
185 tribution of the spatial weights for the same period.

186 Summarizing, our model is feeded by COVID-19 covariates, socio-demographic
187 covariates and human movement-related covariates. COVID-19 covariates include
188 cumulative cases, average number of cases in neighboring health zones, deaths
189 and average number of deaths in neighboring health zones, and spatial weights
190 computed from the daily mobility matrices and infection. A temporal covariate,
191 day of the week, was computed as a factor from the date.

⁵ <https://github.com/FlowmapBlue/flowmapblue.R>

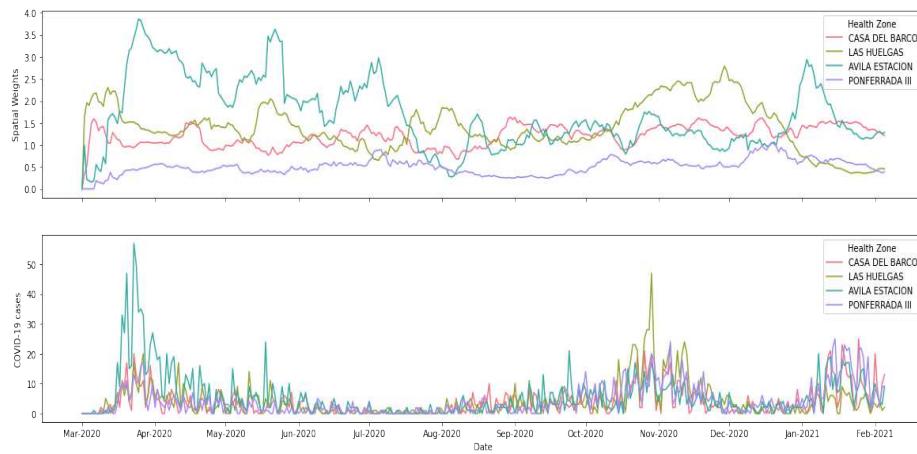


Fig. 3: Spatial weights and COVID-19 cases for the selected health zones

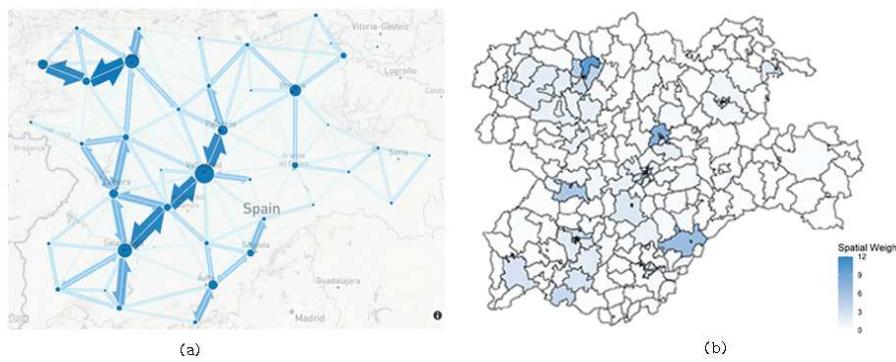


Fig. 4: For the last week of study period 2021-01-30 till 2021-02-05: (a) Flowmap of the study area with the median mobility; (b) Spatial distribution of median values of spatial weights

192 3 A Bayesian LSTM method

193 We use here the term Bayesian LSTM method, to indicate that we use a statisti-
 194 cal model within a Bayesian framework informed by the output of a Long Short
 195 Term Memory (LSTM) neural network method. We aim to model the number of
 196 infections on an areal unit, in our case health zones, based on spatial covariates,
 197 temporal trends, and mobility matrices. Thus our combined model considers tem-
 198 poral and spatial dependence structures, and provides predictions in space and
 199 time of the number of infections.

200 Figure 5 shows a graphical overview of the proposed model which contains
 201 two major components: (a) a deep learning method (LSTM), and (b) a Bayesian
 202 spatial Poisson regression model. The input to the LSTM method are the temporal

series of the cases of infections. The LSTM method learns from these temporal series and predicts the number of cases in the future. Predictions from the LSTM method are embedded into the Poisson regression as an expected value. The spatial correlation structure is modeled using a stochastic partial differential equation (SPDE) method through the INLA approach.

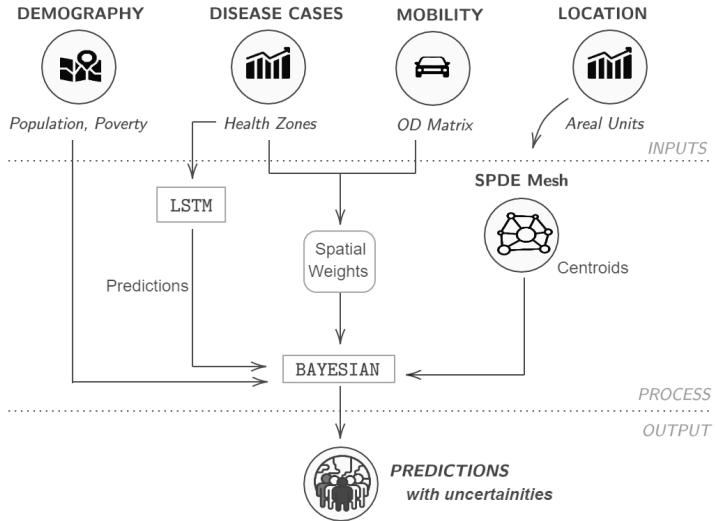


Fig. 5: Graphical overview of the Bayesian LSTM method

3.1 LSTM method

Artificial neural networks are a class of machine learning methods inspired by the functioning of human brain and work on the principle of parallel processing. They consist of layers of interconnected processors known as neurons, which have a vector of weights associated with them. Artificial neural networks models consist of input data also known as input layer, layers of interconnected neurons also known as hidden states, and the output layer which is the output of the model. Fitting an artificial neural network involves estimating the optimal value of these weights which are able to accurately reproduce and mimic some training data. The optimization of these weights is done through the gradient descent method, and the weights assigned to each layer are adjusted proportionally to the derivatives (Bengio et al., 1994).

Among many types of artificial neural networks, recurrent neural networks are arguably the most useful ones for sequential data (as time series) as they have a stack of non-linear units that can learn even long-term dependencies of time series

223 data (Bengio et al., 1994). In recurrent neural networks, the configuration of hidden
 224 states acts as the network memory and the hidden layer state at a time is dependent
 225 on its previous state which enables to learn from past data, thus handling long-
 226 term dependencies (Mikolov et al., 2014). This makes recurrent neural network
 227 an excellent choice for learning and predicting time-dependent data. However,
 228 despite having these advantages, as the recurrent neural networks perform the
 229 gradient descent method with each timestamp of the data, they are likely to fall
 230 into the gradient vanishing problem. Due to this problem, as the recurrent neural
 231 network loops through the networks recurrent connections, the effect of a given
 232 input on hidden layers, and consequently on the output, either decays or explodes
 233 exponentially (Hochreiter, 1991). One alternative approach to tackle this problem
 234 comes from using a LSTM method (Hochreiter & Schmidhuber, 1997), that solves
 235 the gradient vanishing problem by introducing LSTM memory cells instead of the
 236 hidden units. These LSTM cells consist of input, output and forget gates; the
 237 input and output gates are used for the control of the flow of memory cell input
 238 and output into the rest of the model, whereas the forget gates are responsible for
 239 learning the weights that control the rate at which the value stored in the memory
 240 cell decays. With the addition of these gates, the LSTM is able to bypass the
 241 vanishing gradient problem while also learning from the long term dependencies
 242 in the data (Salehinejad et al., 2018).

243 In our case, the LSTM method accounts for the temporal trend of the COVID-
 244 19 spread, learning from the temporal trend of the infected cases on individ-
 245 ual health zones separately, rather than considering the spatial cross-correlation
 246 amongst the regions. Since the LSTM methods are mostly applicable for temporal
 247 series data, it can be assumed that the LSTM method learns more from the tem-
 248 poral trend in the infection on individual health zones separately than from the
 249 spatial relationship between the health zones, which will be further accounted for
 250 in the Bayesian regression model.

251 3.1.1 Architecture

252 We used a four layered LSTM, for which the first layer is the input layer given
 253 by the daily time series of COVID-19. In order to create a supervised learning
 254 problem, the temporal series of infected cases were converted to an input-output
 255 pair which is performed by shifting the data (Brownlee, 2017). Thus, for every
 256 time step t of the time series, one day ahead shifting is done in the data to create
 257 a shifted prediction at $t + 1$. The second layer of the model consists of the 128
 258 LSTM memory cells; similarly, the third and fourth layers consist of 64 and 32
 259 memory cells, respectively. This number of memory cells in each layer comes from
 260 experimentation and also motivated by previous works (Shahid et al., 2020). With
 261 this configuration, the model has 131489 parameters consisting of three stacked
 262 LSTM layers which are recurrently used for the time period T (equal to the total
 263 number of days under study). Finally, a dense layer connects all the recurrent layers
 264 and connects them to the output layer. The dense layer has the linear activation
 265 function. The architecture of the LSTM method is shown in Figure 6. Additional
 266 parameters and hyper-parameters that define the LSTM method are shown in
 267 more detail in Appendix B (Table 6).

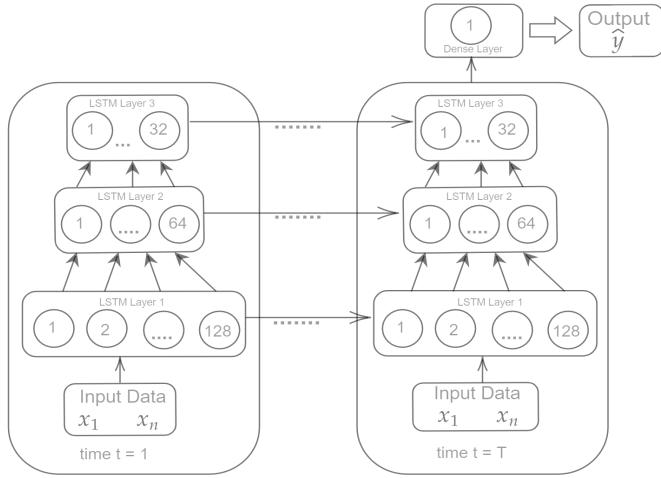


Fig. 6: Architecture of the LSTM method

268 3.2 Spatio-temporal Poisson regression and Bayesian inference

269 To deal with uncertainty, we consider in a second stage a spatio-temporal stochastic
 270 model for the counts of COVID-19 infected cases, which is informed by the output
 271 of LSTM run at a first stage.

Let Y_{it} and E_{it} be the number of observed and expected cases in the i -th area (health zone) and the t -th period (day), $t = 1, \dots, T$. We assume that conditional on the relative risk, ρ_{it} , the number of observed cases follows a Poisson distribution

$$Y_{it} | \rho_{it} \sim Po(\lambda_{it} = E_{it}\rho_{it})$$

272 where, the log-risk is modeled as

$$\log(\rho_{it}) = \beta_0 + Z_{it}^T \beta_{it} + S(x_i) \quad (2)$$

273 with $S(\cdot)$ a spatially structured random effect, and the Z_{it} stand for the covariates
 274 (as mentioned in Section 2). We assigned a vague prior to the vector of coefficients
 275 $\beta = (\beta_0, \dots, \beta_p)$ which is a zero mean Gaussian distribution with precision 0.001.
 276 Finally, all parameters associated to log-precessions are assigned inverse Gamma
 277 distributions with parameters equal to 1 and 0.00005.

278 To compute the joint posterior distribution of model parameters, Bayesian inference
 279 has traditionally relied upon Markov Chain Monte Carlo (MCMC) (Gilks,
 280 1996; Brooks, 2011). This distribution is often in a high dimensional space and thus
 281 it is computationally very expensive. As an alternative computationally faster solution,
 282 Rue et al. (2009) developed a new approximation to the posterior marginal
 283 distributions of model parameters based on a Laplace approximation, and named
 284 it as integrated nested Laplace approximation (INLA). INLA focuses on models
 285 that can be expressed as latent Gaussian Markov random fields (GMRF). In

particular, we use a stochastic partial differential equation (SPDE) method, as introduced by (Lindgren et al., 2011). SPDE consists in representing a continuous spatial process like a Gaussian field (GF) using a discretely indexed spatial random process such as a Gaussian Markov random field (GMRF). In particular, the spatial random process $S(\cdot)$ follows a zero-mean Gaussian process with Matérn covariance function represented as

$$Cov(S(x_i), S(x_j)) = \frac{\sigma^2}{2^{\nu-1} \Gamma(\nu)} (\kappa \|x_i - x_j\|)^{\nu} K_{\nu}(\kappa \|x_i - x_j\|) \quad (3)$$

where $K_{\nu}(\cdot)$ is the modified Bessel function of second order, and $\nu > 0$ and $\kappa > 0$ are the smoothness and scaling parameters, respectively. INLA approach constructs a Matérn SPDE model, with spatial range r and standard deviation parameter σ . The model parameterization is expressed as

$$(\kappa^2 - \Delta)^{(\alpha/2)}(\tau S(x)) = W(x)$$

where $\kappa = \sqrt{8\nu}/r$ is the scale parameter, $\Delta = \sum_{i=1}^d \frac{\partial^2}{\partial x_i^2}$ is the Laplacian operator, $\alpha = (\nu + d/2)$ is the smoothness parameter, τ is inversely proportional to σ and $W(x)$ is a spatial white noise (Blangiardo & Cameletti, 2015). Note that we have $d = 2$ for a two-dimensional process, and we fix $\nu = 1$, so that $\alpha = 2$ in our case.

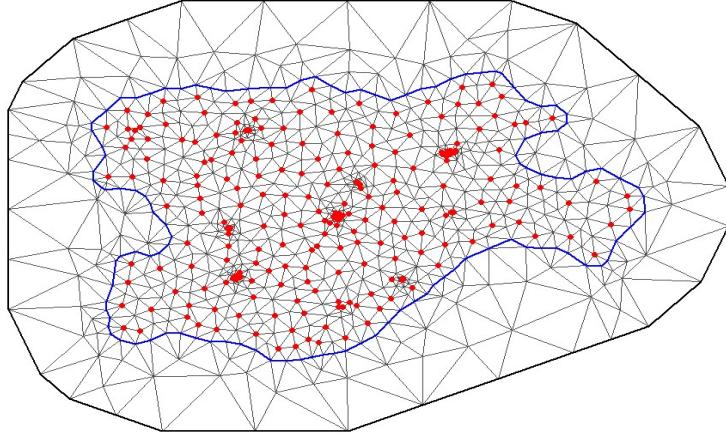


Fig. 7: SPDE triangulation for the study area of Castilla-Leon

We use the centroids of each health zone as the target locations over which we build the mesh. The mesh is formed by smaller triangles within the region of interest, and by larger ones outside the region. The constrained refined Delaunay triangulation is illustrated in Figure 7. The blue line highlights the outline boundary of the study area, with the red dots indicating the centroids of the individual health zones. Note that some few regions show sort of clusters due to the close proximity of health zones. We generate the projection matrix to project the spatially continuous Gaussian random field from the observations to the mesh nodes.

306 Centroids of individual health zones and the triangulations in the mesh are used
 307 to generate the projection matrix. We fixed $r = 0.1$ and $\sigma^2 = 1$. Parameters τ
 308 and κ are renamed as $\theta_1 = \log(\tau)$ and $\theta_2 = \log(\kappa)$, and we assign them zero mean
 309 vague Gaussian independent priors with precisions equal to 0.1.

310 **4 Results**

311 We fitted our Bayesian neural network approach (named as LSTM-INLA through-
 312 out this section) and compared it with two other baseline models, one which is
 313 only using a LSTM method (named as LSTM) and the other one that only fits a
 314 spatial Poisson regression with INLA and no LSTM (named as INLA). We fitted
 315 the models for all the temporal range except for the last week, and used these last
 316 7 days for prediction. The models were evaluated using the averaged Root Mean
 317 Squared Error (RMSE) from all health zones. Additionally, we also considered
 318 the Bayesian metrics Watanabe Akaike information criterion (WAIC) (Watanabe,
 319 2010), deviance information criterion (DIC) (Spiegelhalter et al., 2002) and con-
 320 ditional predictive ordinate (CPO) (Pettit, 1990).

321 Table 4 shows the corresponding metrics, with RMSE evaluated over the train-
 322 ing period (RMSE Training) and over only the prediction period (from 2021-01-30
 323 to 2021-02-05, RMSE Prediction).

Table 4: Metrics for model evaluations

Model	RMSE Training	RMSE Prediction	DIC	WAIC	CPO
INLA	5.33	14.24	373184.93	375164.6	-2.29
LSTM	4.44	6.07	-	-	-
<i>LSTM-INLA</i>	<i>4.14</i>	<i>5.51</i>	<i>354601.13</i>	<i>355510.1</i>	<i>-2.17</i>

*The best model is marked in italics.

324 The RMSE for the LSTM-INLA model is lower than the INLA and LSTM
 325 methods for both the training and prediction periods. We note that although the
 326 RMSE for the training set is quite as good as for the other two methods, the
 327 RMSE for the prediction set for INLA and LSTM is far larger. This suggests that
 328 inclusion of LSTM as an expected value for the spatial Poisson regression plays an
 329 important role. Similarly, the comparison of INLA and LSTM-INLA models with
 330 DIC, WAIC and CPO metrics, shows that the LSTM-INLA combination provides
 331 the best fit. Table 5 shows the correlation between the observed values and the
 332 predicted ones for the prediction period (recall this is the last week of the overall
 333 temporal range). The correlation is largest when using the combined LSTM-INLA
 334 model (0.80) reinforcing the goodness-of-fit of our proposal.

335 Figure 8 depicts the observed cumulative cases of COVID-19 at three selected
 336 weeks within the overall temporal range and chosen at different phases of the
 337 pandemic. We also show the corresponding predictions from the LSTM method and
 338 the combined LSTM-INLA model. In particular, first row of Figure 8 represents
 339 the cumulative number of cases on the initial week of COVID-19 spread in Spain,

Table 5: Correlation between observed and predicted cases (2021-01-30 to 2021-02-05)

Model	Correlation
INLA	0.77
LSTM	0.75
<i>LSTM-INLA</i>	<i>0.80</i>

340 2020-03-22 to 2020-03-28, second row is for the week 2020-10-18 to 2020-10-24, and
 341 third row stands for the 7-days prediction ahead period, from 2021-01-30 to 2021-
 342 02-05. A map depicting the prediction from the LSTM-INLA model and observed
 343 cases for the final week of the study period is published in an R-Shiny app, which
 344 can be accessed through the link⁶. A sample view of the shiny app is presented in
 345 Figure 13 in Appendix A.

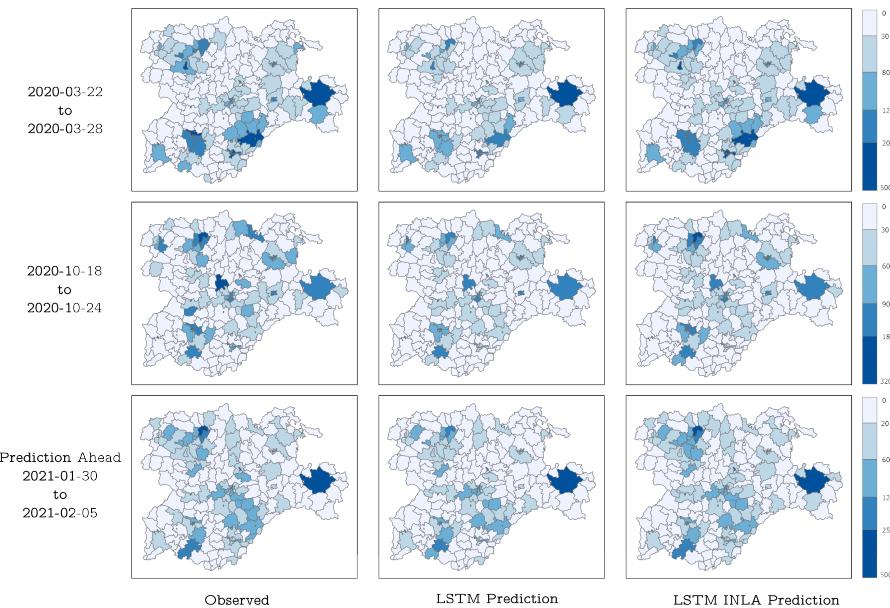


Fig. 8: Spatial distribution of the observed cases (left column) of COVID-19 for three selected weeks. Prediction from the LSTM method (central column) and from LSTM-INLA model (right column)

346 To visualize the temporal trends, Figure 9 shows the observed cases together
 347 with the predicted ones for four selected health zones (Avila Estacion, Las Huel-
 348 gas, Casa del Barco and Ponferrada-II). In particular, we note that we can draw,
 349 together with the predictions under LSTM-INLA, the corresponding 95% credible

⁶ <https://poshan-niraula.shinyapps.io/CYLCovidPrediction/>

350 interval, providing a measure of the uncertainty associated to the prediction, thing
 351 that we can not obtain under LSTM alone. Comparing the prediction from the
 352 LSTM method (green lines), the LSTM-INLA prediction with 95% credible inter-
 353 val (blue lines) with the observed cases (red lines), we note the better prediction
 354 results when using LSTM-INLA. Figure 14 in Appendix D shows the correspon-
 355 ding residual plots. They suggest the better behavior of the LSTM-INLA model as
 356 they are lower in magnitude and symmetrically distributed around the zero line.
 357 This is also true to the prediction ahead case.

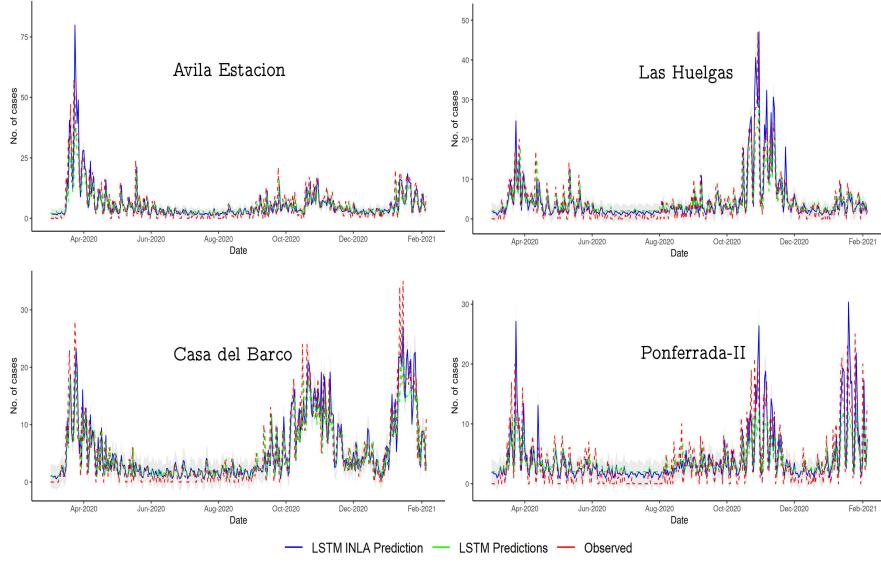


Fig. 9: Temporal trend plots of the observed and predicted cases with LSTM and LSTM-INLA models for four selected health zones. The grey band stands for the 95% credible interval under the LSTM-INLA model

358 Having in mind the model described in equation 2, we now put in place some
 359 information related to the posterior distribution of fixed and random effects. In
 360 particular, Figure 10 depicts the marginal posterior mean and 95% credible inter-
 361 vals of spatial random effect $S(\cdot)$. ID in the X-axis of Figure 10 represents 799
 362 triangulation nodes of the SPDE mesh used in the model. A stronger and signifi-
 363 cative spatial effect is observed basically on the nodes of smaller triangles within
 364 the region of interest (as shown in Figure 7). The nodes outside the region show
 365 no spatial effect.

366 Additionally, Figure 11 and Table 7 in Appendix C depict the marginal pos-
 367 terior distributions of all fixed effects including the intercept (β_0) and the other
 368 covariates. We note that four covariates, namely number of people demanding for
 369 employment, number of commercial offices, number of industrial units and num-
 370 ber of office units in the urban areas, have no influence in our model. The positive

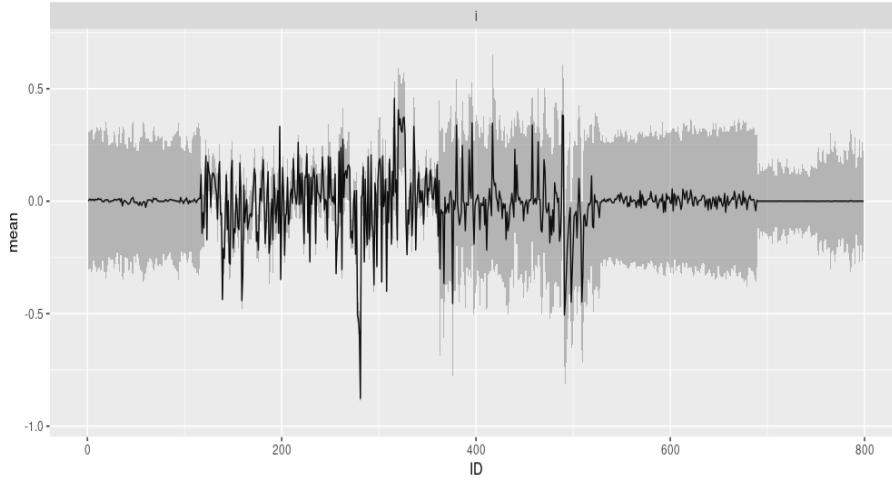


Fig. 10: Marginal posterior mean of the spatial random effect $S(\cdot)$

371 mean values for covariates such as average cases in neighbouring health zones,
 372 cumulative cases, or deaths indicate positive influence in the model. The covariate
 373 associated to daily movement (spatial weight) has the highest positive mean value
 374 which indicates strong positive influence of human mobility on the model.

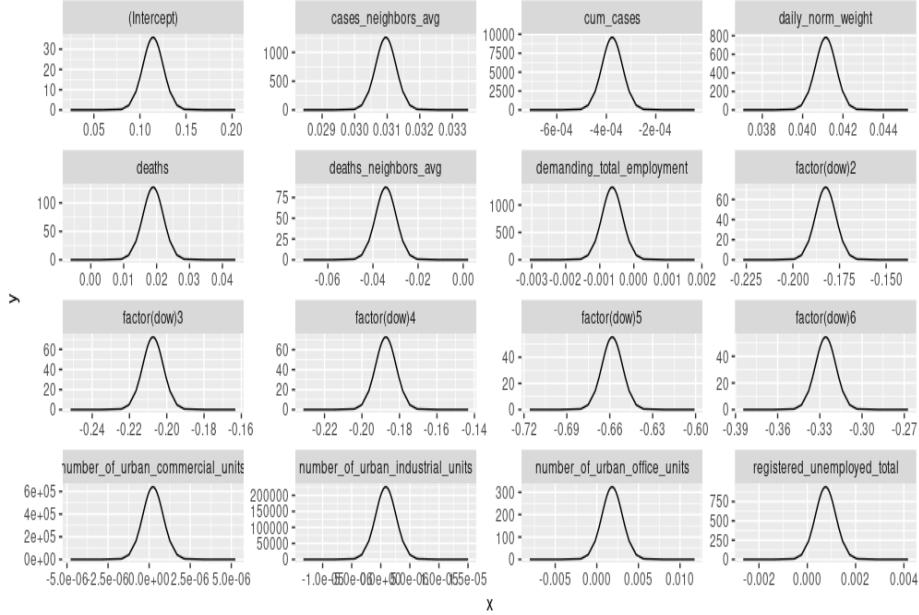


Fig. 11: Marginal posterior distributions of covariate coefficients

375 Finally, Figure 12 shows the marginal posterior Gaussian distributions of the
 376 two hyperparameters for the spatial random field θ_1, θ_2 . Mean and variance for
 377 the two hyperparameters are $\theta_1 = (-3.10, 0.142)$, and $\theta_2 = (3.35, 0.099)$.

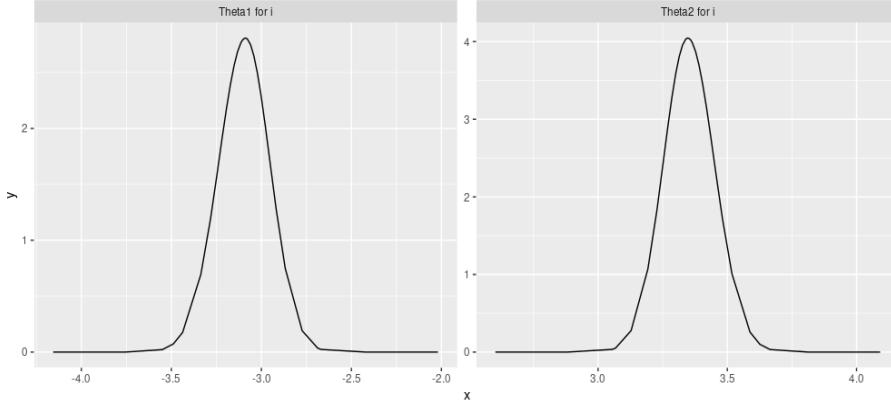


Fig. 12: Hyperparameters θ_1 and θ_2 for the spatial random field $S(\cdot)$

378 5 Conclusions and discussion

379 For modeling the spread and outbreak of infectious diseases, a model comprising
 380 the combination of neural network and Bayesian inference for a spatio-temporal
 381 Poisson regression has been proposed. This model is able to provide good pre-
 382 dictions of further cases of COVID-19 while handling uncertainties. In particular,
 383 our model has two components, a LSTM neural network, which learns from the
 384 temporal patterns, and a spatial Poisson regression with expected values the pre-
 385 dictions coming from the LSTM. The spatio-temporal Poisson regression considers
 386 various spatial and temporal covariates. It is noteworthy that we consider daily
 387 matrices of population movement that are transformed into spatial weights and
 388 act as additional covariates in the model.

389 The proposed model was evaluated with COVID-19 daily infected cases in
 390 Castilla-Leon (Spain), consisting of 245 health zones, and within a temporal range
 391 running from March 1, 2020 to February 5, 2021. The combined model was able
 392 to predict the number of daily infections in each health zone, outperforming two
 393 other cases, one with only a neural network method and the other with only a
 394 spatio-temporal Poisson regression. A key and novel aspect is the introduction
 395 as a spatial weight of the population movement, being highly influential in the
 396 overall fit. However, we note that sudden increasing peaks or abrupt decreasing
 397 magnitudes can not be finely fitted by our model. We believe this is due to typos,
 398 errors or under-reporting actions, and they clearly mean a challenge for modeling
 399 purposes of this sort of data.

400 Clearly, the accuracy of prediction may be improved by the addition of other
401 variables relevant to the disease of study which may include the weather conditions
402 and preventive measures. The phenomenon of infectious disease spread has a lot
403 of complexities and is dependent on numerous factors. These factors include the
404 organism causing the disease, the mode of transmission, human behaviors, envi-
405 ronmental conditions, and most importantly, some potential preventive measures
406 applied. All of these factors are not quantifiable but a maximum number of these
407 factors are to be considered while modeling the diseases. In this study, one of
408 the most relevant considered factors is human mobility. Some socio-demographic
409 variables were considered but we believe more variables associated with the socio-
410 demography and climatic conditions can be introduced. Similarly, the variables
411 related to human behavior and preventive measures such as social distancing and
412 personal hygiene should be incorporated in future works.

413 The focus of this work is on the combination of neural networks and Poisson
414 regression within a Bayesian framework. The predictions from neural networks
415 were used as expected values for the Poisson regression which can be improved
416 by transferring the predictions to a prior distribution and use them as prior in-
417 formation in the Bayesian inference. Here we followed a two-stage procedure, but
418 ideally it would be better a joint solution such as spatio-temporal recurrent neural
419 networks able to predict results with uncertainties. Finally, the proposed method
420 is applied only in one scenario of COVID-19 infection for a short period. Thus,
421 data with a longer period and different spatial scales should be used to test the
422 versatility of the model.

423 The model is believed to be useful for the governments in monitoring any
424 infectious diseases. The results from the model can be used in formulating health-
425 related policies such as the application of preventive measures or vaccination.
426 The contribution of this work is that it is able to take advantage of the neural
427 network methods in learning complex dependencies from the data, as well as from a
428 Bayesian paradigm to associate the uncertainties in the predictions. In conclusion,
429 this work is able to present a model that can provide accurate predictions of
430 infectious diseases and help in a way to mitigate the impacts.

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621 Conflict of interest

622 The authors declare that they have no conflict of interest.

623 Data availability statement

624 The pre-processed data, the Python and R code for implementing the proposed
625 model can be made available upon request.

626 Appendices

627 A Shiny App

Castilla-Leon (Spain) health zones: COVID-19 observed cases and predicted ones

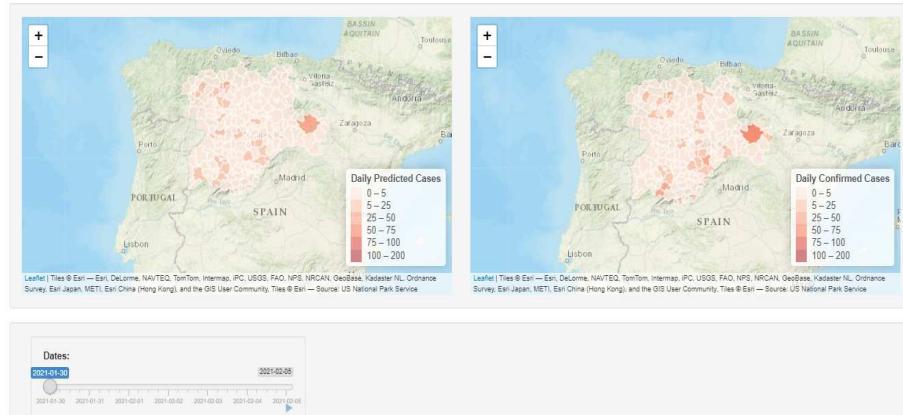


Fig. 13: View of R-Shiny App visualizing observed and predicted COVID-19 cases.
See: <https://poshan-niraula.shinyapps.io/CYLCovidPrediction/>

628 B LSTM method Parameters

Table 6: Summary of parameters and hyperparameters in the LSTM model

Parameter	Value
Number of LSTM layers	3
Hidden Units in LSTM layers	Layer 1: 128 Layer 2: 64 Layer 3: 32
Number of dense layers	1
Activation function of dense layer	Linear
Number of epochs	100
Loss Function	Mean Squared error
Optimizer	ADAM Learning Rate: 0.001 β_1 : 0.9 β_2 : 0.999
Batch Size	10

 629 **C Marginal posterior distributions of covariate coefficients**

Table 7: Marginal posterior mean and credible interval of fixed effects

Covariate	Mean	Credible interval
Monday	-0.182	-0.193, -0.172
Tuesday	-0.207	-0.218, -0.197
Wednesday	-0.187	-0.198, -0.177
Thursday	-0.658	-0.672, -0.644
Friday	-0.326	-0.340, -0.312
Saturday	-0.206	-0.212, -0.192
Sunday	-0.321	-0.336, -0.310
Average cases in neighboring health zones	0.031	0.030, 0.032
Cumulative cases	0.025	0.019, 0.031
Deaths	0.019	0.013, 0.025
Average deaths in neighboring health zones	-0.034	-0.043, -0.025
Daily normal weight (spatial weight)	0.041	0.040, 0.042
Number of people demanding employment	-0.001	-0.001, 0.000
Total registered unemployment	0.001	0.000, 0.002
Number of urban commercial units	0.000	0.000, 0.000
Number of urban industrial units	0.000	0.000, 0.000
Number of urban office units	0.002	-0.001, 0.004

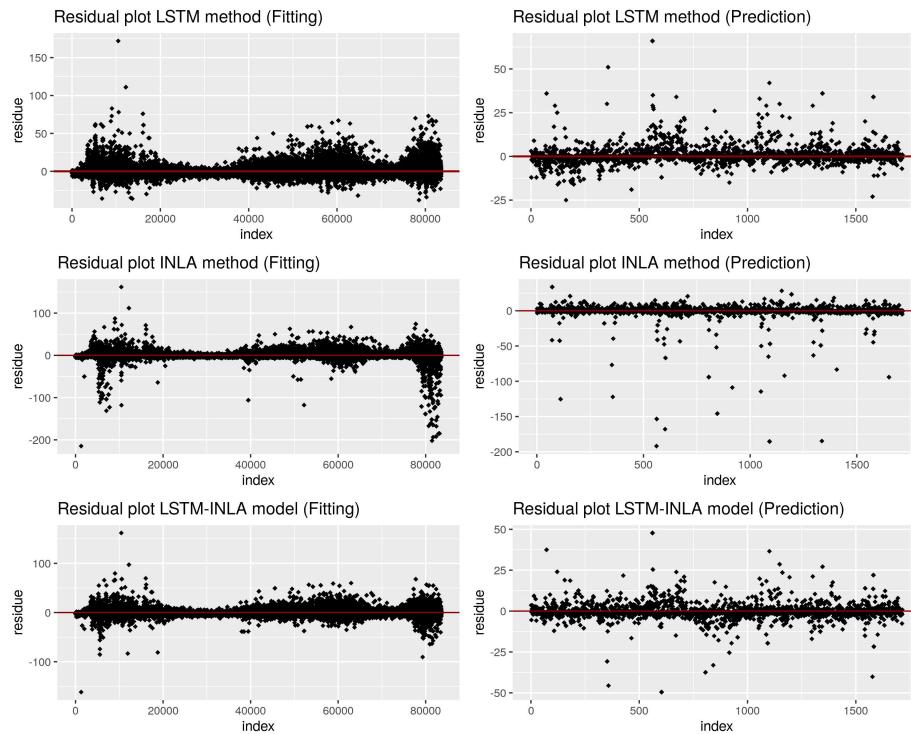
630 D Residual plots of fitted models and their predictions

Fig. 14: Residual plot of the fitted models (left) and predictions (right)