



# Space-time geostatistical modelling of malaria risk



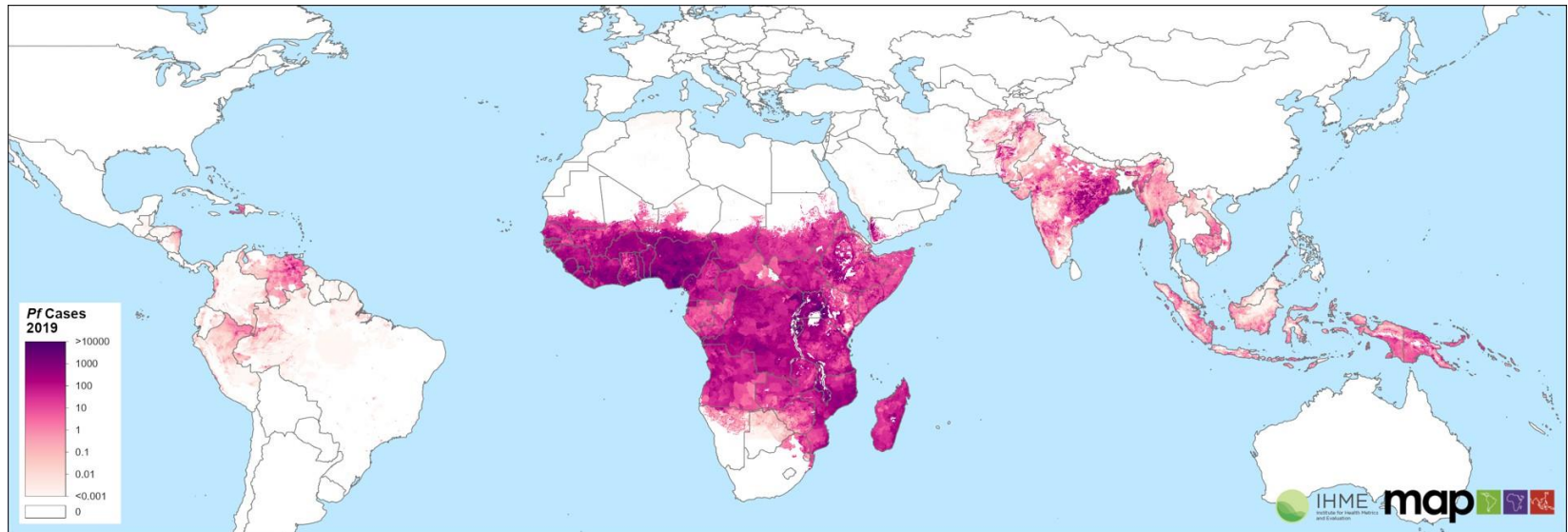
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# SPACE-TIME GEOSTATISTICAL MODELLING AND MAPPING OF OF MALARIA RISK

## Global malaria picture

Malaria for instance is a global challenge which continuous to affect deprived communities especially in sub-Saharan Africa

### PLASMODIUM FALCIPARUM CLINICAL CASES IN ALL AGE GROUPS IN 2019



Source: <https://malariaatlas.org/trends/region/MAP/GLOBAL>



# MALARIA AS A GLOBAL CHALLENGE

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- Maps have often been developed based on point reference data from demographic and health survey data
  - (Fobil et al, 2012)
  - (Yankson et al, 2019)
  - (Gemperli et al, 2006)
- Such data are however not routine, hence updating such maps is not feasible within shorter time periods
- They fail to examine the temporal trends



# DHIMS DATA FOR SURVEILLANCE

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- Routine surveillance managed through health information management systems can rather be helpful
- Despite such data are routinely collected, they have poor spatial and temporal resolution.
- For instance the
  - **DHIMS** is currently deployed by many developing countries to store and manage aggregated morbidity data
  - Accessible spatial resolution is at the district levels
  - Accessible temporal resolution is monthly

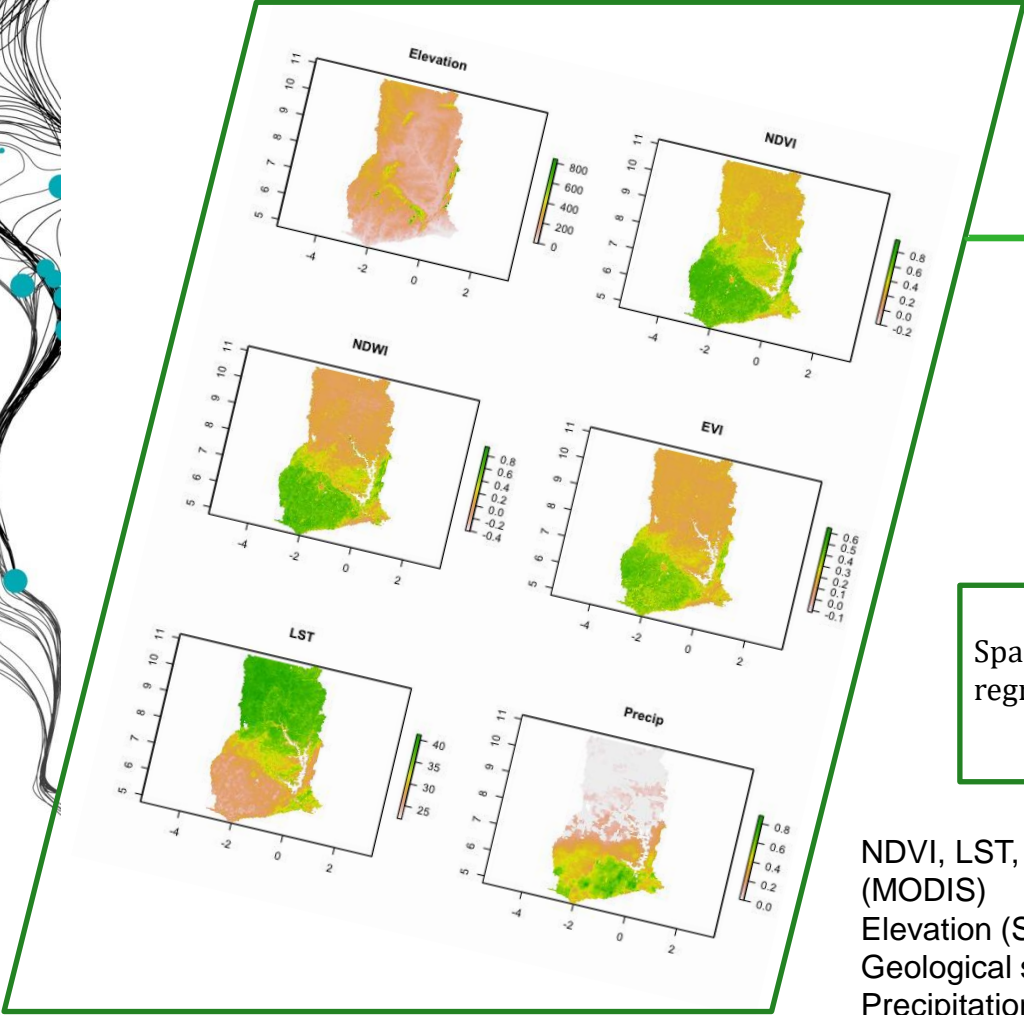


# LOCAL ACTION GAPS

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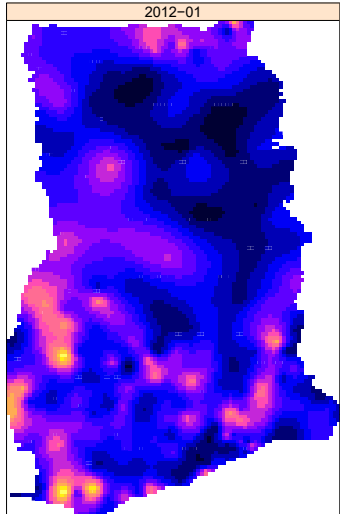
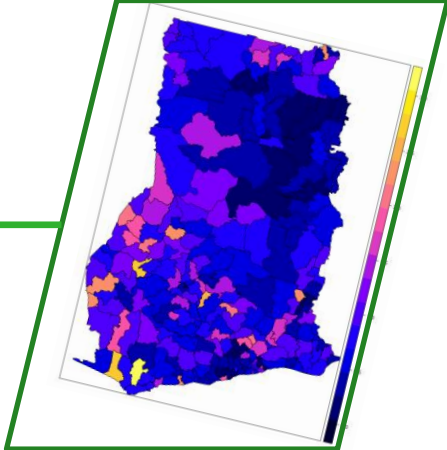
- Malaria
  - They remain a major public health burden in developing countries, especially in sub-Saharan Africa
  - Most studies
    - either focus on the spatial patterns at a particular point in time
    - or the temporal patterns for an entire geographic area
  - Why?
    - data challenges and/or
    - unavailable easy to implement statistical methods

# STATIC AND DYNAMIC REMOTELY SENSED COVARIATES (1 KM X 1 KM)



Space-time Poisson regression kriging

NDVI, LST, NDWI, EVI (MODIS)  
Elevation (SRTM, US Geological survey)  
Precipitation (Climate hazard group infrared precipitation)







# THE POISSON DATA GENERATING PROCESS

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- We consider malaria counts and population data
  - $\{y_{it}, n_{it}\}$
- As spatio-temporal outcomes of malaria and population data disaggregated by
  - districts  $i = 1, \dots, m = 216$
  - months  $t = 1, \dots, T = 55$  (from Jan 2012 to July 2016)
- Such sampling models are typically realizations from the Poisson process
  - $y_{it} | r_{it} \sim Poi(\lambda_{it})$
  - $\log \lambda_{it} = \log n_{it} + \log r_{it}$
  - $\log r_{it} = \beta_0 + \sum_{k=1}^K \beta_{kt} x_{ikt} + \sum_{p=1}^P \omega_k z_{ik}$
  - $x_{ikt}$  represent the dynamic covariates/predictors
  - $z_{ik}$  represent the static covariates/predictors



# STANDARD SEMIVARIANCE ESTIMATION

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- Here, the maximum likelihood estimate of the residual risk is

- $\tau_{it} = \frac{(y_{it} - \lambda_{it})}{n_{it}}$

- Assuming the risk was not a ratio variable, then
- The spatio-temporal variogram represents the semi-variance between any pair of risks estimates which are separated by spatial lag  $h$  and/or temporal lag  $u$ :

- $\gamma(h, u) = \frac{1}{2} \mathbb{E}(\tau_{it} - \tau_{i+h, t+u})^2$

- $\gamma^*(h, u) = \frac{1}{2} \sum_{N(h, u)} [\tau_{it} - \tau_{i+h, t+u}]^2$





# MODIFIED SEMI-VARIANCE ESTIMATION

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- The standard nonparametric geostatistical methods for semivariogram estimation used for continuous data are not directly applicable for the current model due to the
  - Heteroscedasticity (varying variance inversely proportional to the population)
  - non-stationarity of the process of counts

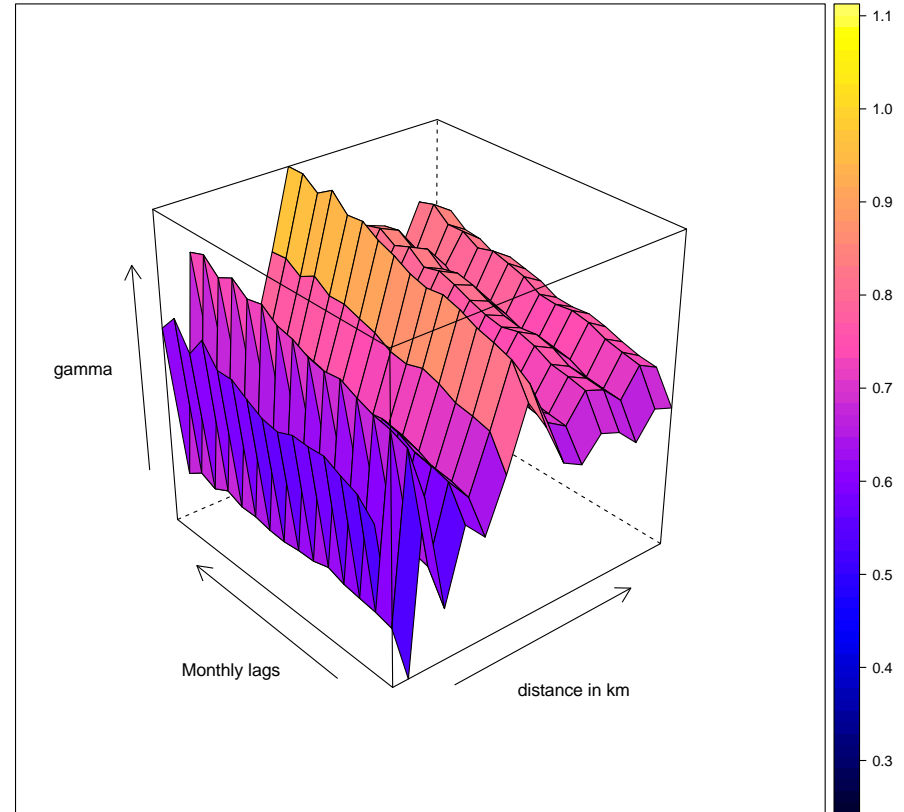
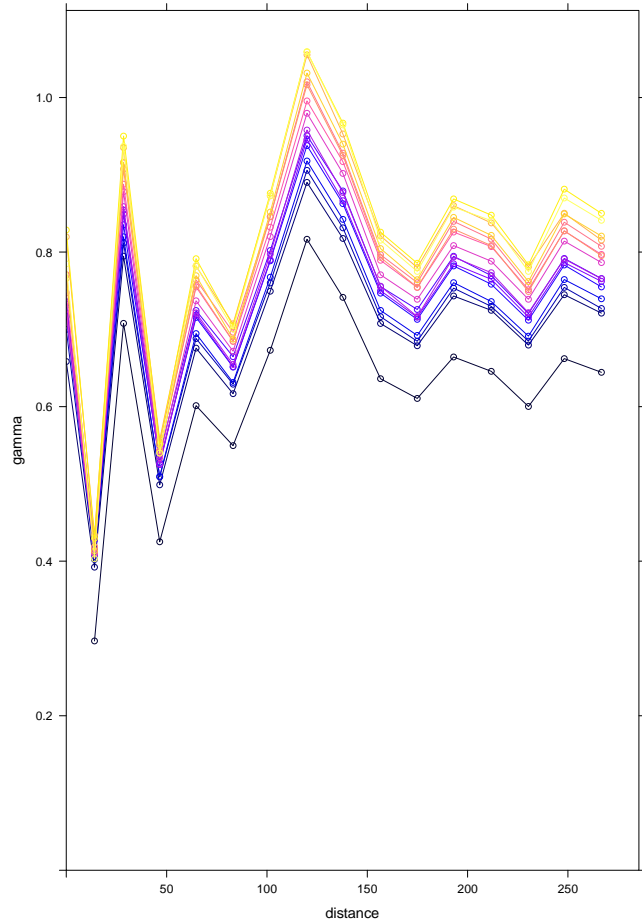
- $$\gamma^*(h, u) = \frac{\sum_{N(h,u)} \left\{ \frac{n_{it} \cdot n_{i+h,t+u}}{n_{it} + n_{i+h,t+u}} [\tau_{it} - \tau_{i+h,t+u}]^2 - \bar{\tau} \right\}}{2 \sum_{N(h,u)} \frac{n_{it} \cdot n_{i+h,t+u}}{n_{it} + n_{i+h,t+u}}}$$

- Where the population weighted average risk is

- $$\bar{\tau} = \frac{\sum n_{it} \cdot \tau_{it}}{\sum n_{it}}$$

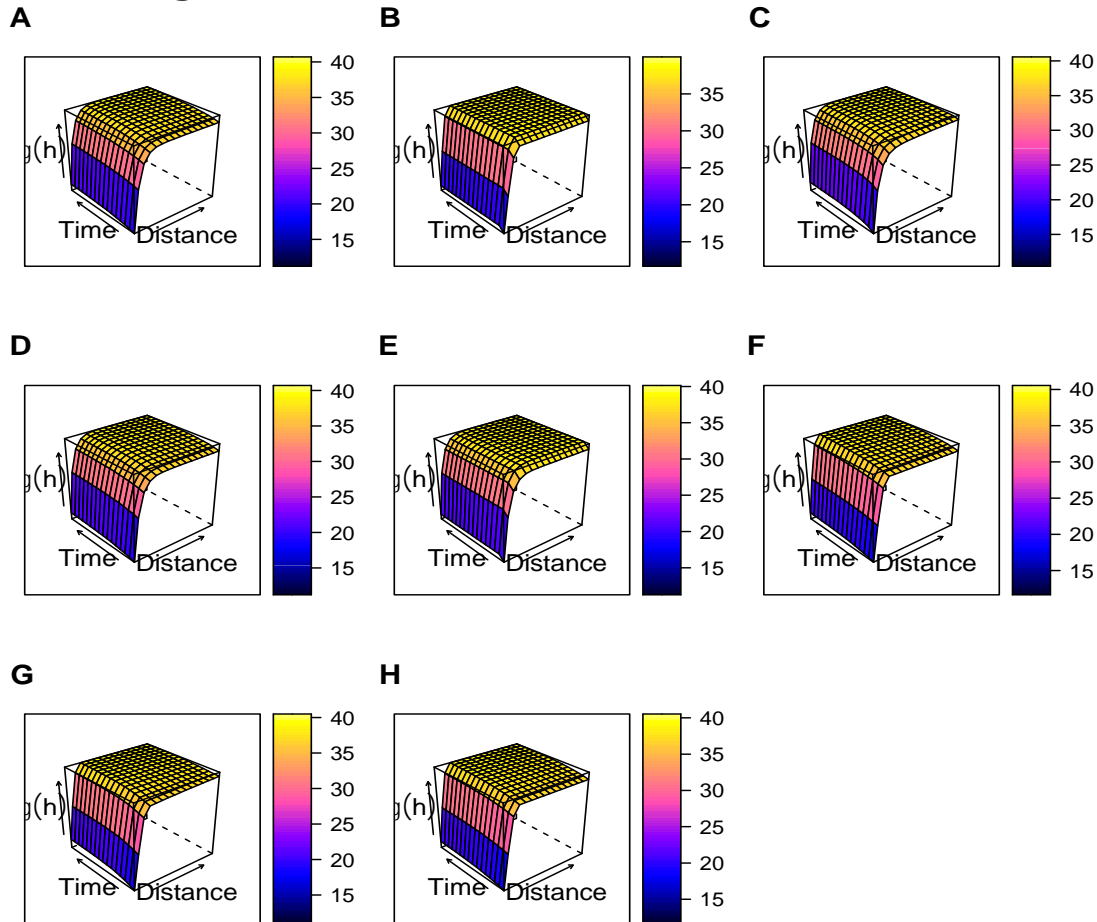
# EMPIRICAL SPACE-TIME SEMI-VARIOGRAM

The space-time variogram clearly shows indication of space-time correlation



# SPACE-TIME SEMI-VARIOGRAM MODELS

The combination of exponential, spherical and Gaussian space-time variogram models



# SPACE-TIME SEMI-VARIOGRAM MODEL PARAMETERS AND ASSOCIATED SSErr

	Estimates							
Parameters	EEE	SSS	ESE	EES	ESS	SES	SSE	SEE
Parameters	A	B	C	D	E	F	G	H
$\tau_s^2$	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
$\tau_t^2$	0.08	1.48	0.51	0.08	0.53	0.06	0.02	0.00
$\tau_{st}^2$	13.05	13.44	12.28	13.05	13.05	13.44	12.83	13.46
$\sigma_s^2$	23.75	23.27	23.55	23.75	23.75	23.27	23.06	23.26
$\sigma_t^2$	2.13	0.06	0.01	2.13	1.03	2.13	0.02	2.14
$\sigma_{st}^2$	0.00	0.00	2.37	0.00	0.00	0.00	2.80	0.00
$\phi_s$	8.86	22.33	8.87	8.86	8.86	22.33	22.25	22.38
$\phi_t$	5.44	0.4	0.4	5.45	0.4	5.45	0.4	5.13
$\phi_{st}$	1.69	1.56	4.49	1.48	1.54	1.58	5.54	6.92
<b>SSErr</b>	8.18	7.08	8.23	8.18	8.41	<b>6.85</b>	<b>6.87</b>	<b>6.85</b>

A (spatial: Exponential, temporal: Exponential, joint: Exponential); B (spatial: spherical, temporal: spherical, joint: spherical); C (spatial: exponential, temporal: spherical, joint: exponential); D (spatial: exponential, temporal: exponential, joint: spherical); E (spatial: exponential, temporal: spherical, joint: spherical); **F (spatial: spherical, temporal: exponential, joint: spherical); G (spatial: spherical, temporal: spherical, joint: exponential); H (spatial: spherical, temporal: exponential, joint: exponential)**



# SPACE-TIME PREDICTION

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- The prediction equation is a weighted average of neighboring risks
  - $\hat{\tau}_{i_0,t_0} = \sum_{it}^{m^T} \lambda_{it} \tau_{it}$
- Therefore
  - $\hat{\tau}_{i_0,t_0} = \exp(\beta_0 + \sum_{k=1}^K \hat{\beta}_{kt} x_{i_0,kt} + \sum_{p=1}^P \hat{\omega}_k z_{i_0,k}) + \hat{\tau}_{i_0,t_0}$
- The kriging weights express the strength of the association between observation locations and the prediction location, estimated as
  - $\lambda_{it} = \gamma_0^T \gamma^{-1}$
- The kriging weights takes into consideration the the spatial covariances between all data points as well as the covariances between the prediction locations and observation location



# UNCERTAINTIES AND CROSS VALIDATION

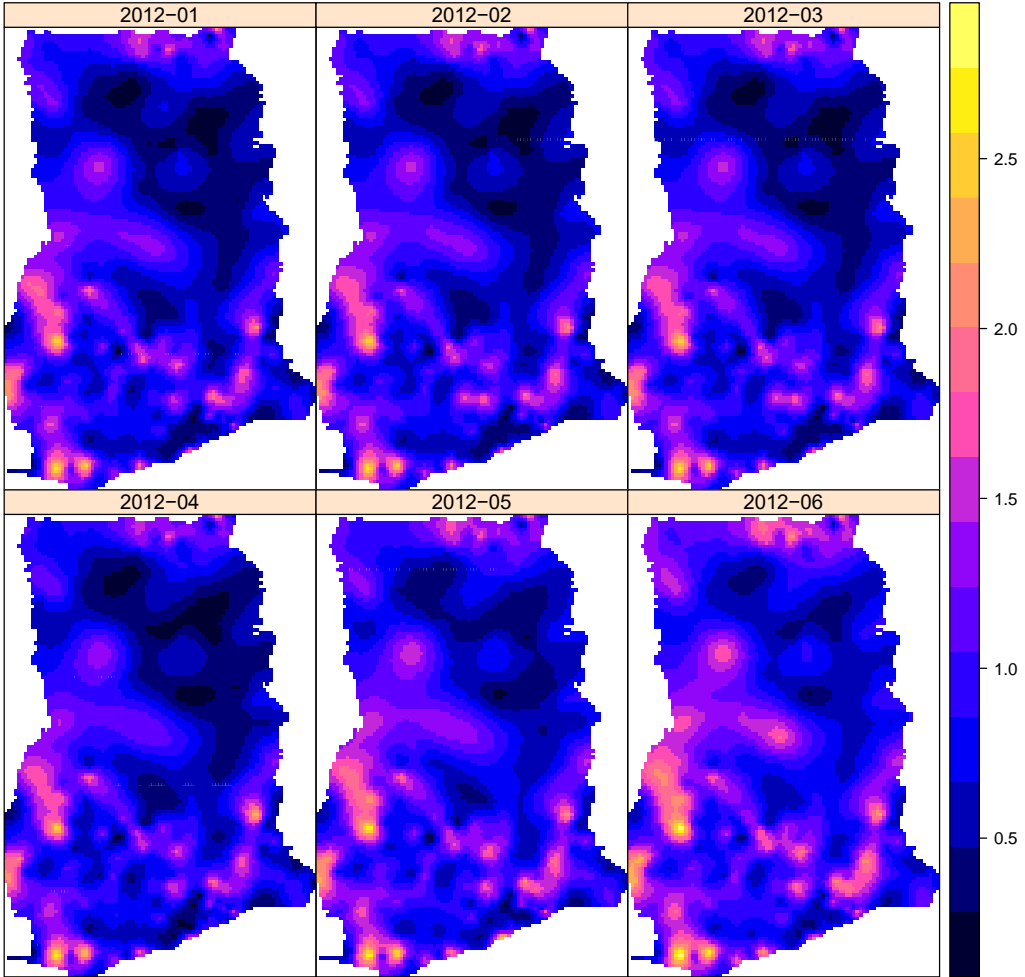
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- Prediction uncertainties were assessed through the kriging variance
  - $\sigma^2 (s_0, t_0) = \gamma_0^T \gamma^{-1} \gamma_0$
- We further used the leave-one-out cross validation (LOOCV) to assess the prediction accuracy.
- The Root Mean Square Error (RMSE) is then used to assess the accuracy of the kriging predictions.

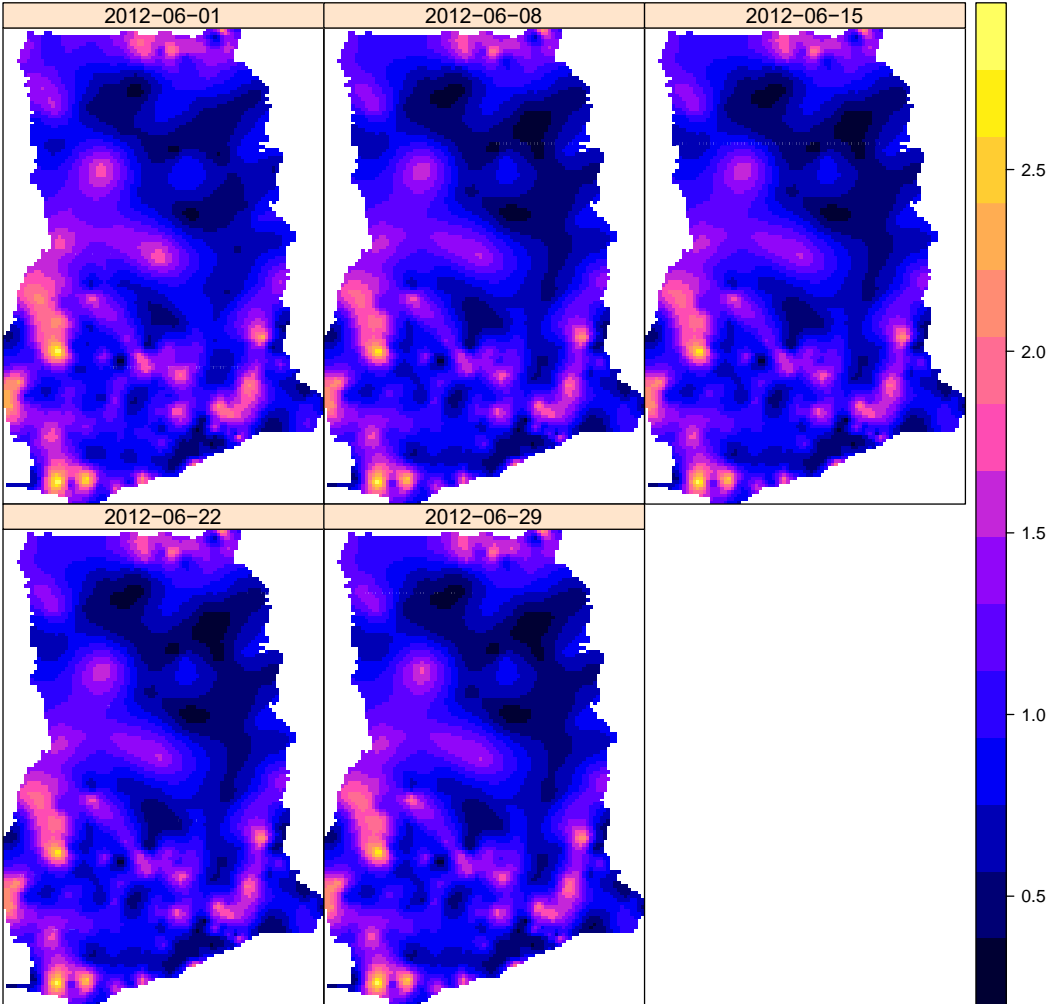
- $$RMSE = \sqrt{\frac{1}{m \times T} \sum_{s=1}^m \sum_{t=1}^T (\tau_{it} - \hat{\tau}_{it})^2}$$



# RESULTS: JAN-JUNE-2012



# RESULTS: WEEKS IN JUNE 2012





# CONTRIBUTIONS/CONCLUSIONS

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- **Methodology**

- Developed a spatial statistical framework for space-time prediction of malaria
- The method is transferable to other diseases with similar data structure

- **Epidemiologically**

- Observed spatial correlation of malaria risk, reflecting spatial clustering
- The spatial clustering is heterogenous across times/different months
- The spatial patterns and clusters are somehow spatially persistent

- **Further/Future work**

- Account for over-dispersion since the Poisson assumption has no variance parameter
- Improve the prediction by incorporating high temporal resolution external information for the covariates



# CONCLUSIONS

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- We have shown here that spatial statistics can be employed for analyzing and modelling geographically referenced health data
  - Either point pattern, lattice, or geostatistical data in terms of the underlying data generating process
  - Points and area data in terms of the feature representation
- The methods demonstrated in the sample studies are only few. With more challenging data structure, different relevant methods can be adapted and/or emerge
- This affirms that the beautiful marriage between spatial statistics and public health conceives GeoHealth

# THANK YOU

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- Q \$ A