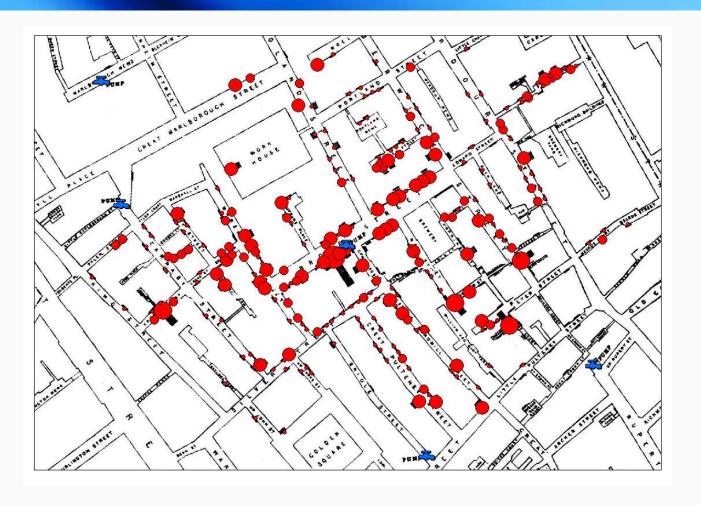
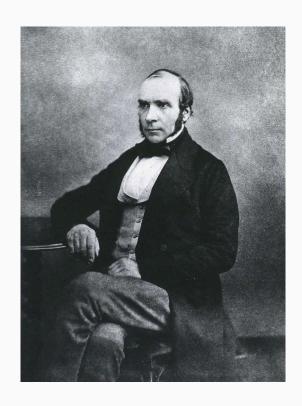


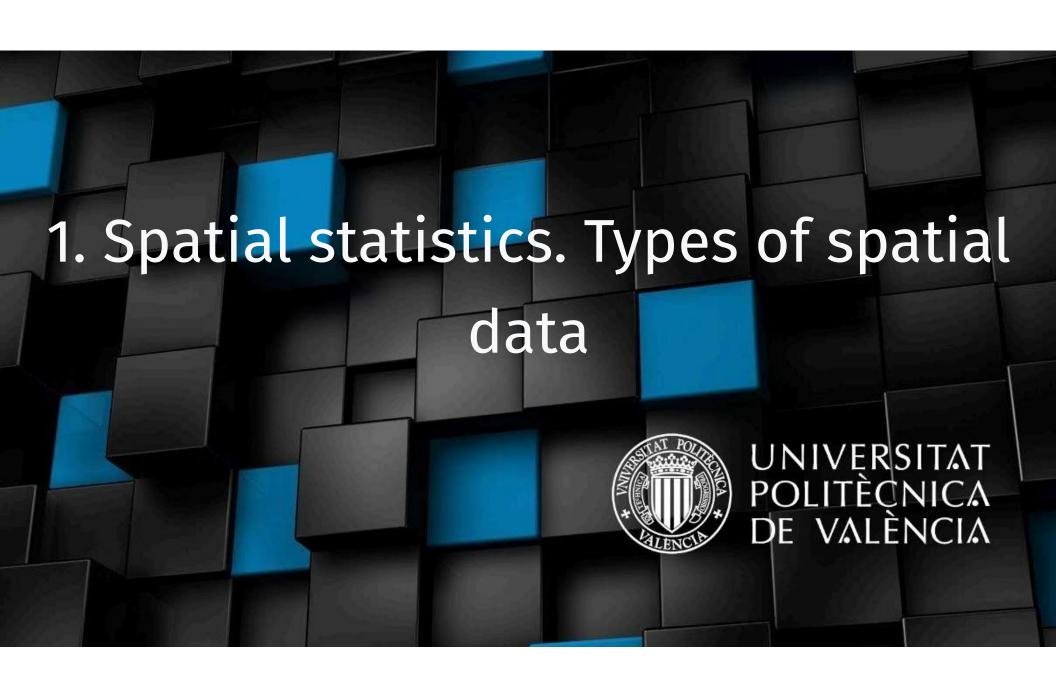
John Snow's Cholera Map in London 1854'





Outline

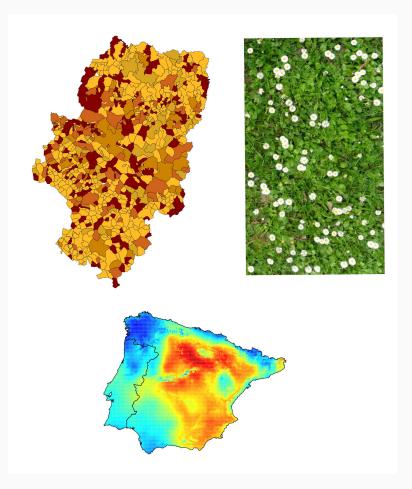
- 1. Spatial statistics. Types of spatial data
- 2. Disease mapping
- 3. Geostatistics
- 4. Penalized complexity priors (PC-priors)
- 5. References

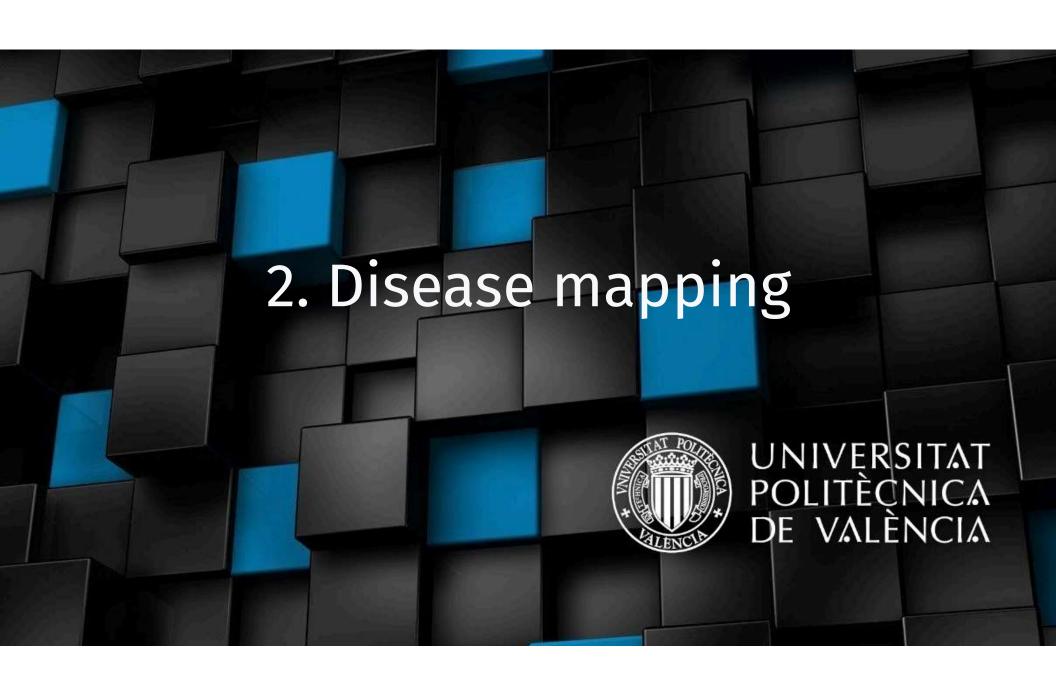


Spatial statistics. Types of spatial data

Spatial statistics is defined as the part of statistics which deal with spatial data and study spatial patterns.

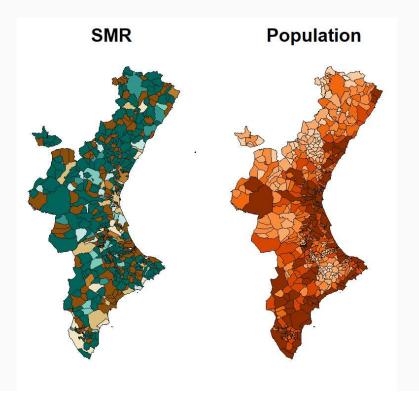
- Lattice or areal data: observations are taken at a finite number of sites whose whole constitutes the entire study region (discrete space), e.g. number of sick people by provinces.
- **Point pattern**: the interest is study the process which generates the points. e.g. distribution of trees in a mountain.
- Geostatistical data: consist of a collection of data in a fixed set locations over a continuous spatial field, e.g. amount of fish in the ocean or presence/absence of a plant in a country.





Oral Cancer mortality in Valencian Region

- In this analysis, we study **oral cancer mortality in the municipalities of the Valencian Region** using a disease mapping model. The aim is to understand the spatial distribution of risks and identify high-risk areas while accounting for variability due to population size and random noise. The variables are:
- **Obs**: the number of observed deaths from oral cancer in the study period.
- **Exp**: the number of expected deaths, based on population size and age-specific rates.
- SMR: the standardized mortality ratio, calculated as ${
 m SMR}={
 m \frac{Obs}{Exp}}\cdot 100$
 - **SMR = 100**: Risk is equivalent to the standard population.
 - SMR > 100: excess risk.
 - SMR < 100: reduced risk.



The model

• A conditional independent **Poisson** likelihood function is assumed:

$$y_i \sim \mathrm{Poisson}(\lambda_i), \;\; \lambda_i = E_i
ho_i, \;\; \log(
ho_i) = \eta_i \;, i = 1, \dots 32$$

• We assume that $\eta_i=eta_0+u_i+v_i$, being $m{u}$ the independent random effect and $m{v}$ the spatially structured random effect:

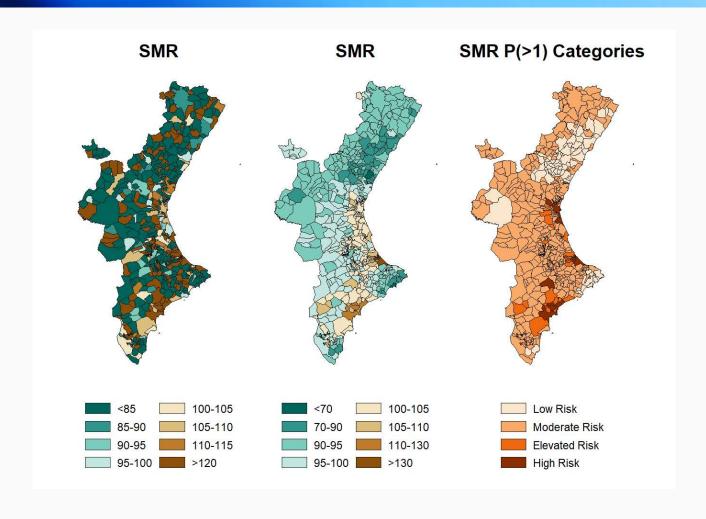
$$u_i \sim \mathcal{N}\left(0, au_{m{u}}^{-1}
ight), \; v_i \mid m{v}_{-i} \sim \mathcal{N}\left(rac{1}{n_i} \sum_{i \sim j} v_j, rac{1}{n_i au_{m{v}}}
ight) \; .$$

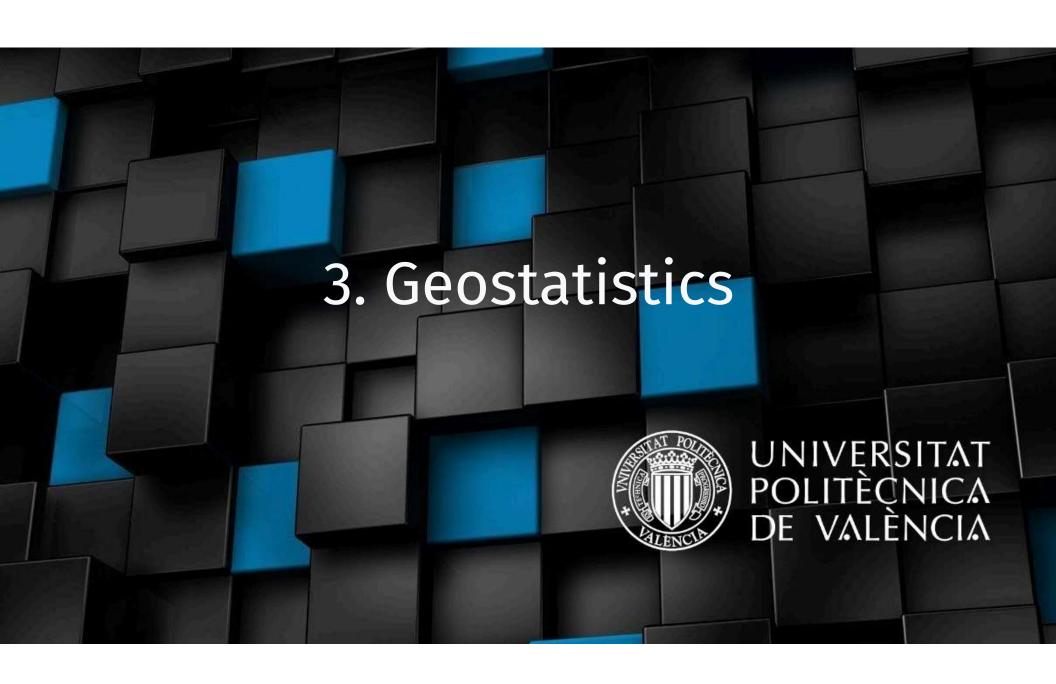
In this case $m{ heta}=(v_1,\ldots,v_{32},u_1,\ldots,u_{32})$, and $m{ heta}\mid m{\psi}$ is Gaussian distributed.

• **Hyperpriors** for the standard deviation parameters σ_u and σ_v follow uniform priors:

$$\sigma_u,\sigma_v \sim \mathrm{Uniform}(0,\infty)$$

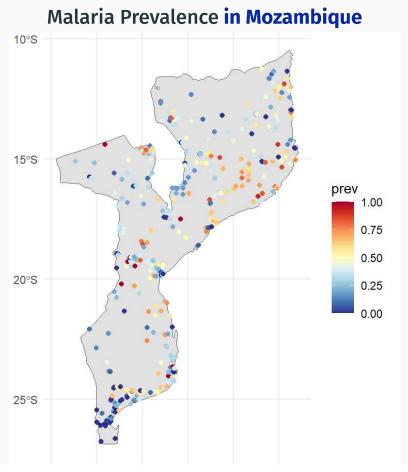
Predicting Risk in Valencia Region





Continuous spaces

- Sometimes, the assumption that the observations have been collected over discrete time points have to be removed.
- The same happen in **space**.
 - If we are studying the presence of a disease,
 pollution in an area or the temperature of a country, the locations where the phenomenon of interest is measured are not frequently allocated in a lattice.
- Then, we are dealing with continuous spaces in 1D and 2D

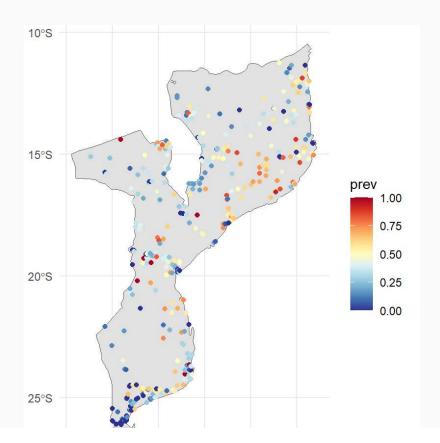


Malaria Prevalence in Mozambique

- This analysis studies **malaria prevalence in Mozambique** using a spatial Bayesian model. The goal is to predict malaria risk and evaluate the effects of environmental and demographic covariates.
- **Examined**: the number of individuals examined for malaria.
- **Positive**: the number of individuals testing positive for malaria.

Covariates:

- **Altitude**: Elevation of the study location (in meters).
- **Temperature**: Average temperature (in °C).
- Proximity to water bodies: Distance to the nearest water source (in kilometers).



Geostatistics. Basis

- Geostatistical models assume that the observations are correlated.
- They are based on the following principle

Everything is related to everything else, but near things are more related than distant things

• So, two close locations tend to **co-vary** more than those far from each other.

Let's be a bit more formal

- A random spatial effect w(s) at a location $s \in \mathcal{D}$ can be considered as a **stochastic process** characterized by a spatial index s which varies continuously in the fixed domain \mathcal{D} , where \mathcal{D} is a fixed subset of r-dimensional Euclidean space.
- The spatial process w(s) is Gaussian if for any $n\geq 1$ and any set of sites $s=\{s_1,\ldots,s_n\}$, $w=\{w(s_1),\ldots,w(s_n)\}$ has a multivariate normal distribution with mean $\mu=E(w(s))$ and a structured covariance matrix Σ . Usually μ is assumed to be ${\bf 0}$. In the literature, this process is widely known as a **Gaussian field (GF)**.
- The key issue in spatial statistics is the covariance function C, which determines the covariance between random variables in two different points. If s_i and s_j are two locations in space, then the **covariance** function is defined as

$$\mathcal{C}(w(s_i), w(s_j)) = Cov(w(s_i), w(s_j))$$

ullet It defines the covariance matrix $oldsymbol{\Sigma}$ of the GF. Each element of the matrix $oldsymbol{\Sigma}_{ij}$ is defined as:

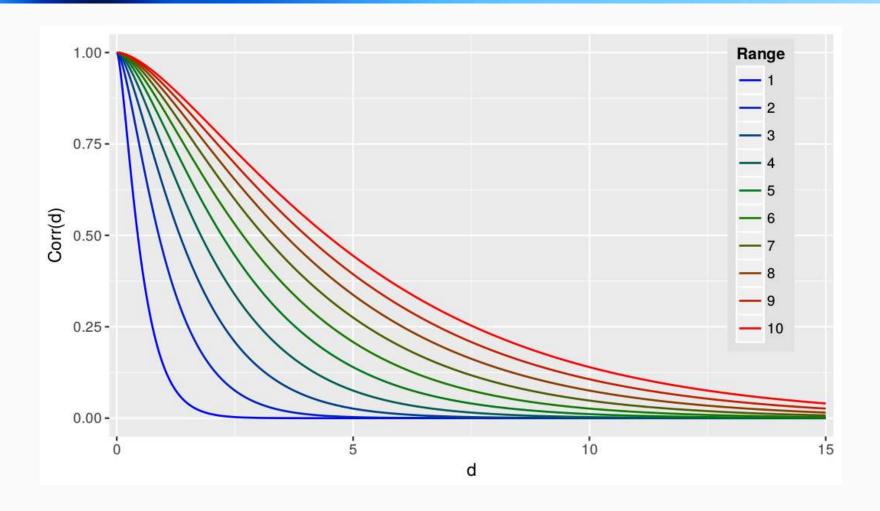
$$\mathbf{\Sigma}_{ij} = \mathcal{C}(w(s_i), w(s_j))$$

Matérn

- Stationarity. We say that the GF is second-order stationary if $\mu(s)=\mu$ and $Cov(w(s),w(s+h))=\mathcal{C}(h)$ for all $h\in\mathcal{R}$ such that s and s+h lie within \mathcal{D} . The covariance function in two different locations depends on the distance vector between these two locations.
 - An example could be the spread of a pathogen in plants. If there is a road close to the crop, maybe this pathogen could spread faster along the road in cars or trucks than in the crop, it would depend on the direction.
- **Isotropy**. We say that the GF is isotropic if the covariance function depends only on the Euclidean distance between points, i.e., Cov(w(s), w(s+h)) = C(||h||).
 - For instance, if we think again in the spread of a pathogen in a crop, it would mean that the spread does not depend on the direction, just on the distance.
- Matérn correlation function is very common.

$$\mathcal{C}(||h||) = \sigma_{m{w}}^2 \left(rac{\sqrt{8}}{\phi}||h||
ight) K_1 \left(rac{\sqrt{8}}{\phi}||h||
ight)$$

Matérn correlation function



Geostatistics in the context of LGMs

Likelihood

A conditional independent Binomial likelihood function is assumed:

$$y_i \mid \pi_i \sim \mathrm{Binomial}(n_i, \pi_i), \; \eta_i = \mathrm{logit}(\pi_i) = eta_0 + eta_1 Temp + w_i \;, i = 1, \dots 447$$

Latent Gaussian field

$$oldsymbol{w} \sim \mathcal{N}(0, oldsymbol{\Sigma}(\sigma_{oldsymbol{w}}, \phi)), \; eta_j \sim \mathcal{N}(0, au = 0.001)$$

 $m{ heta} = (eta_0, eta_1, w_1, \dots, w_{447})$, and $m{ heta} \mid m{\psi}$ is Gaussian distributed.

- $w \sim \mathcal{N}(\mathbf{0}, \mathbf{\Sigma}(\sigma_w, \phi))$, i.e., the spatial effect is assumed to be a **continuous Gaussian field (GF)** with Matérn covariance structure, where:
- $\Sigma(\sigma_w, \phi)$ is a **covariance matrix** depending on the distance between locations, σ_w is the **variance** of the spatial effect, and ϕ is the **range** of the spatial effect.

Hyperparameters $oldsymbol{\psi} = (\sigma_{oldsymbol{w}}, \phi)$

Problem: INLA can not fit continuous GFs

Solution: approximate the continuous GFs using the Stochastic Partial Differential Equation approach (SPDE)

The SPDE approach

Likelihood

$$y_i \mid \pi_i \sim \mathrm{Ber}(\pi_i)$$

$$\operatorname{logit}(\pi_i) = \beta_0 + \beta_1 Temp + w_i$$

Latent Gaussian field

$$oldsymbol{eta} \sim \mathcal{N}(oldsymbol{0}, au = ext{0.0001})$$

$$oldsymbol{w} \sim \mathcal{N}(oldsymbol{0}, oldsymbol{\Sigma}(oldsymbol{\sigma_w}, oldsymbol{\phi}))$$

Hyperparameters

$$p(\sigma_{\boldsymbol{w}}, \phi)$$

Likelihood

$$y_i \mid \pi_i \sim \mathrm{Ber}(\pi_i)$$

$$\operatorname{logit}(\pi_i) = eta_0 + eta_1 Temp + w_i$$

Latent Gaussian field

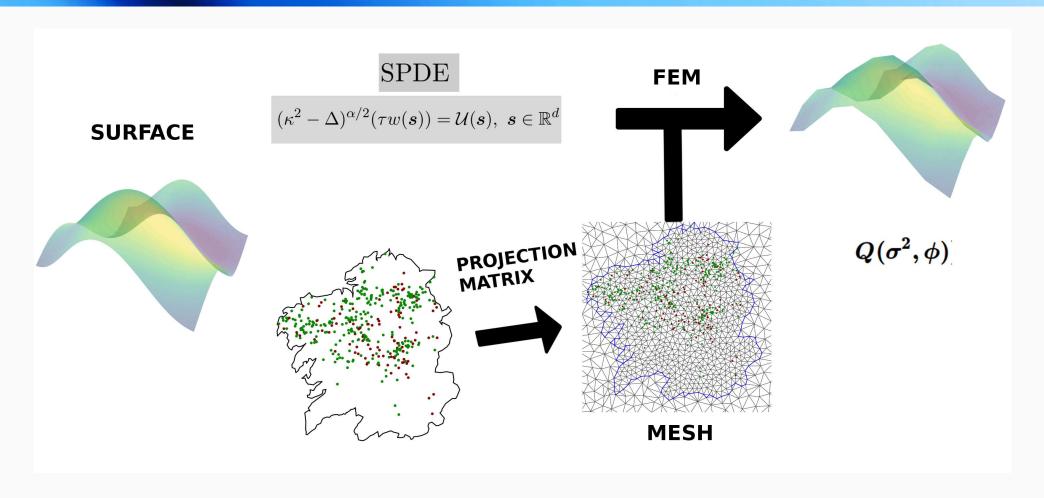
$$oldsymbol{eta} \sim \mathcal{N}(oldsymbol{0}, au = 0.0001)$$

$$oldsymbol{w} \sim \mathcal{N}(oldsymbol{0}, oldsymbol{Q}^{-1}(oldsymbol{\sigma_w}, oldsymbol{\phi}))$$

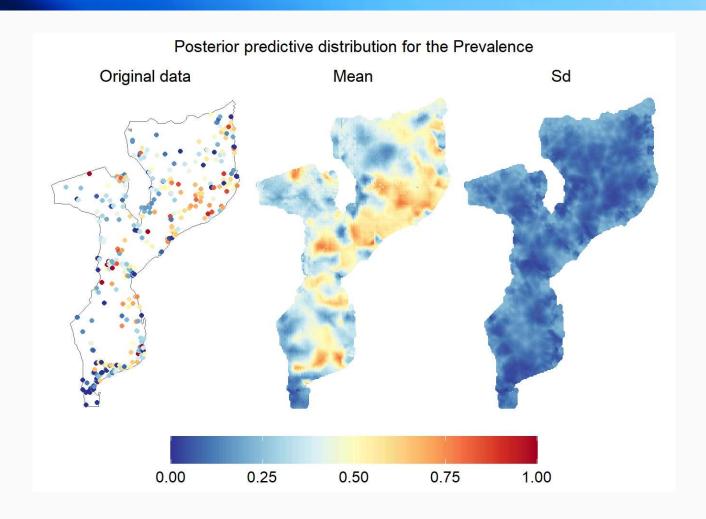
Hyperparameters

$$p(\sigma_{\boldsymbol{w}}, \phi)$$

How is the approximation conducted?



Malaria Prevalence in Mozambique





Penalizing departure from the base model

- Simpson et al. (2017) propose priors that penalize departure from a base model and for this reason they are called **Penalized Complexity (PC) priors**.
- The prior favors the base model unless evidence is provided against it, following the principle of parsimony.
- Distance from the base model is measured using the **Kullback-Leibler** distance, and penalization from the base model is done at a **constant rate on the distance**.
- Finally, the PC prior is defined using **probability statements** on the model parameters in the appropriate scale.

Hyperpriors for the standard deviation in an iid

• The **PC-prior for the precision** au has density:

$$p(au) = rac{\lambda}{2} au^{-3/2} \exp(-\lambda au^{-1/2}) \,, \; au > 0 \,,$$

where

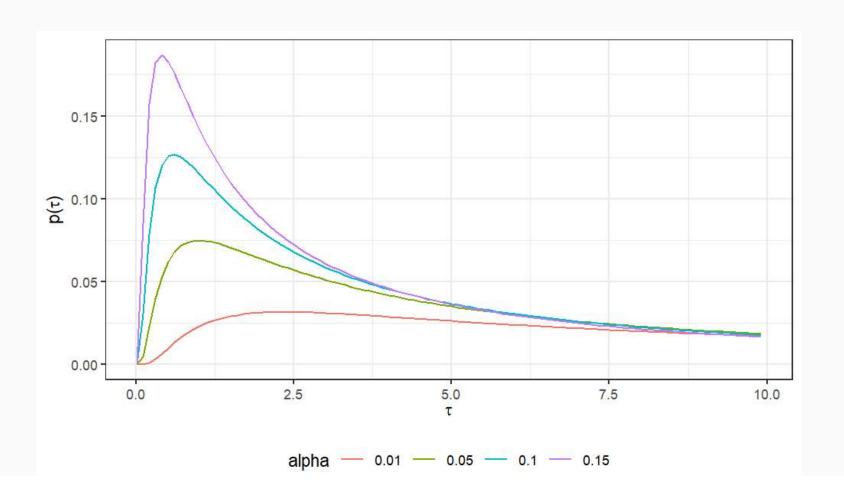
$$\lambda = -rac{ln(lpha)}{u}\,,$$

and (u, α) are the parameters to this prior. The interpretation of (u, α) is that:

$$Prob(\sigma > u) = \alpha$$
, $u > 0$, $0 < \alpha < 1$.

- Functions inla.pc.{d,p,q,r}.prec allow us to **deal with this priors**.
- If we want to plot the prior in terms of the **standard deviation** σ , remember that using function inla.tmarginal we can go from the τ parameter to σ parameter.

Hyperpriors for the standard deviation in an iid.



Spatial effect: priors

ullet The PC-prior for the **range** is defined in terms of ϕ_0 and p_1 so that

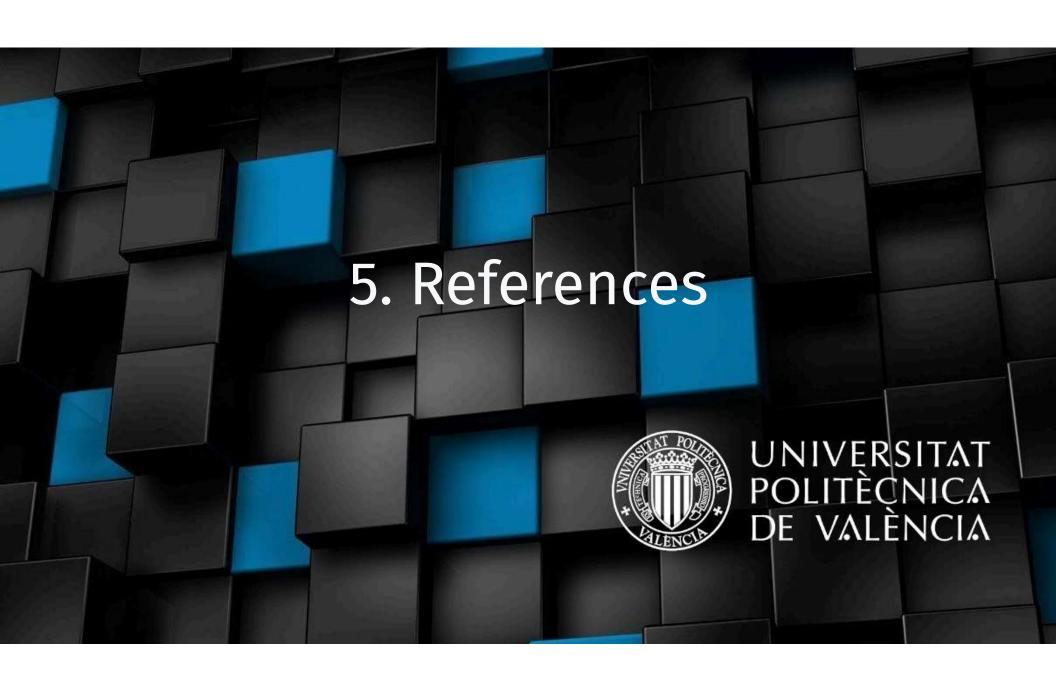
$$Prob(\phi < \phi_0) = p_1$$

ullet The PC-prior for the **standard deviation** is defined in terms of σ_0 and p_2 so that

$$Prob(\sigma_{m{w}} > \sigma_0) = p_2$$

• In order to define the SPDE using PC-priors, the following command have to be used:

```
spde ← inla.spde2.pcmatern(
mesh = ...,
prior.range = c(phi0, p1),
prior.sigma = c(sigma0, p2))
```



This material has been constructed based on:

- Moraga, P., Dean, C., Inoue, J., Morawiecki, P., Noureen, S. R., & Wang, F. (2021). Bayesian spatial modelling of geostatistical data using INLA and SPDE methods: A case study predicting malaria risk in Mozambique. Spatial and Spatio-temporal Epidemiology, 39, 100440.
- Blangiardo, M., & Cameletti, M. (2015). Spatial and spatio-temporal Bayesian models with R-INLA. John Wiley & Sons.
- Fuglstad, G. A., Simpson, D., Lindgren, F., & Rue, H. (2019). Constructing priors that penalize the complexity of Gaussian random fields. Journal of the American Statistical Association, 114(525), 445-452.
- INLA tutorials
- INLA book by Virgilio Gómez-Rúbio
- INLA book by Paula Moraga
- SPDE book by Krainski et al.



Master's Degree in Data Analysis, Process Improvement and Decision Support Engineering

Joaquín Martínez-Minaya, 2024-12-16

VAlencia BAyesian Research Group Statistical Modeling Ecology Group Grupo de Ingeniería Estadística Multivariante

jmarmin@eio.upv.es

