

Session 2.2: Spatial models for small area data: disease mapping and ecological regression

Geospatial analytics using R and R-INLA

MRC
Centre for Environment & Health



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London



Learning Objectives

After this session you should be able to:

- Explain the main ideas underlying the use of Bayesian methods for producing spatially smoothed estimates of disease risk in small areas
- Describe different priors for spatial random effects
- Explore aetiological hypothesis between a health outcome and exposure based on disease mapping
- Describe Poisson regression with spatial random effects for continuous and categorical covariates
- Use R-INLA to produce maps of smoothed estimates of disease risk, carry out spatial smoothing of disease risk and specify ecological regression models

The topics treated in this lecture are covered in:

- Chapter 5-6 of the book **Spatial and Spatio-Temporal Bayesian models with R-INLA**
- Chapter 5 of the book **Geospatial Health Data: Modeling and Visualization with R-INLA and Shiny**
<https://www.paulamoraga.com/book-geospatial/index.html>

Outline

1. Spatial Structure
2. Example: suicides in London
3. Ecological regression with spatial random effects

Spatial patterning

- Poisson-logNormal model is based on the assumption that the observations in the data set are identically distributed and independent
- However, data that occur close together in space (or time) are likely to be correlated
→ **Dependence between observations is a more realistic assumption**
- This spatial patterning, or **spatial autocorrelation**, may be treated as useful information about unobserved influences
- Formally, spatial autocorrelation measures the correlation of a variable with itself through space. If Z_i is the attribute Z observed at location i , and then Z_j is the attribute Z observed at location j , the term spatial autocorrelation refers to the correlation between Z_i and Z_j
- In other words, it quantifies the degree of which observations, at spatial locations, are similar to nearby observations
- Ignoring this dependence can lead to biased and inefficient inference
→ **Smooth in space** prior distribution for the random effects should allow for spatial correlation

Spatial structure

Building a spatial model

- We specify the distribution of each random effect considering the values of the spatial random effects in **neighbouring areas**
- We have a conditional specification since we are conditioning on knowing the neighbours
- We need a rule for determining the neighbours of each area (the most popular one is based on contiguity)
- We specify a conditional autoregressive (CAR) model to capture the spatial structure in the data

Spatial neighbours

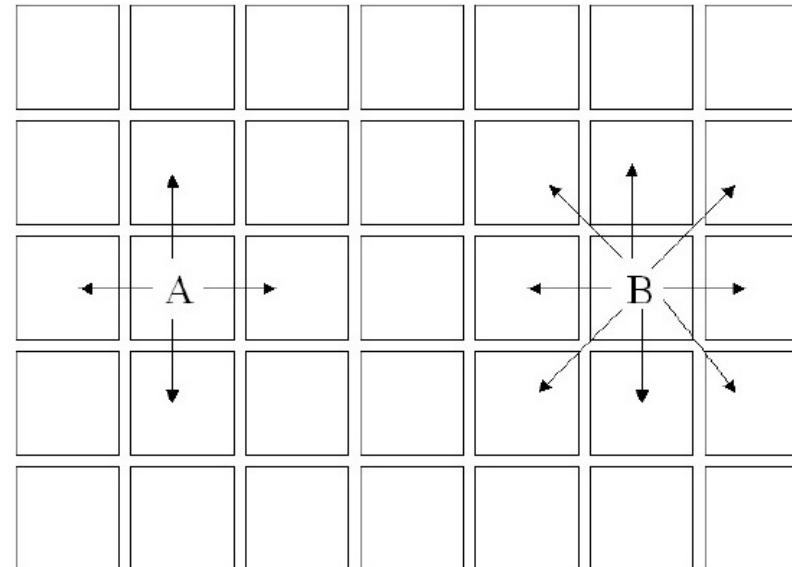
- Key to analyse areal data is the concept of **spatial connectivity** or **spatial proximity**
- Let i and j index two members of the lattice (i.e. two locations such as two countries, or two districts etc.)
- With each pair of sites, we associate a **weight** w_{ij} , so that the spatial weights express the neighbour structure between the observations
- Now, let N be the total number of areal units. The spatial relationship between the areas is represented as an adjacency matrix \mathbf{W} with dimensions $N \times N$, where the entries w_{ij} of the matrix are the spatial weights:

$$\mathbf{W} = \begin{bmatrix} w_{11} & w_{12} & \cdots & w_{1N} \\ w_{21} & w_{22} & \cdots & w_{2N} \\ \vdots & \vdots & \ddots & \vdots \\ w_{N1} & w_{N2} & \cdots & w_{NN} \end{bmatrix}$$

- In its simple form, $w_{ij} = 1$ if areas i and j are adjacent, 0 otherwise
- The diagonal elements of \mathbf{W} are zero, that is $w_{ii} = 0$

Weights based on contiguity

Operationally, we can distinguish between a **rook (A)** and a **queen (B)** criterion of contiguity between areas (in analogy to the moves allowed for the such-named pieces on a chess board):



- The **rook** criterion defines neighbours by the existence of a common edge between two spatial units
- The **queen** criterion defines neighbours as spatial units sharing a common edge or a common vertex

Example of neighbours computation in R

We need to define:

- Neighbour connectivity (who is neighbour?)
- Neighbour weights (how much does the neighbour matter?)
- To do so, we can work with the package `spdep` and we use the function `poly2nb` to define neighbour connectivity according to rook or queen criterion (contiguity neighbours)
- Also, the function `nb2listw` defines spatial weights for neighbours lists, while `nb2mat` defines spatial weights matrices

Example of neighbourhood structure

- We compute a neighbourhood structure for Luxembourg, one of the smallest country in Europe. We use the shapefile for Luxembourg available in the R package raster

```
> library(raster); library(spdep); library(mapview)
>
> # Read in Luxembourg shapefile from R package raster
> lux = shapefile(system.file("external/lux.shp", package="raster"))
> summary(lux)
```

Object of class SpatialPolygonsDataFrame

Coordinates:

	min	max
x	5.74414	6.528252
y	49.44781	50.181622

Is projected: FALSE

proj4string : [+proj=longlat +datum=WGS84 +no_defs]

Data attributes:

ID_1	NAME_1	ID_2	NAME_2	AREA
Min. :1.000	Length:12	Min. : 1.00	Length:12	Min. : 76.0
1st Qu.:1.000	Class :character	1st Qu.: 3.75	Class :character	1st Qu.:187.2
Median :2.000	Mode :character	Median : 6.50	Mode :character	Median :225.5
Mean :1.917		Mean : 6.50		Mean :213.4
3rd Qu.:3.000		3rd Qu.: 9.25		3rd Qu.:253.0
Max. :3.000		Max. :12.00		Max. :312.0

- We plot of Luxembourg divided into 12 cantons and display their name

```
> par(mar=c(0,0,0,0)) # par sets or adjusts plotting parameters,  
> # while the parameter mar stands for margin size  
> plot(lux, border=3, col=terrain.colors(length(lux)), axes=F)  
> text(lux, "NAME_2", cex=0.5)
```



Note that we can also generate a vector of contiguous colors using the functions `rainbow(n)`, `heat.colors(n)`, `terrain.colors(n)`, `topo.colors(n)`, and `cm.colors(n)`.

- We compute contiguity-based neighbors using poly2nb function (here rook's move contiguity is used)

```
> w.rook = poly2nb(lux, row.names=lux$ID_2, queen=FALSE)
> w.rook
```

Neighbour list object:

Number of regions: 12

Number of nonzero links: 46

Percentage nonzero weights: 31.94444

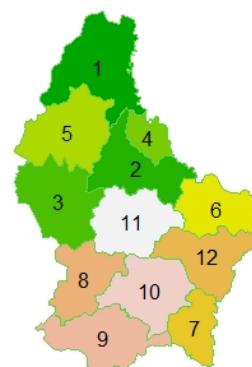
Average number of links: 3.833333

- The nb object w.rook lists for each polygon the neighboring polygons. For example, to see the neighbors for the first polygon in the object, type:

```
> w.rook[[1]]
```

[1] 2 4 5

```
> par(mar=c(0,0,0,0))
> plot(lux, border=3,
+       col=terrain.colors(length(lux)),
+       axes=F)
> text(lux, "ID_2", cex=1)
```



- Finally, we assign the weights to each neighboring polygon. The argument style can take on a number of character values: W=row standardized, B=binary, C=globally standardized. With zero.policy=TRUE we insert zero into the weights matrix where there is no connection

```
> w.rook.l <- nb2listw(w.rook, style="B", zero.policy=TRUE)
> w.rook.l$weights[1] # Check the weight of the first polygon's three neighbors
```

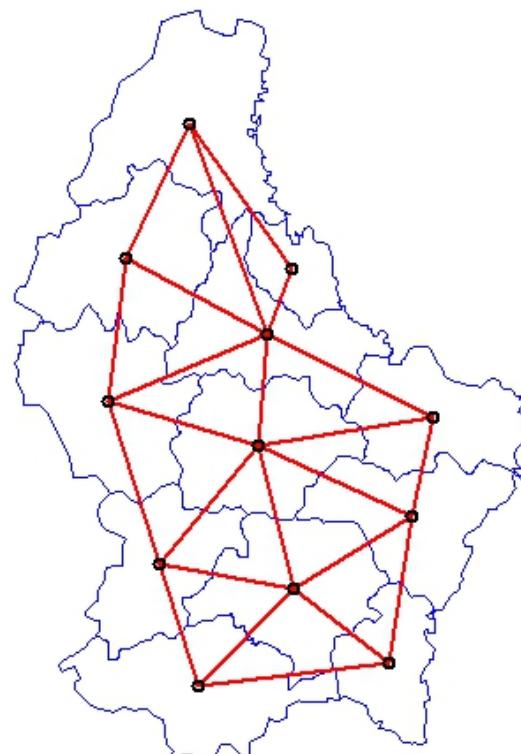
```
[[1]]
[1] 1 1 1
```

```
> w.rook.m <- nb2mat(w.rook, style="B", zero.policy = TRUE) # spatial weights matrix
> w.rook.m
```

	[,1]	[,2]	[,3]	[,4]	[,5]	[,6]	[,7]	[,8]	[,9]	[,10]	[,11]	[,12]
1	0	1	0	1	1	0	0	0	0	0	0	0
2	1	0	1	1	1	1	0	0	0	0	0	1
3	0	1	0	0	1	0	0	0	1	0	0	1
4	1	1	0	0	0	0	0	0	0	0	0	0
5	1	1	1	0	0	0	0	0	0	0	0	0
6	0	1	0	0	0	0	0	1	0	0	0	1
7	0	0	0	0	0	0	0	1	0	1	1	0
12	0	0	0	0	0	1	1	0	0	0	1	1
8	0	0	1	0	0	0	0	0	0	1	1	1
9	0	0	0	0	0	0	1	0	1	0	1	0
10	0	0	0	0	0	0	1	1	1	1	0	1
11	0	1	1	0	0	1	0	1	1	0	1	0

- We now plot the neighbour relationship based on rook move adjacency

```
> # Compute coordinates at centroids of each canton  
> coords <- coordinates(lux)  
>  
> # Plot  
> par(mai=c(0,0,0,0))  
> plot(lux, col='white', border='blue')  
> plot(w.rook, coords, col='red', lwd=2, add=TRUE)
```



Modelling

Intrinsic Conditional Autoregressive (ICAR) model

General definition

Common definition

Remarks

$$\mathbf{u} \sim \text{ICAR}(\mathbf{W}, \sigma_u^2)$$

with

- \mathbf{W} matrix defining the neighbours (weights)
- σ_u^2 conditional variance parameter of \mathbf{u}

$$u_i \mid u_j \quad j \neq i \sim \text{Normal} \left(\frac{\sum_j W_{ij} u_j}{\sum_j W_{ij}}, \frac{\sigma_u^2}{\sum_j W_{ij}} \right)$$

Intrinsic Conditional Autoregressive (ICAR) model

General definition

Common definition

Remarks

$$\mathbf{u} \sim \text{ICAR}(\mathbf{W}, \sigma_u^2)$$

Let $\partial_i = \text{set of areas adjacent to } i$, $w_{ij} = 1$ for $j \in \partial_i$, 0 otherwise

$$u_i \mid u_j \quad j \neq i \sim \text{Normal} \left(\frac{\sum_{j \in \partial_i} u_j}{n_i}, \frac{\sigma_u^2}{n_i} \right)$$

- u_i is smoothed towards mean risk in a set of neighbouring areas
- Conditional variance inversely proportional to the number of neighbours (so more neighbours, less variability)

Intrinsic Conditional Autoregressive (ICAR) model

General definition	Common definition	Remarks
• ICAR model is improper: the overall mean of the \mathbf{u} is not defined. So an additional constraints needs to be imposed: sum-to-zero constraint : $\sum_i u_i = 0$		
• The parameter σ_u^2 represents the conditional variance of the random effects (and not the marginal one) and its magnitude determines the amount of spatial variation		
• No closed-form expression available for the marginal between-area variance of the spatial effects → estimate marginal spatial variance empirically		

$$s_{\text{u.marginal}}^2 = \sum_i (u_i - \bar{u})^2 / (N - 1)$$

Combining ICAR with unstructured random effects

- ICAR model makes a strong spatial assumption; it cannot take a limiting form that allows non-spatial variability
- Besag, York and Mollie (BYM) recommended combining the ICAR prior and the standard normal prior to allow for both
 - spatially unstructured latent covariates v modelled as iid → global smoothing
 - spatially correlated latent covariates u modelled as ICAR → local smoothing

BYM

BYM2

$$\begin{aligned}y_i &\sim \text{Poisson}(\lambda_i = \rho_i E_i) \\ \eta_i &= \log \rho_i = b_0 + v_i + u_i \\ v_i &\sim \text{Normal}(0, \sigma_v^2) \\ \mathbf{u} &\sim \text{ICAR}(\mathbf{W}, \sigma_u^2)\end{aligned}$$

- Need to specify hyperprior distributions for:
- σ_v^2 (between-area unstructured marginal variance), e.g. $1/\sigma_v^2 \sim \text{Gamma}(1, 0.001)$
- σ_u^2 (between-area spatial conditional variance), e.g. $1/\sigma_u^2 \sim \text{Gamma}(1, 0.001)$
- b_0 (mean log relative risk), e.g. $b_0 \sim \text{Normal}(0, 0.0001)$

Combining ICAR with unstructured random effects

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 - spatially correlated latent covariates \mathbf{u} modelled as ICAR → local smoothing

BYM

BYM2

$$\begin{aligned}y_i &\sim \text{Poisson}(\lambda_i = \rho_i E_i) \\ \eta_i &= \log \rho_i = b_0 + b_i \\ \mathbf{b} &= \frac{1}{\sqrt{\tau_b}} (\sqrt{1-\phi} \mathbf{v}_* + \sqrt{\phi} \mathbf{u}_*)\end{aligned}$$

where \mathbf{v}_* and \mathbf{u}_* are standardised versions of \mathbf{u} and \mathbf{v} .

- Need to specify hyperprior distributions for:
 - $\tau_b > 0$, which is the precision parameter controlling the marginal variance of the random effect
 - ϕ , which is the mixing parameter measuring the proportion of the marginal variance explained by the structured effect \mathbf{u}_* . The BYM2 model is equal to an only spatial model when $\phi = 1$, and an only unstructured spatial noise when $\phi = 0$.

Priors for BYM2

Under the BYM2 specification the hyperparameters τ_b and ϕ are modelled using **Penalised Complexity (PC) priors** (Simpson, Rue, Riebler, Martins, and Sorbye, 2017):

- Regularise inference while not forcing too strong information
- Penalise departure from a "base" model (e.g., typically characterised by a fixed value of the relevant parameter)
- Prior tends to favour the base model → need fairly strong evidence to move away from it
- The distance (d) between the **base** model and an **alternative**, more complex model, is measured by the Kullback-Leibler divergence
- The penalization from the base model is done at a constant rate on the distance by assigning an exponential distribution to d :

$$p(d) = \lambda \exp(-\lambda d) \sim \text{Exponential}(\lambda)$$

- PC prior defined using probability statements on the model parameters to determine the value of λ using "reasonable" information.
- For PC priors, see:
 - Simpson, Rue, Riebler et al. (2017)
 - section 5.4 of Gomez-Rubio' book **Bayesian inference with INLA** (<https://becarioprecario.bitbucket.io/inla-gitbook/index.html>)
 - sections 4.3:4.4 of Moraga's book **Geospatial Health Data: Modeling and Visualization with R-INLA and Shiny** (<https://www.paulamoraga.com/book-geospatial/index.html>)

Priors for BYM2

Using probability statements, we can define the PC priors for the two hyperparameters as:

Prior on τ_b

$$P((1/\sqrt{\tau_b}) > U_1) = \alpha_1$$

which can be interpreted as *the probability that the standard deviation of the random effect is larger than U_1 is equal to α_1*

Prior on ϕ

$$P(\phi < U_2) = \alpha_2$$

which can be interpreted as *the probability that the spatial random effect explains less than U_2 of the total variability is equal to α_2*

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$$\text{where } \lambda = \frac{-\log(\alpha_k)}{U_k}$$

- We need to define U_1, U_2 and α_1, α_2 ; following Simpson, Rue, Riebler et al. (2017):

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 - A marginal standard deviation (sd) not too large (e.g. 0.5);

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 - A marginal standard deviation (sd) not too large (e.g. 0.5);
 - $\alpha_1 = 0.01$ (we want to allow for a small probability)

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- We need to define U_1, U_2 and α_1, α_2 ; following Simpson, Rue, Riebler et al. (2017):
 - A marginal standard deviation (sd) not too large (e.g. 0.5);
 - $\alpha_1 = 0.01$ (we want to allow for a small probability)
 - $\alpha_2 = 2/3$ (we expect a higher probability that the variability to be explained by the spatial random effect is lower than 50%)

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- We need to define U_1, U_2 and α_1, α_2 ; following Simpson, Rue, Riebler et al. (2017):
 - A marginal standard deviation (sd) not too large (e.g. 0.5);
 - $\alpha_1 = 0.01$ (we want to allow for a small probability)
 - $\alpha_2 = 2/3$ (we expect a higher probability that the variability to be explained by the spatial random effect is lower than 50%)
- Therefore, following Simpson, Rue, Riebler et al. (2017) the priors are as follows:
 1. Assuming $U_1 = 0.5/0.31$ translates into $P(\sigma_{\tau_b} > 1.62) = 0.01$, using the rule of thumb in Simpson, Rue, Riebler et al. (2017)
 2. Assuming $U_2 = 0.5$ we get $P(\phi < 0.5) = 2/3$

Poisson model with BYM random effects

- Choice of the adjacency matrix (neighbours): 2 areas are neighbours if they share a common border
→ Adjacency matrix implemented in INLA
- An area cannot be specified as its own neighbour
- Adjacency matrix must be symmetric

Poisson model with BYM random effects

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→ Adjacency matrix implemented in INLA
- An area cannot be specified as its own neighbour
- Adjacency matrix must be symmetric
- $\text{RR}_i = \exp(b_0 + b_i)$: RR in area i relative to the age/sex structure (used to estimate the E_i)
- residual $\text{RR}_i = \exp(b_i)$: residual RR in area i relative to the region average after adjusting for the overall risk

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- residual $\text{RR}_i = \exp(b_i)$: residual RR in area i relative to the region average after adjusting for the overall risk
- σ_b^2 reflects the marginal variability of the REs
- ϕ represents the weight of the spatial structure

Example: Suicides in London

Suicides in Greater London, M+F, 1989-1993, Boroughs

- 32 boroughs in Greater London
- Interest: mapping the RR in each borough
- Methods with no spatial structure: SMR, non spatial smoothing
- Spatial smoothing using the BYM model

$$y_i \sim \text{Poisson}(\rho_i E_i)$$

$$\log \rho_i = b_0 + b_i$$

$$v_i \sim \text{Normal}(0, \sigma_v^2)$$

$$\mathbf{u} \sim \text{ICAR}(\mathbf{W}, \sigma_u^2)$$

- Data: \mathbf{y} and \mathbf{E}
- Priors: $\sigma_v^2, \sigma_u^2, b_0$
- Parameters of interest:
 - residual RR ($\text{resRR}_i = \exp(b_i)$)
 - marginal variance ($1/\tau_b$)
 - percent of total variation in the log RR due to spatial effects (ϕ)

Adjacency matrix in INLA

- It is possible to produce a graph from a shapefile
- Upload the shapefile using sf package

```
> library(sf)
> london.gen = st_read("LDNSuicides.shp")
> london.gen$ID = seq(1,32)
```

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> london.gen$ID = seq(1,32)
```

- Use poly2nb and nb2INLA from the spdep package to transform the shapefile into adjacency matrix

```
> library(spdep)
> nb2INLA("LDN.graph",poly2nb(london.gen))
> LDN.adj = paste(getwd(),"/LDN.graph",sep="")
```

Adjacency matrix in INLA

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> nb2INLA("LDN.graph", poly2nb(london.gen))
> LDN.adj = paste(getwd(), "/LDN.graph", sep="")
```

- Now LDN.graph has been saved in the working directory and can be called when specifying the BYM model (see later)

Spatial distributions for area level data in INLA

We introduce here the specification of the ICAR and BYM2 models in INLA, which are done through f():

ICAR in INLA

BYM2 in INLA

```
> formula.ICAR = y ~ f(ID, model="besag", graph=LDN.adj)
```

- ID is the area identifier
- graph=LDN.adj identifies the adjacency structure constructed as seen before
- model=besag specifies the intrinsic conditional autoregressive structure as described before

On the example:

```
> formula = y ~ 1 + f(ID, model="besag", graph=LDN.adj,  
+                         hyper=list(  
+                           prec=list(  
+                             prior="loggamma", param=c(1, 0.0005))))
```

Spatial distributions for area level data in INLA

We introduce here the specification of the ICAR and BYM2 models in INLA, which are done through f():

ICAR in INLA

BYM2 in INLA

```
> formula.BYM2 = y ~ f(ID, model="bym2", graph=LDN.adj)
```

- ID is the area identifier
- graph=LDN.adj identifies the adjacency structure constructed as seen before
- model=BYM2 specifies the combination of the intrinsic conditional autoregressive structure and unstructured random effect as described before

On the example:

```
> formula = y ~ 1 + f(ID, model="bym2", graph="LDN.graph",
+   hyper=list(prec = list(
+     prior = "pc.prec",
+     param = c(0.5 / 0.31, 0.01)),
+     phi = list(
+       prior = "pc",
+       param = c(0.5, 2 / 3))))
```

Running the model in INLA

To run the model in INLA

```
> mod.suicides = inla(formula,family="poisson",
+                      data=data.suicides,E=E,
+                      control.compute=list(dic=TRUE, waic=TRUE))
```

R-INLA estimates the parameters $\theta = \{b_0, \mathbf{b}, \mathbf{u}\}$ and the hyper-parameters $\psi = \{\tau_b, \phi\}$.

How to get information from random effects

- The random effect are obtained through

```
> mod.suicides$summary.random$ID
```

	ID	mean	sd	0.025quant	0.5quant	0.975quant	mode	kld
1	1	-0.08520571	0.10806503	-0.29842674	-0.08479728	0.125710349	-0.08479143	4.431102e-09
2	2	-0.17682950	0.08551914	-0.34571400	-0.17647405	-0.009958857	-0.17647179	6.363872e-09
3	3	-0.21879690	0.09676412	-0.40988824	-0.21839964	-0.029969446	-0.21840105	6.372153e-09
4	4	0.11867926	0.08262032	-0.04260947	0.11836740	0.281743916	0.11836790	5.466778e-09
5	5	-0.14137329	0.08418355	-0.30667107	-0.14136764	0.023888688	-0.14136876	2.925415e-09
6	6	0.40094149	0.08265620	0.23906629	0.40082157	0.563541877	0.40083400	2.468795e-09

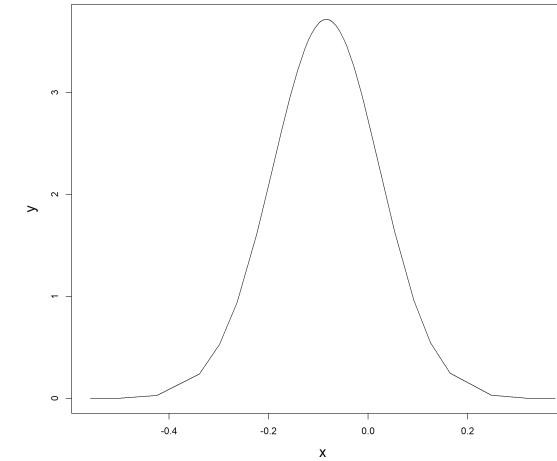
which is a matrix formed by $2n$ rows:

- $1 : n$ rows include information on the area specific residuals b_i
- $n + 1 : 2n$ rows are the spatially structured residual u_i

How to get information from random effects

- All these parameters are on the logarithmic scale; to transform the marginal back to the natural scale:

```
> b = mod.suicides$marginals.random$ID[1:Nareas]
```



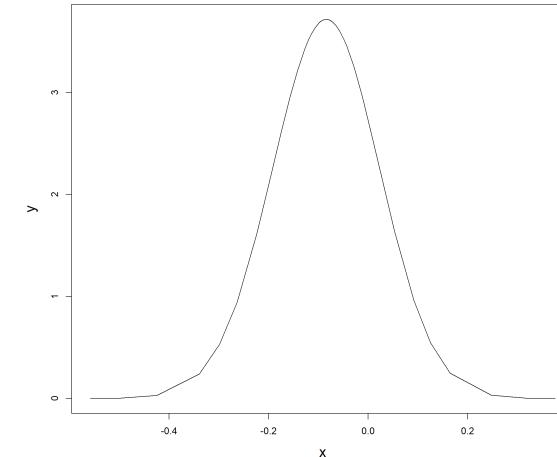
this returns a list with Nareas number of elements,
each representing the posterior marginal of b_i for
that area

How to get information from random effects

- All these parameters are on the logarithmic scale; to transform the marginal back to the natural scale:

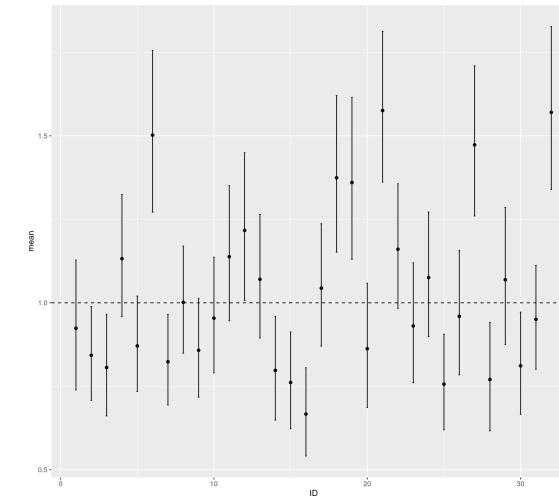
```
> b = mod.suicides$marginals.random$ID[1:Nareas]
```

this returns a list with Nareas number of elements, each representing the posterior marginal of b_i for that area



- Then we can get the posterior mean and 95% credible intervals:

```
> zeta = lapply(b,function(x) inla.emarginal(exp,x))
> zeta_CI = lapply(b,function(x)
+   inla.qmarginal(c(0.025,0.975),
+     inla.tmarginal(exp,x)))
```



Identification of spatial patterns (exceedance probabilities)

- We can now calculate the probabilities that the smoothed RR estimates are greater than a given threshold value
- These probabilities are called **exceedance probabilities** and are useful to assess unusual elevation of disease risk (Richardson, Thomson, Best, and Elliott, 2004)
- The probability that the smoothed RR of area i is higher than a value c can be written as $P(resRR_i > c)$. This can be computed by subtracting $P(resRR_i \leq c)$ to 1, that is:

$$P(resRR_i > c) = 1 - P(resRR_i \leq c)$$

- In R-INLA, we can compute the probability $1 - P(resRR_i \leq c)$ using the `inla.pmarginal()` function with arguments equal to the marginal distribution of the $resRR_i$ and the threshold value c . Then, the exceedance probability $P(resRR_i > c)$ can be computed by subtracting this probability to 1:

```
> 1 - inla.pmarginal(q = c, marginal = marg)
```

where `marg` is the marginal distribution of the residual relative risks, and c is the threshold value.

- For reference, see section 6.6 of Moraga's book <https://www.paulamoraga.com/book-geospatial/index.html>

Posterior probability in R-INLA for suicides in London analysis

- Remember the parametrisation $\zeta = \exp(b_i)$
- We now visualize $p(\zeta_i > 1 \mid \mathbf{y}) = p(b_i > 0 \mid \mathbf{y})$ using the built-in function `inla.pmarginal`:

```
> c = 0  
> prob.b = lapply(b, function(x) {1 - inla.pmarginal(c, x)})
```

Posterior probability in R-INLA for suicides in London analysis

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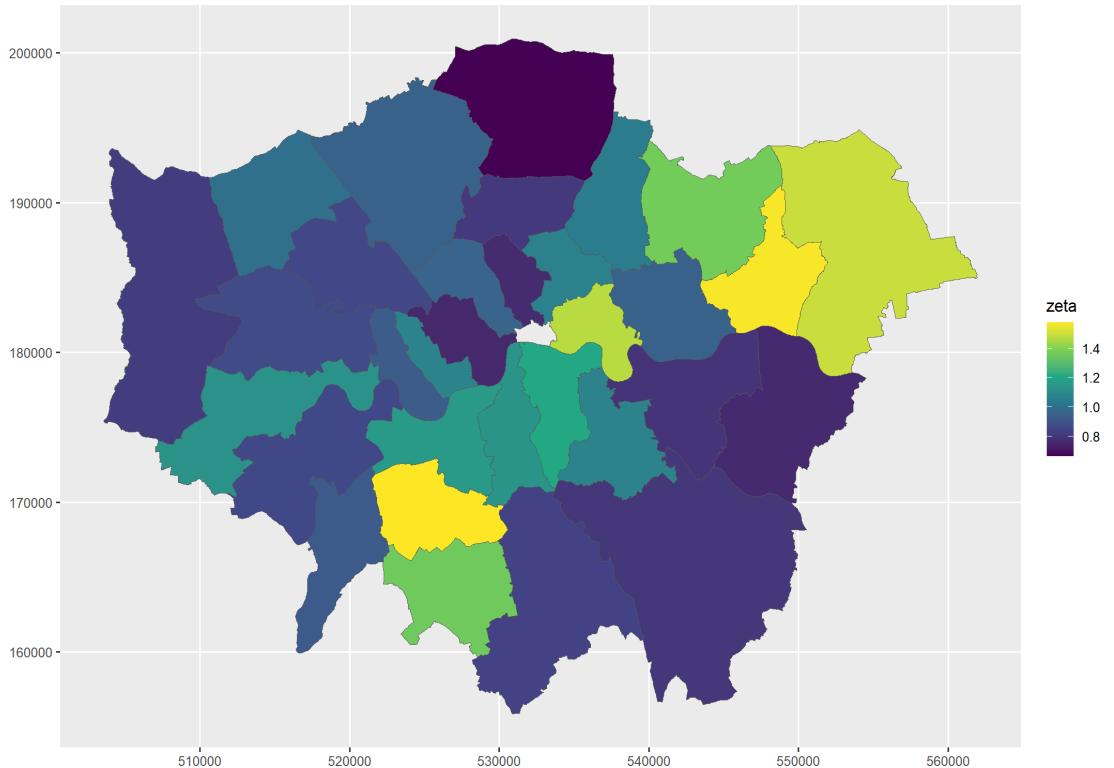
- Create an object with all the info to map

```
> RR_BYM = tibble(zeta=unlist(zeta),prob=unlist(prob.b), ID=seq(1,32))
> out_map = left_join(london.gen,datasuicides, by="ID") %>% left_join(., RR_BYM, by="ID")
> out_map$pp_breaks = cut(out_map$prob,
+                         breaks = c(0, c(0.2, 0.8),
+ 1), include.lowest = T)
```

...and then create some maps

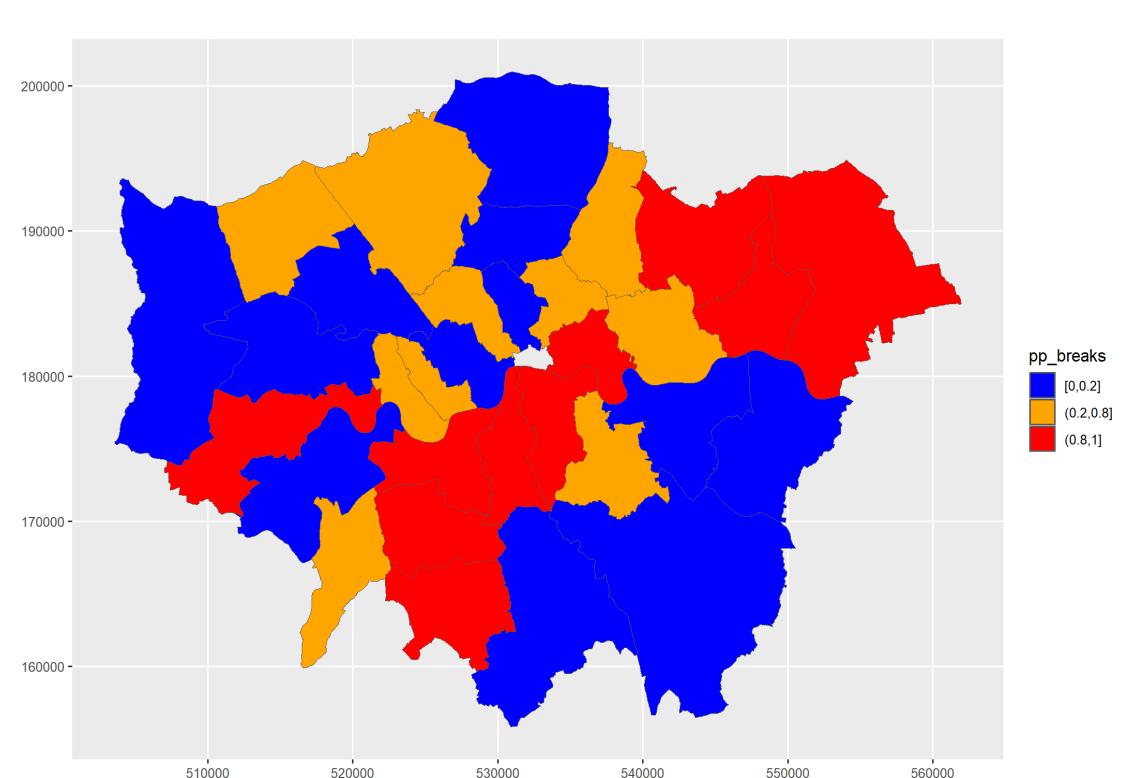
Map of posterior mean of b_i

```
> ggplot() + geom_sf(data = out_map,  
+                      aes(fill = zeta)) +  
+                      scale_fill_viridis_c()
```



Map of posterior probability of $b_i > 0$

```
> ggplot() + geom_sf(data = out_map,  
+                      aes(fill = pp_breaks)) +  
+                      scale_fill_manual(values = c("blue", "orange",
```



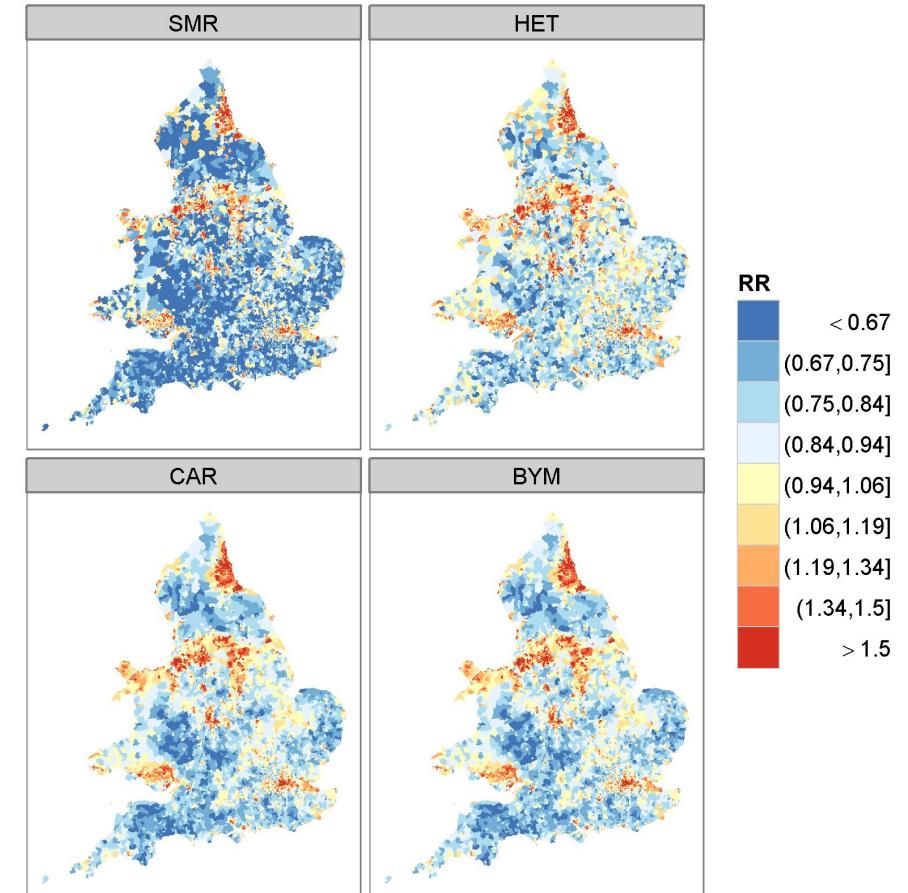
Output from different models

Comparing maps

Shrinkage

Interpretation

- SMR non smoothed RR
- HET non spatially smoothed residual RR: $\exp(v)$
- CAR spatially smoothed residual RR: $\exp(u)$
- BYM spatially and non spatially smoothed residual RR: $\exp(b) = \exp(u + v)$

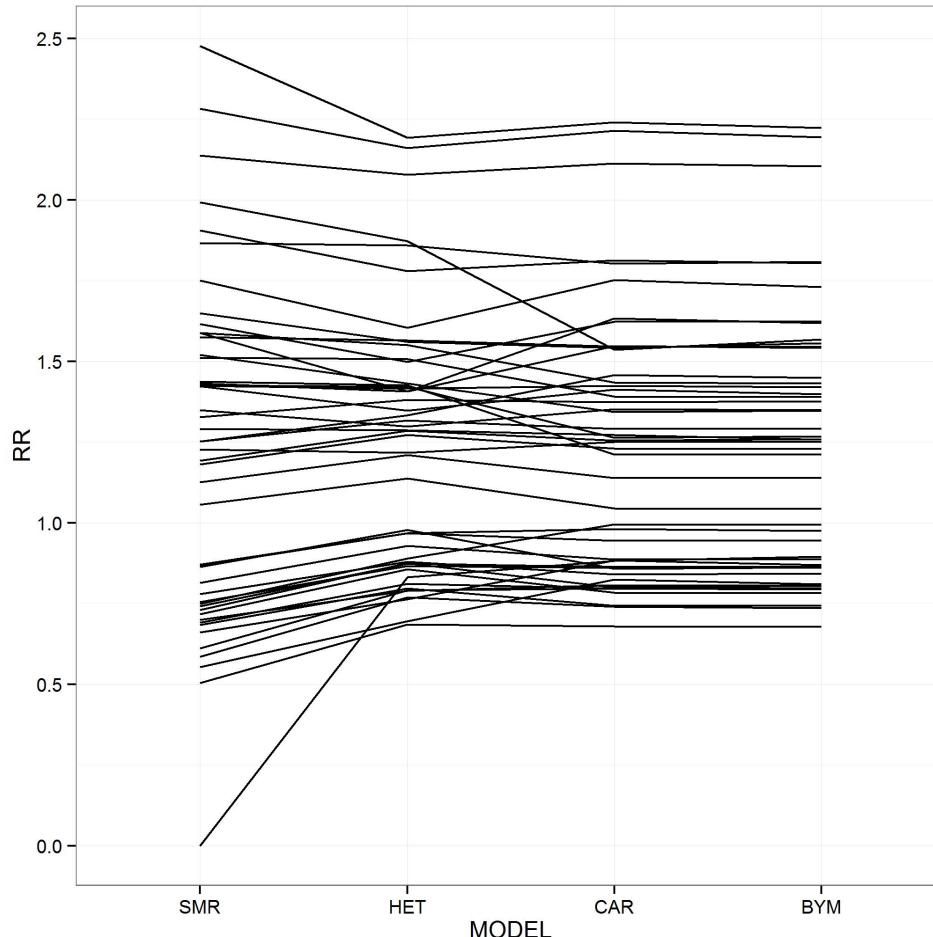


Output from different models

Comparing maps

Shrinkage

Interpretation



- Shrinkage towards the mean due to the **borrowing of strength**

Output from different models

Comparing maps

Shrinkage

Interpretation

- Smoothed relative risks are more stable (precise than observed)

→ geographical patterns of risk are easier to detect using smoothed maps

- Smoothed relative risks have higher specificity:
 - Possible "false positive" values shrunk towards mean
 - But in danger of over-smoothing (false negatives)
 - Visual impact of maps can be very dependent on the choice of colours and cut-points used to shade each region
- Care must be taken not to over-interpret any patterns identified

Ecological regression with spatial random effects (Generalized linear mixed-effect models)

Disease Mapping vs Ecological Regression

Disease mapping studies

- Focus is on description
- Level of inference is at the aggregate (small area) level

Ecological correlation studies

- Focus is on **explanation**
 - Used for investigating specific exposure-disease hypotheses at small-area scale
 - Poisson regression can be used to model relationship between any area-level exposure measure and incidence/prevalence of disease
 - Such area-level exposure measures include average annual pollution level, proportion of population who smoke, proportion of population living with x km of a landfill site, etc.

Poisson regression with random effects

Straightforward extension of disease mapping model:

Ecological regression with BYM structure

$$\begin{aligned}y_i &\sim \text{Poisson}(E_i\rho_i); \quad i = 1, \dots, N \\ \log\rho_i &= b_0 + \beta_1 x_i + u_i + v_i \\ \text{residual RR}_i &= \exp(b_i) = \exp(u_i + v_i) \\ \boldsymbol{b} &= \frac{1}{\sqrt{\tau_b}} (\sqrt{1-\phi} \boldsymbol{v}_* + \sqrt{\phi} \boldsymbol{u}_*) \\ v_i &\sim \text{Normal}(0, \sigma_v^2) \quad \boldsymbol{u} \sim \text{ICAR}(\mathbf{W}, \sigma_u^2)\end{aligned}$$

where

- O_i and E_i : Observed and expected nb of cases in each area i
- λ_i : unknown RR
- x area-level covariate of interest
- β_1 : parameter associated with the covariate
- \boldsymbol{v}_* : standardised version of unstructured random effects, i.i.d.
- \boldsymbol{u}_* : standardised version of random effects with spatial structure, conditional distribution

Interpretation of the parameters

- $\exp(\beta_1)$ is the change in risk associated with a unit change in exposure x
- b_i is the random effect in area i
- $\exp(b_i)$ is the residual or adjusted relative risk of disease in area i
after accounting for the effects of measured covariates and the overall mean risk
- The variance of the random effects reflects the amount of overdispersion in the data (total residual variance = Poisson variance + random effects variance)

Poisson regression with random effects - INLA code

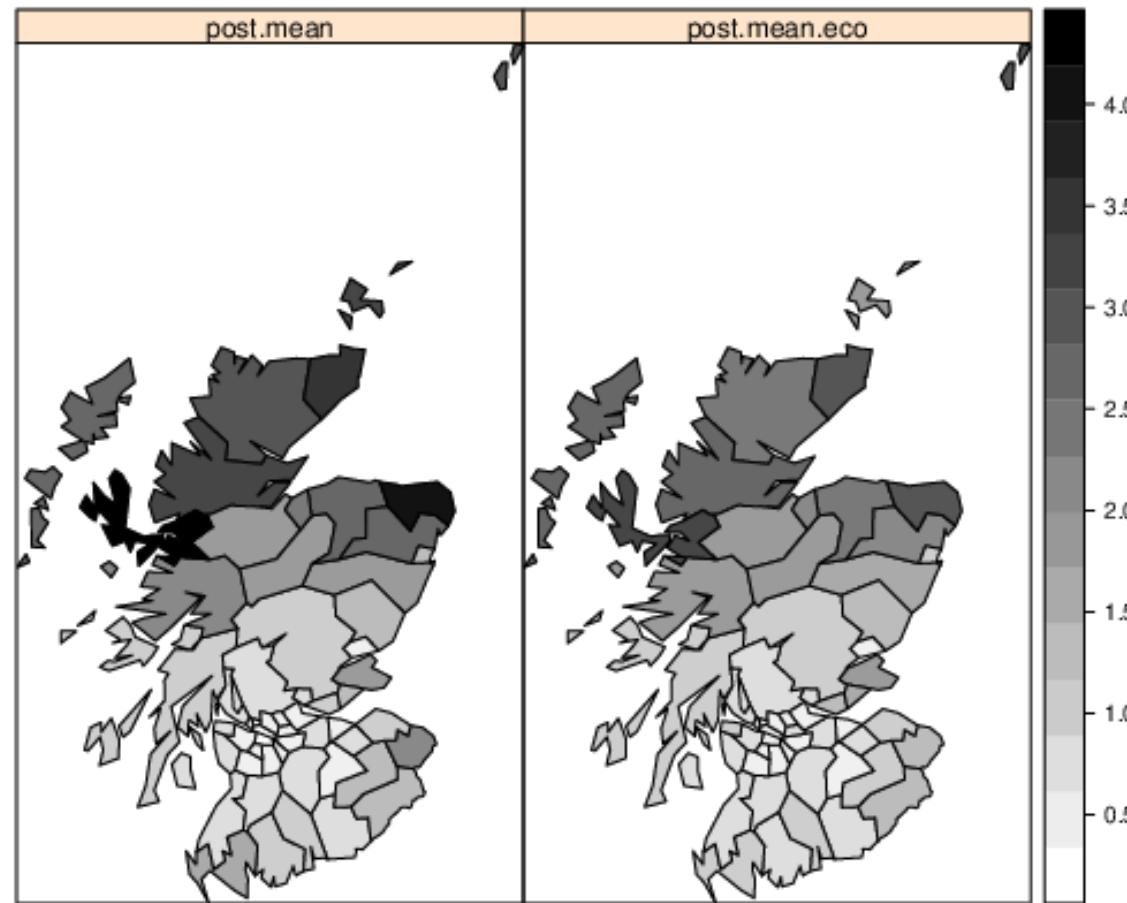
- Continuous covariate

```
> formula.ecoreg.inla = y ~ 1 + x +
+     f(id,model="bym", graph=graph,
+         hyper=list(prec.spatial=list(
+             prior="loggamma",param=c(0.01,0.01))))
```

- Categorical covariate

```
> formula.ecoreg.inla = y ~ 1 + cut(x,breaks=c(0,7,10,24),
+     include.lowest=TRUE) +
+     f(id,model="bym", graph=graph,
+         hyper=list(prec.spatial=list(
+             prior="loggamma",param=c(0.01,0.01))))
```

Comparison between disease mapping and ecological regression



- Less extreme values when covariates are included

→ part of the spatial variability is explained by the covariates

Poisson regression with random effects

Extension to several variables

$$\begin{aligned}y_i &\sim \text{Poisson}(E_i\rho_i); \quad i = 1, \dots, N \\ \log\rho_i &= b_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + b_i \\ &\dots\end{aligned}$$

- $\exp(\beta_1)$ is the relative risk of disease/death associated with a unit increase in exposure x_1 , after adjustment for x_2
- $\exp(\beta_2)$ is the relative risk of disease/death associated with a unit increase in exposure x_2 , after adjustment for x_1
- $\exp(b_i)$ is the residual or adjusted relative risk of disease/death in area i after accounting for the effects of measured covariates and the overall mean risk

Summary

- Hierarchical models allow "borrowing of strength" across units
 - posterior distribution of ρ_i for each unit borrows strength from the likelihood contributions for all the units, via their joint influence on the posterior estimates of the unknown hyper-parameters
- Judgements of exchangeability need careful assessment
 - units suspected a priori to be systematically different might be modelled by including relevant covariates so that residual variability more plausibly reflects exchangeability
 - subgroups of prior interest should be considered separately
- Mapping geographical variations in disease risk is an important epidemiological technique for suggesting aetiological hypotheses
- When combined with data on geographical variations in exposure, disease mapping techniques can be used to investigate and quantify **ecological** associations between disease risk and potential exposures

A few notes on ecological bias and atomistic fallacy

- In the today's session, we worked with aggregated data.
- Aggregation has implications for the type of inference that are possible.
- Ecological inference is the process where aggregated data are used to infer individual level relationships.
Reasons:
 - individual data are not available for confidentiality reasons
 - individual data are not reliable or too expensive to be collected
- **Ecological (or aggregation) bias** can occur. It is the difference between the estimates of relationships obtained using grouped data and those estimates obtained using individual data (e.g. the association observed at the area level do not hold for the individuals within areas).
- The converse, using individual level estimates uncritically to infer group level relationships (ignoring the possibility of group level or contextual effects) is called as **atomistic or individualistic fallacy**.

References

- Richardson, S., A. Thomson, N. Best, et al. (2004). "Interpreting posterior relative risk estimates in disease-mapping studies". In: *Environmental Health Perspectives* 112.9, pp. 1016-1025.
- Simpson, D., H. Rue, A. Riebler, et al. (2017). "Penalising Model Component Complexity: A Principled, Practical Approach to Constructing Priors". In: *Statistical Science* 32.1, pp. 1 - 28. DOI: [10.1214/16-STS576](https://doi.org/10.1214/16-STS576). URL: <https://doi.org/10.1214/16-STS576>.