## SpGPCW: Spatially Varying Gaussian Process Modeling for Critical Window Estimation

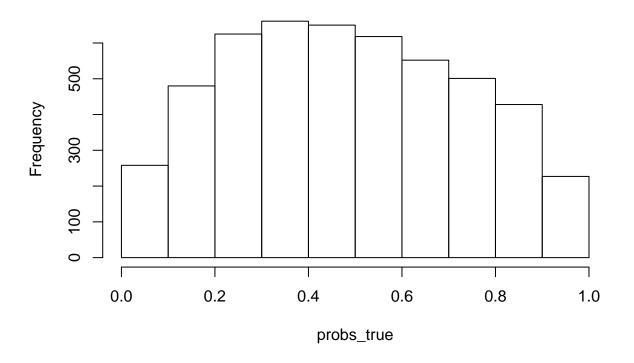
## SpGPCW\_Example

- [1] Simulate data from the proposed model:
  - Setting the reproducibility seed and initializing packages for data simulation:

```
set.seed(2365)
library(SpGPCW)
library(mnormt) #Multivariate normal distribution
library(boot) #Inverse logit transformation
library(spdep) #Creating a grid
## Loading required package: sp
## Loading required package: Matrix
## Loading required package: spData
## To access larger datasets in this package, install the spDataLarge
## package with: `install.packages('spDataLarge',
## repos='https://nowosad.github.io/drat/', type='source'))`
  • Setting the global data values:
n<-5000 #Sample size
m<-25 #Number of exposure time periods
g<-4 #Size of square spatial grid
s<-g^2 #Number of spatial locations
grid<-cell2nb(nrow=g,
              ncol=g,
              type="rook",
              torus=FALSE) #Evenly spaced grid
neighbors<-nb2mat(grid,
                  zero.policy=TRUE,
                  style="B") #Adjacency matrix
MCAR<-diag(rowSums(neighbors)) -</pre>
      neighbors
site_id<-rep(s, times=n)</pre>
for(j in 1:s){
   site_id[(1 + floor(n/s)*(j-1)):(floor(n/s)*j)] < -j
z<-matrix(0, nrow=n, ncol=m)</pre>
for(j in 1:s){
   z[(site_id == j),]<-matrix(rnorm(n=sum(site_id == j)),</pre>
                              nrow=sum(site_id == j),
                               ncol=m,
                               byrow=TRUE) #Exposure design matrices
for(j in 1:m){
  z[,j] < -(z[,j] - median(z[,j]))/IQR(z[,j]) #Data standardization (interquartile range)
  }
```

```
x<-matrix(1,</pre>
          nrow=n,
          ncol=2)
                  #Covariate design matrix
x[,2] < -rnorm(n)
beta_true<- c(-0.10, 0.20)
sigma2_theta_true<-0.35
phi theta true<-0.01
sigma2_eta_true<-0.10
phi_eta_true<-0.10
Sigma_theta_true<-sigma2_theta_true*chol2inv(chol(temporal_corr_fun(m, phi_theta_true)[[1]]))
Sigma_eta_true<-sigma2_eta_true*chol2inv(chol(temporal_corr_fun(m, phi_eta_true)[[1]]))
theta_true<-rmnorm(n=1,
                   mean=rep(0, times=m),
                   varcov=Sigma_theta_true)
theta_true<-theta_true -
            mean(theta_true)
rho_true<-0.45
eta_true<-rmnorm(n=1,
                 mean=rep(0, times=(m*s)),
                 varcov=chol2inv(chol(kronecker((rho_true*MCAR + (1 - rho_true)*diag(s)),
                                                  chol2inv(chol(Sigma_eta_true))))))
eta_true<-eta_true -
          mean(eta_true)
logit_p_true<-rep(0, times=n)</pre>
for(j in 1:s){
   logit_p_true[site_id == j]<-x[(site_id == j),]%*%beta_true +</pre>
                                z[(site_id == j),]%*%theta_true +
                                z[(site_id == j),]%*%eta_true[(1 + (j-1)*m):(j*m)]
probs_true<-inv.logit(logit_p_true)</pre>
hist(probs_true)
```

## Histogram of probs\_true



• Simulating the analysis dataset:

[2] Fit SpGPCW to estimate spatially varying critical windows of susceptibility:

```
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## **********************
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## phi_theta Acceptance: 32%
```

```
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## Progress: 70%
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```

## rho Acceptance: 38%

```
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## Progress: 80%
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## ************
## Progress: 100%
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## rho Acceptance: 39%
## phi_eta Acceptance: 23%
## *********
[3] Analyzing Output:
par(mfrow=c(2,2))
plot(results$beta[1, 1001:10000],
    type="1",
    ylab="beta0",
    xlab="Sample")
abline(h=beta_true[1],
      col="red",
      lwd=2) #True value
plot(results$beta[2, 1001:10000],
    type="1",
    ylab="beta1",
    xlab="Sample")
abline(h=beta_true[2],
      col="red",
      lwd=2) #True value
plot(rowMeans(results$theta[,1001:10000]),
    theta_true)
abline(0, 1)
```

```
eta<-simplify2array(results$eta)
eta_post_means<-rep(0, times=(s*m))
counter<-0
for(j in 1:s){
    for(k in 1:m){
        counter<-counter + 1
        eta_post_means[counter]<-mean(eta[j,k,1001:10000])
        }
    }
plot(eta_post_means,
    eta_true)
abline(0, 1)</pre>
```

