

# Decomposition of variance for spatial Cox processes

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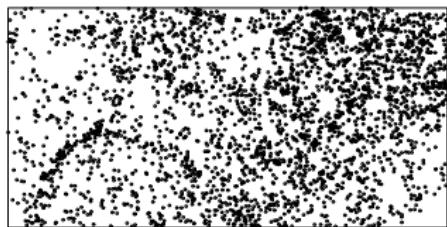
## Background: Tropical rain forest ecology

Fundamental questions: which factors influence the spatial distribution of rain forest trees and what is the reason for the high biodiversity of rain forests ?

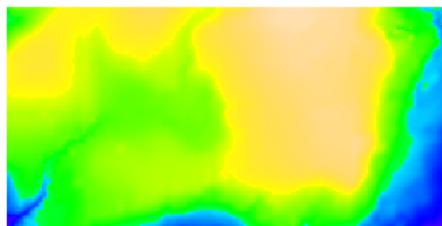
Key factors:

- ▶ environment: topography, soil composition,...
- ▶ seed dispersal limitation: by wind, birds or mammals...

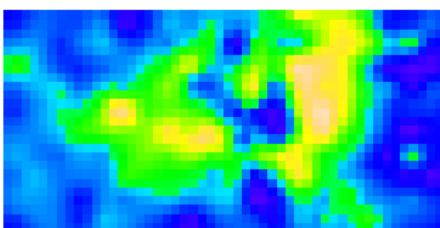
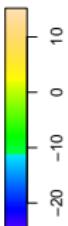
## Example: *Capparis Frondosa*



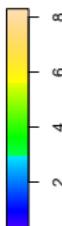
- ▶ *observation window  $W = 1000 \text{ m} \times 500 \text{ m}$*
- ▶ seed dispersal ⇒ *clustering*
- ▶ environment ⇒ *inhomogeneity*



Elevation



Potassium content in soil.

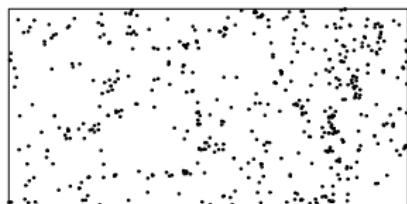


Quantify dependence on environmental variables and seed dispersal using statistics for spatial point processes.

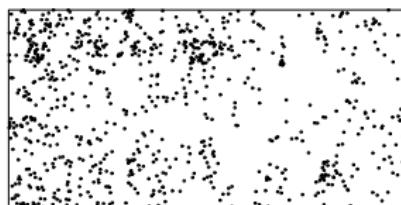
## Example: modes of seed dispersal and clustering

Three species with different modes of seed dispersal:

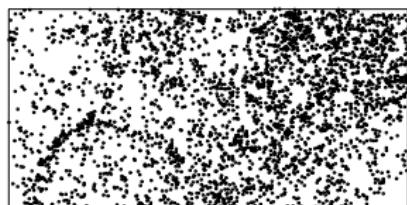
*Acalypha Diversifolia* explosive capsules



*Loncocharpus Heptaphyllus* wind



*Capparis Frondosa* bird/mammal



Is degree of clustering related to mode of seed dispersal ?

Quantify how much of the spatial variation is due to respectively environment and seed dispersal ?

Approach: Cox process model for joint effects of environment and seed dispersal.

## Cox processes

$\mathbf{X}$  is a *Cox process* driven by the non-negative random intensity function  $\Lambda$  if, conditional on  $\Lambda = \lambda$ ,  $\mathbf{X}$  is a Poisson process with intensity function  $\lambda$ .

Intensity function

$$\rho(u) = \mathbb{E}\Lambda(u)$$

Second-order product density

$$\rho^{(2)}(u, v) = \mathbb{E}\Lambda(u)\Lambda(v) = \text{Cov}[\Lambda(u), \Lambda(v)] + \rho(u)\rho(v)$$

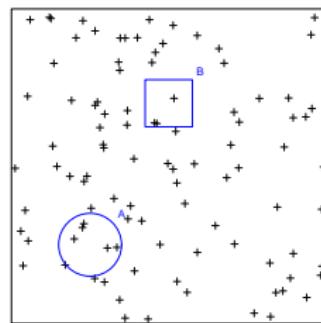
Pair correlation function

$$g(u, v) = \frac{\rho^{(2)}(u, v)}{\rho(u)\rho(v)}$$

# Mean and covariances of counts

$A$  and  $B$  subsets of the plane

$$\mathbb{E}N(A) = \int_A \rho(u)du$$



$$\begin{aligned}\text{Cov}[N(A), N(B)] &= \int_{A \cap B} \mathbb{E}\Lambda(u)du + \int_A \int_B \text{Cov}[\Lambda(u), \Lambda(v)]dudv \\ &= \int_{A \cap B} \rho(u)du + \int_A \int_B \rho(u)\rho(v)[g(u, v) - 1]dudv \\ &= \text{Poisson variance} + \text{extra variance due to } \Lambda\end{aligned}$$

## Decomposition of variance for a count

Prediction of count  $N(B)$  given  $\Lambda$  :

$$\hat{N}(B) = \mathbb{E}[N(B)|\Lambda] = \int_B \Lambda(u)du$$

$\mathbb{V}\text{ar}N(B) =$

variation =

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$$\text{Var}\Lambda(u) = \text{Var}\hat{\Lambda}(u) + \text{Var}[\Lambda(u) - \hat{\Lambda}(u)] \quad \hat{\Lambda}(u) = \mathbb{E}[\Lambda(u)|Z]$$

variation of  $\Lambda$  = variation due to environment + other sources

Measure of influence of environmental covariates  $Z$ :

$$R^2 = \frac{\text{Var}\hat{\Lambda}(u)}{\text{Var}\Lambda(u)} = \frac{\text{Var}\mathbb{E}[\Lambda(u)|Z]}{\text{Var}\Lambda(u)}$$

(right hand side does not depend on  $u$  in case of stationary environment)

Analogy to linear regression:

$$SSR = \mathbb{E} \int_W [\hat{\Lambda}(u) - \rho(u)]^2 du = |W| \text{Var}\hat{\Lambda}(u)$$

$$SST = \mathbb{E} \int_W [\Lambda(u) - \rho(u)]^2 du = |W| \text{Var}\Lambda(u)$$

Then

$$R^2 = \frac{SSR}{SST}$$

## Additive model for $\Lambda$

Additive model:

$$\Lambda(u) = \tilde{Z}(u) + \Lambda_0(u)$$

$\tilde{Z}$  contribution from environment:

$$\tilde{Z}(u) = \beta Z(u)^T \quad Z(u) = (Z_1(u), \dots, Z_p(u))$$

$\Lambda_0$ : structured variation (e.g. seed dispersal) independent of environment.

Cox process **X** union of independent Cox processes with random intensity functions  $\tilde{Z}(u)$  and  $\Lambda_0(u)$ .

Assume  $\tilde{Z}$  and  $\Lambda_0$  non-negative and stationary.

## $R^2$ for additive model

$$R^2 = \frac{\sigma_{\tilde{Z}}^2}{\sigma_{\tilde{Z}}^2 + \sigma_0^2}$$

$$\sigma_{\tilde{Z}}^2 = \text{Var} \tilde{Z}(u) \quad \text{and} \quad \sigma_0^2 = \text{Var} \Lambda_0(u)$$

## Log-linear model

$$\Lambda(u) = \Lambda_0(u) \exp[\tilde{Z}(u)]$$

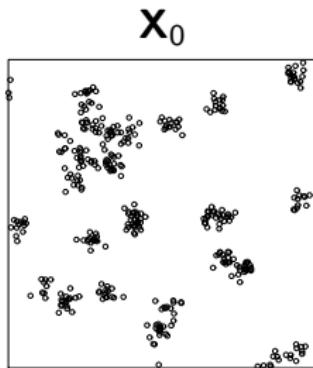
Interpretation in terms of survival of seedlings:

$\mathbf{X}_0$  seedlings: stationary Cox process with random intensity function  $\Lambda_0$ .

$\mathbf{X}$  thinning of  $\mathbf{X}_0$  with survival depending on environment  $\tilde{Z}$ .

$\tilde{Z}$  does not need to be non-negative.

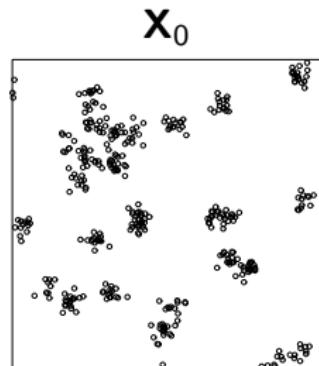
## Example: Cox/cluster process: Inhomogeneous Thomas process



$\Lambda_0$  shot-noise process  $\Rightarrow \mathbf{X}_0$  cluster process:

Offspring distributed according to Gaussian density around Poisson parents

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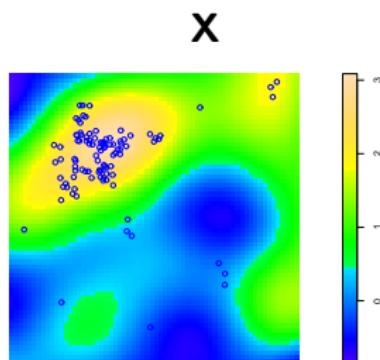
$\Lambda_0$  shot-noise process  $\Rightarrow \mathbf{X}_0$  cluster process:

Offspring distributed according to Gaussian density around Poisson parents

Inhomogeneity: offspring in  $\mathbf{X}_0$  survive according to probability

$$p(u) \propto \exp(Z(u)\beta^T)$$

depending on covariates (independent thinning).



## $R^2$ for log-linear model

$$R^2 = \frac{\sigma_{\exp \tilde{Z}}^2}{\sigma_{\exp \tilde{Z}}^2 + \sigma_0^2 [\sigma_{\exp \tilde{Z}}^2 + \rho_{\exp \tilde{Z}}^2]}$$

$$\sigma_{\exp \tilde{Z}}^2 = \text{Var} \exp[\tilde{Z}(u)]$$

$$\rho_{\exp \tilde{Z}} = \mathbb{E} \exp[\tilde{Z}(u)]$$

$$\sigma_0^2 = \text{Var} \Lambda_0(u)$$

## Models for $c_0(u - v) = \text{Cov}[\Lambda_0(u), \Lambda_0(v)]$

NB: any positive definite function is a covariance function but not necessarily for a non-negative random process  $\Lambda_0$ . Use covariance functions from explicit constructions of  $\Lambda_0$ .

Log-Gaussian:

$$\Lambda_0(u) = \exp[Y(u)] \quad c_0(u - v) = \rho_0^2 \exp[\text{Cov}(Y(u), Y(v))] - \rho_0^2$$

where  $Y$  Gaussian field and  $\rho_0 = \mathbb{E}\Lambda_0(u)$ .

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Shot-noise:

$$\Lambda_0(u) = \sum_{u \in C} \alpha k(u - v) \quad c_0(u - v) = \kappa \alpha^2 \int_{\mathbb{R}^2} k(w) k(w + v - u) dw$$

where  $C$  homogeneous Poisson with intensity  $\kappa$  and  $k(\cdot)$  probability density.

## Matérn class

$\Lambda_0$  shot-noise process: sum of  $K_{(\nu-1)/2}$  Bessel densities  $k(\cdot)$  centered around points of homogeneous Poisson process.

$$c_0(h) = \kappa\alpha^2 \int_{\mathbb{R}^2} k(w)k(w+v-u)dw = \sigma_0^2 \frac{(\|h\|/\eta)^\nu K_\nu(\|h\|/\eta)}{2^{\nu-1}\Gamma(\nu)}$$

With  $\nu = 1/2$  (exponential covariance function)

$$c_0(h) = \sigma_0^2 \exp(-\|h\|/\eta)$$

With ' $\nu = \infty$ ' ('Gaussian' covariance function)

$$c_0(h) = \sigma_0^2 \exp[-(\|h\|/\eta)^2]$$

## Parameter estimation ( $\beta$ )

$\mathbf{X}$  observed within  $W \subset \mathbb{R}^2$

Estimate  $\beta$  and  $\psi = (\sigma_0^2, \eta, \nu)$  using  $\mathbf{X}|Z$ .

First-order composite likelihood:

$$\text{CL}_1(\beta) = \sum_{u \in \mathbf{X} \cap W} \log \rho(u|Z; \beta) - \int_W \rho(u|Z; \beta) du$$

$\rho(\cdot|Z, \beta)$  intensity function for  $\mathbf{X}|Z$ .

For additive model:

$$\rho(u|Z, \beta) = \rho_0 + \beta Z(u)^T \quad \rho_0 = \mathbb{E}\Lambda_0(u)$$

## Estimation of $\psi$

Second-order composite likelihood (given  $\hat{\beta}$ ):

$$\begin{aligned} \text{CL}_2(\psi|\hat{\beta}) = & \sum_{\substack{u,v \in \mathbf{X} \cap W \\ \|u-v\| \leq R}}^{\neq} \log \rho^{(2)}(u, v | Z; \hat{\beta}, \psi) \\ & - \iint_{\|u-v\| \leq R} \rho^{(2)}(u, v | Z; \hat{\beta}, \psi) du dv \end{aligned}$$

$\rho^{(2)}(\cdot, \cdot | Z; \beta, \psi)$  second-order product density for  $\mathbf{X}|Z$ .

For additive model:

$$\rho^{(2)}(\cdot, \cdot | Z; \beta, \psi) = \rho(u | Z, \beta) \rho(v | Z, \beta) + c_0(u - v; \psi)$$

## Environmental variances

$Z$  observed on grid  $G = \{u_i\}_{i=1,\dots,M}$  and  $\widehat{\bar{Z}}(u) = \widehat{\beta}Z(u)^\top$ .

$$\widehat{\sigma}_{\tilde{Z}}^2 = \frac{1}{M} \sum_{u \in G} \left\{ \widehat{\bar{Z}}(u) - \widehat{\rho}_{\tilde{Z}} \right\}^2$$

and

$$\widehat{\sigma}_{\exp \tilde{Z}}^2 = \frac{1}{M} \sum_{u \in G} \left\{ \exp [\widehat{\bar{Z}}(u)] - \widehat{\rho}_{\exp \tilde{Z}} \right\}^2$$

where

$$\widehat{\rho}_{\tilde{Z}} = \frac{1}{M} \sum_{u \in G} \widehat{\bar{Z}}(u) \quad \widehat{\rho}_{\exp \tilde{Z}} = \frac{1}{M} \sum_{u \in G} \exp [\widehat{\bar{Z}}(u)]$$

## Results for rain forest data

Covariates elevation and potassium for Acalypha and Capparis.  
Nitrogen and phosphorous for Lonchocarpus.

Qualitative similar results for additive and log-linear model regarding dependence on covariates ( $\beta$ ).

Species	model for $\Lambda$	$c_0$	$CL_2(\hat{\psi} \hat{\beta})$	$R^2$
Acalypha	log-linear	'Gaussian'	-1239.8	0.01
		Matérn	-1221.0	0.02
		LG-Matérn	-1204.6	0.01
	additive	'Gaussian'	-1641.7	0.01
		Matérn	-1623.4	0.01
		LG-Matérn	-1623.4	0.01

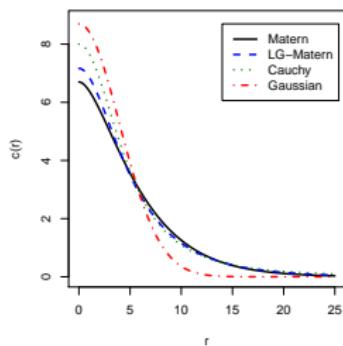
LG-Matérn:  $c_0(h) = \rho_0^2[\exp(c(h)) - 1]$  where  $c(\cdot)$  Matérn.

## Results continued

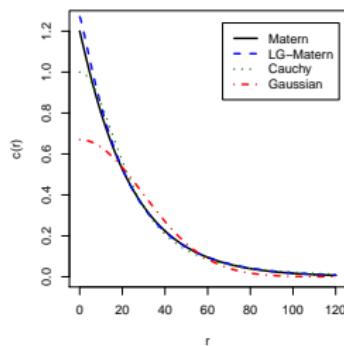
Species	model for $\Lambda$	$c_0$	$CL_2(\hat{\psi} \hat{\beta})$	$R^2$
Loncocharpus	log-linear	'Gaussian'	-3204	0.17
		Matérn	-3156	0.10
		LG-Matérn	-3153	0.10
	additive	'Gaussian'	-4081	0.11
		Matérn	-4055	0.07
		LG-Matérn	-4055	0.07
Capparis	log-linear	'Gaussian'	-76657	0.37
		Matérn	-76325	0.19
		LG-Matérn	-76311	0.19
	additive	'Gaussian'	-81139	0.23
		Matérn	-80685	0.16
		LG-Matérn	-80685	0.16

# Fitted covariance functions (log-linear model)

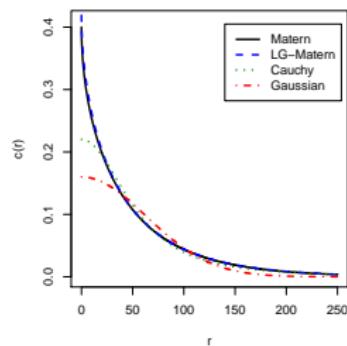
Acalypha



Loncocharpus



Capparis



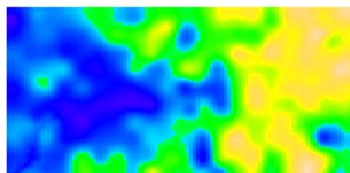
## Some conclusions

- Estimated  $R^2$  sensitive to choice of model: especially  $c_0$ .
- Best fit with log-linear model (interpretation in terms of survival).
- Best fit with (LG)-Matérn (heavy tails for covariance/cluster density).
- Largest  $R^2$  for Capparis (bird/mammal seed dispersal), smallest for Acalypha (explosive capsules).

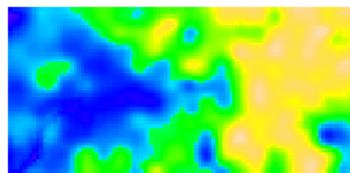
# Stationary environment ?

$\hat{\tilde{Z}}$ :

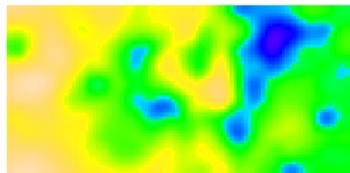
Acalypha (additive)



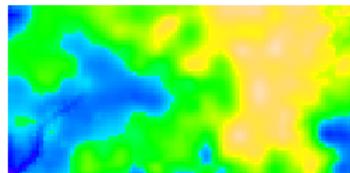
Acalypha (log-linear)



Loncho (log-linear)



Capparis (log-linear)



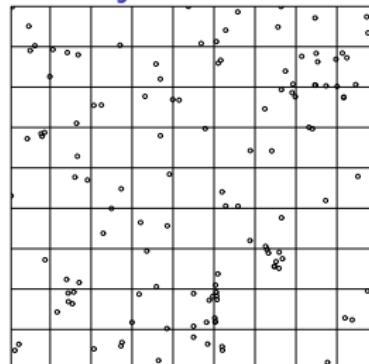
Handling possible non-stationarity topic for further research.

## Composite likelihood obtained from binary random field

Random count variables:  $N_i = \#\mathbf{X} \cap C_i$   
number of points in  $C_i$ .

Disjoint subdivision  $W = \cup_{i=1}^n C_i$  in  
'cells'  $C_i$ .

$X_i = 1[N_i > 0]$  binary random variable.  
 $P(X_i = 1) = \rho(u_i; \beta)|C_i|$ .



Bernoulli composite likelihood

$$\prod_{i=1}^n P(X_i = 1)^{X_i} (1 - P(X_i = 1))^{1-X_i} \equiv \prod_{i=1}^n \rho(u_i; \beta)^{X_i} (1 - \rho(u_i; \beta)|C_i|)^{1-X_i}$$

has limit ( $|C_i| \rightarrow 0$ )

$$\text{CL}_1(\beta) = \sum_{u \in \mathbf{X} \cap W} \log \rho(u; \beta) - \int_W \rho(u; \beta) du$$

Second-order composite likelihood obtained from binary variables associated with joint presence of points in pairs of cells  $C_i$  and  $C_j$ :

$$X_{ij} = 1[N_i > 0, N_j > 0] \quad (= N_i N_j \text{ when } C_i \text{ and } C_j \text{ small })$$

where

$$P(X_{ij} = 1) = \mathbb{E}N_i N_j = \rho^{(2)}(u, v; \beta, \psi) |C_i||C_j|$$

## Alternative: GEE

Observation vector

$$Y = [N_1, \dots, N_m]$$

Mean vector

$$\mu = [\rho(u_1)|C_1|, \dots, \rho(u_m)|C_m|]$$

Working covariance matrix

$$V = [V_{ij}]_{ij}$$

$$V_{ii} = \text{Var}N_i = \rho(u_i)|C_i| + \rho(u_i)^2[g(u_i, u_i; \psi) - 1]$$

$$V_{ij} = \text{Cov}[N_i, N_j] = \rho(u_i)\rho(u_j)[g(u_i, u_j; \psi) - 1]$$

where  $g(\cdot; \psi)$  working pair correlation function.

## GEE

$$[Y - \mu] V^{-1} D \quad D = \frac{d\mu^T}{d\beta}$$

has limiting form (using Neuman series for  $V^{-1}$ )

$$\begin{aligned} & \sum_{u \in \mathbf{X} \cap W} \frac{\rho'(u; \beta)}{\rho(u)} - \int_W \rho'(u; \beta) \frac{\rho(u; \beta)}{\rho(u)} du \\ & + \sum_{v_0 \in \mathbf{X} \cap W} \frac{1}{\rho(v_0)} \sum_{k=1}^{\infty} \int_{W^k} \prod_{l=1}^k (\rho(v_{l-1}) [g(v_{l-1}, v_l) - 1]) \rho'(v_k; \beta) dv_1 \cdots dv_k \\ & - \int_W \frac{\rho(v_0; \beta)}{\rho(v_0)} \sum_{k=1}^{\infty} \int_{W^k} \prod_{l=1}^k (\rho(v_{l-1}) [g(v_{l-1}, v_l) - 1]) \rho'(v_k; \beta) dv_1 \cdots dv_k \end{aligned}$$

Topic of current research: truncate infinite sum or approximate integral ?