# Drug Abuse by State

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#### Reading in and Handling Shapefiles

1. Read in the drug abuse shapefile.

```
#load dataset
#myShp <- readOGR(dsn="~/R programming/STAT_469/Unit4",layer='DrugAbuse')</pre>
#library(broom) #contains tidy() function which converts polygons to data.frame
#myShp@data$id <- rownames(myShp@data) #Assign ID to each polygon</pre>
#myShp.df <- tidy(myShp, region = "id") #Convert polygon info to data.frame()</pre>
\#myShp.df \leftarrow merge(myShp.df, myShp@data, by = "id") \#Merge data w/polygon data.frame
myShp <- st_read("~/R programming/STAT_469/Unit4/DrugAbuse.shp")</pre>
## Reading layer 'DrugAbuse' from data source
     'C:\Users\jilli\OneDrive\Documents\R programming\STAT_469\Unit4\DrugAbuse.shp'
     using driver 'ESRI Shapefile'
## Simple feature collection with 49 features and 6 fields
## Geometry type: MULTIPOLYGON
## Dimension:
                  XY
## Bounding box:
                  xmin: -124.7328 ymin: 24.95638 xmax: -66.96927 ymax: 49.37173
## Geodetic CRS:
                  NAD83
```

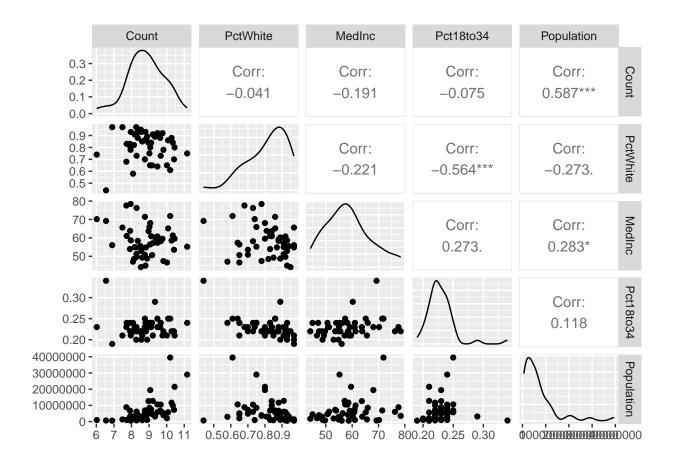
2. Reformat the shapefile data into a dataframe suitable for use with ggplot().

```
data <- data.frame(myShp) %>% select(-geometry)
#data <- data.frame(myShp.df)
#data <- data %>% dplyr::select(-c(id, long, lat, order, hole, piece, group))
data_red <- data %>% dplyr::select(-c(State))
data_red <- data_red %>% mutate(Count = log(Count))
```

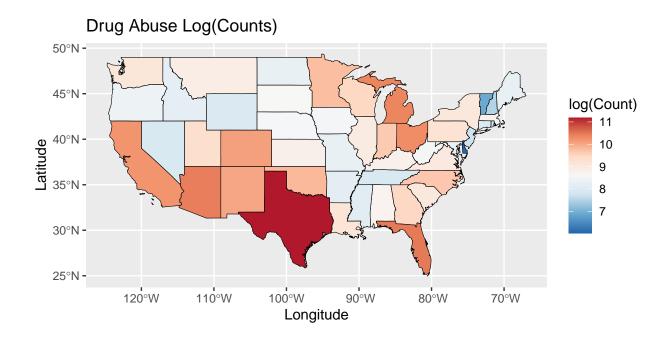
### **Exploratory Data Analysis**

1. Create a pairs plot to assess the relationship between log(Count) and the explanatory variables (note we are using log(Count) here as the response because Poisson regression is log-linear).

```
ggpairs(data_red)
```



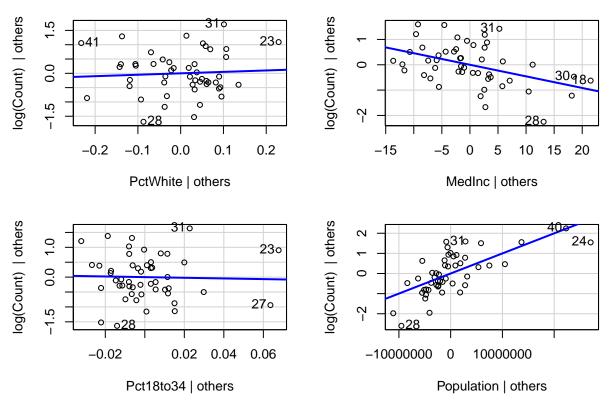
2. Create a choropleth map of log(Count).



3. Fit a lm() of log(Count) using Population, PctWhite, MedInc and Pct18to34 as explanatory variables. Perform a Moran's I test on the residuals to see if there is spatial correlation in the residuals.

data\_lm <- lm(log(Count)~ PctWhite + MedInc + Pct18to34 + Population, data)
avPlots(data\_lm, ask=FALSE)</pre>

### Added-Variable Plots



```
moran.test(x= resid(data_lm), listw=nb2listw(poly2nb(myShp)))
```

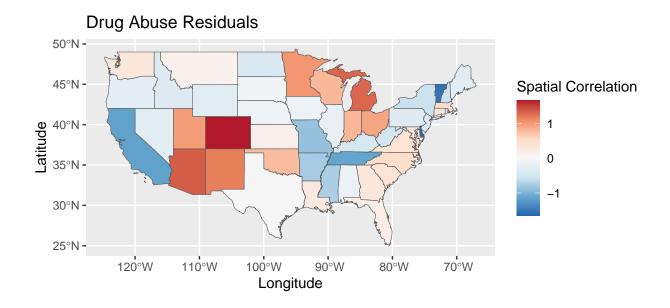
```
##
##
   Moran I test under randomisation
##
## data: resid(data_lm)
##
  weights: nb2listw(poly2nb(myShp))
##
## Moran I statistic standard deviate = 2.5921, p-value = 0.004769
  alternative hypothesis: greater
  sample estimates:
##
  Moran I statistic
                           Expectation
                                                 Variance
##
         0.230238791
                          -0.020833333
                                              0.009381812
```

4. Perform a Geary's C test on your residuals from #3 above to double check if there is spatial correlation in the residuals.

```
geary.test(x= resid(data_lm), listw=nb2listw(poly2nb(myShp)))
```

```
##
## Geary C test under randomisation
##
## data: resid(data_lm)
## weights: nb2listw(poly2nb(myShp))
```

5. Map the residuals from the lm() fit to see if there is spatial correlation.



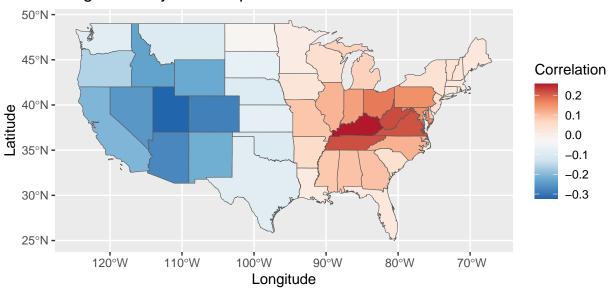
### **Defining Spatial Basis Functions**

1. Create the adjacency matrix.

```
A <- nb2mat(poly2nb(myShp), style="B")
```

2. Create the Moran spatial basis and plot the first basis in a chloropleth map.

### Drug Abuse by Moran Spatial Basis



3. Merge the Moran spatial bases into your myShpDF data frame for use in fitting models later.

```
data <- bind_cols(data, M) #cbind() but efficient</pre>
```

### Spatial GLM Model Fitting

1. Fit a spatial GLM model with Count as the response and using PctWhite, MedInc, Population, Pct18to34 AND your spatial bases as explanatory variables. Print a summary() of the model to see your coefficient table.

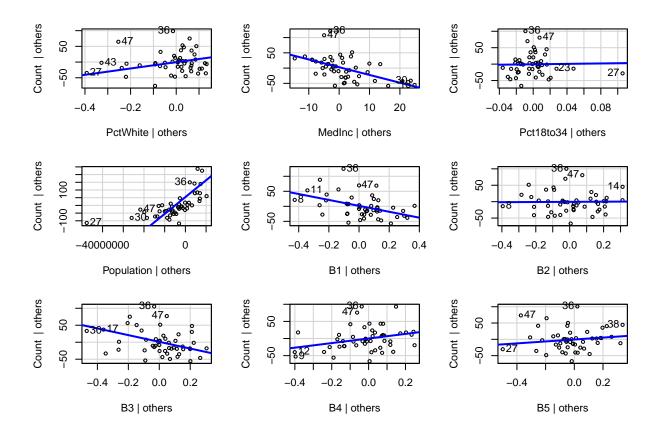
```
data_glm <- glm(formula=Count~.-State-Count, data=data, family=poisson)
summary(data_glm)</pre>
```

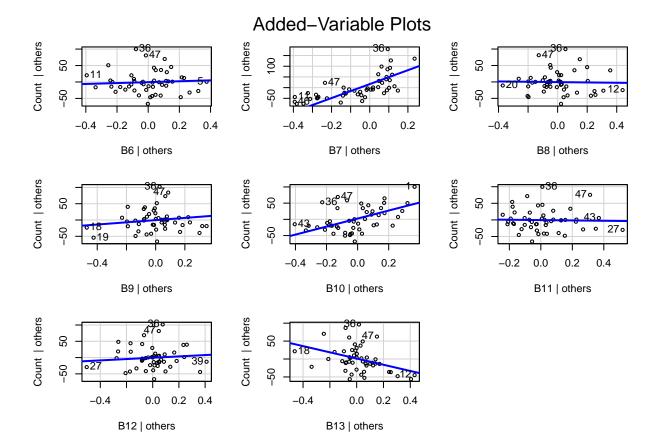
```
##
## Call:
## glm(formula = Count ~ . - State - Count, family = poisson, data = data)
##
## Deviance Residuals:
                     Median
##
      Min
                 1Q
                                   3Q
                                           Max
                      -1.849
## -71.238 -23.869
                               11.556
                                        90.700
##
## Coefficients:
##
                 Estimate Std. Error z value Pr(>|z|)
## (Intercept) 9.856e+00
                          2.699e-02
                                     365.161
                                              < 2e-16 ***
## PctWhite
                          1.798e-02
                                      35.327
                                              < 2e-16 ***
               6.351e-01
## MedInc
               -3.299e-02
                          2.491e-04 -132.425
                                              < 2e-16 ***
## Pct18to34
              -1.863e-01
                          9.051e-02
                                      -2.059
                                              0.03954 *
                          1.877e-10 427.517
                                              < 2e-16 ***
## Population
              8.025e-08
## B1
               -1.202e+00
                          1.098e-02 -109.469
                                              < 2e-16 ***
## B2
              -1.085e-02
                          1.188e-02
                                      -0.914 0.36089
## B3
              -9.081e-01
                          1.017e-02
                                     -89.267
                                              < 2e-16 ***
                          1.374e-02
                                      71.047
## B4
               9.765e-01
                                              < 2e-16 ***
## B5
               1.921e-01
                          1.062e-02
                                      18.084
                                              < 2e-16 ***
## B6
               1.199e-02 1.143e-02
                                       1.049 0.29437
## B7
               3.149e+00 1.123e-02 280.304
                                              < 2e-16 ***
              -3.388e-02 1.232e-02
                                      -2.750 0.00595 **
## B8
## B9
               3.500e-01
                          1.195e-02
                                      29.282
                                              < 2e-16 ***
## B10
               1.164e+00
                          1.010e-02 115.231
                                              < 2e-16 ***
## B11
              -1.620e-01
                          1.207e-02
                                     -13.423
                                              < 2e-16 ***
## B12
               1.128e-01
                          1.173e-02
                                       9.619
                                              < 2e-16 ***
              -1.264e+00 1.258e-02 -100.454 < 2e-16 ***
## B13
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
## (Dispersion parameter for poisson family taken to be 1)
##
##
       Null deviance: 496777
                             on 48 degrees of freedom
## Residual deviance: 53329 on 31 degrees of freedom
## AIC: 53887
##
## Number of Fisher Scoring iterations: 5
```

#### Validating Spatial MLR Model Assumptions and Predictions

1. Check the assumption of linearity using added-variable plots.

```
avPlots(data_glm, ask=FALSE)
```





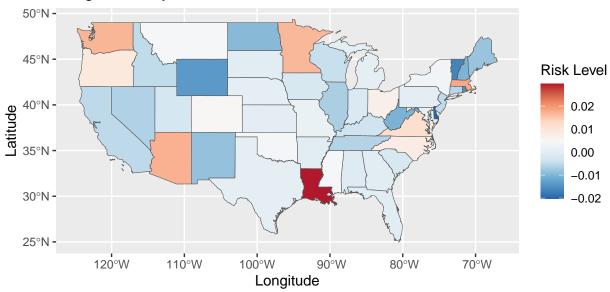
2. Check the assumption of independence by decorrelating residuals and performing Moran's I or Geary's C tests to make sure there is no more spatial correlation.

```
sres <- stdres.gls(data_glm)</pre>
moran.test(x= sres, listw=nb2listw(poly2nb(myShp)))
##
##
    Moran I test under randomisation
##
  data: sres
##
##
   weights: nb2listw(poly2nb(myShp))
##
## Moran I statistic standard deviate = -1.2132, p-value = 0.8875
   alternative hypothesis: greater
   sample estimates:
## Moran I statistic
                            Expectation
                                                  Variance
##
        -0.136699956
                           -0.020833333
                                              0.009120747
geary.test(x= sres, listw=nb2listw(poly2nb(myShp)))
##
    Geary C test under randomisation
##
##
```

## data: sres

3. Draw a choropleth map of the standardized and decorrelated residuals to visually verify that the residuals are no longer spatially correlated.

## Drug Abuse by decorrelated residuals

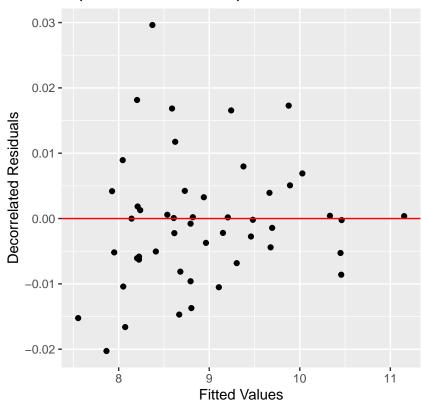


4. Check the assumption of equal variance by plotting the standardized and decorrelated residuals vs. the log(fitted values).

```
ggplot(mapping = aes(x=log(fitted(data_glm)), y=sres)) +
geom_point() +
xlab('Fitted Values') +
```

```
ylab('Decorrelated Residuals') +
ggtitle('Equal Variance Assumption Check:') +
geom_hline(yintercept = 0, col = "red") +
theme(aspect.ratio = 1)
```

### **Equal Variance Assumption Check:**



### Statistical Inference

1. Print out the summary of the GLM model fit and identify the estimates and 95% confidence intervals of your explanatory variables.

### summary(data\_glm)

```
##
## Call:
## glm(formula = Count ~ . - State - Count, family = poisson, data = data)
##
## Deviance Residuals:
##
       Min
                 1Q
                      Median
                                   3Q
                                           Max
##
  -71.238
           -23.869
                      -1.849
                               11.556
                                        90.700
##
## Coefficients:
                 Estimate Std. Error z value Pr(>|z|)
##
## (Intercept) 9.856e+00 2.699e-02 365.161 < 2e-16 ***
```

```
## PctWhite
               6.351e-01 1.798e-02
                                      35.327 < 2e-16 ***
## MedInc
              -3.299e-02 2.491e-04 -132.425
                                             < 2e-16 ***
## Pct18to34
                                             0.03954 *
              -1.863e-01
                          9.051e-02
                                      -2.059
## Population
              8.025e-08
                          1.877e-10
                                    427.517
                                             < 2e-16 ***
## B1
              -1.202e+00
                          1.098e-02 -109.469
                                             < 2e-16 ***
## B2
              -1.085e-02 1.188e-02
                                      -0.914 0.36089
## B3
              -9.081e-01 1.017e-02
                                     -89.267 < 2e-16 ***
                                      71.047 < 2e-16 ***
## B4
               9.765e-01
                         1.374e-02
## B5
               1.921e-01
                          1.062e-02
                                      18.084
                                             < 2e-16 ***
## B6
               1.199e-02
                          1.143e-02
                                       1.049 0.29437
## B7
               3.149e+00
                          1.123e-02
                                     280.304 < 2e-16 ***
                          1.232e-02
                                      -2.750 0.00595 **
## B8
              -3.388e-02
## B9
               3.500e-01
                          1.195e-02
                                      29.282 < 2e-16 ***
## B10
               1.164e+00 1.010e-02 115.231 < 2e-16 ***
## B11
              -1.620e-01 1.207e-02
                                     -13.423 < 2e-16 ***
## B12
               1.128e-01
                          1.173e-02
                                       9.619 < 2e-16 ***
              -1.264e+00 1.258e-02 -100.454 < 2e-16 ***
## B13
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
## (Dispersion parameter for poisson family taken to be 1)
##
##
      Null deviance: 496777 on 48 degrees of freedom
## Residual deviance: 53329 on 31 degrees of freedom
## AIC: 53887
## Number of Fisher Scoring iterations: 5
```

2. Create a chloropleth map of the spatially correlated residuals (just the  $\mathbf{b}'_i\hat{\theta}$  part) to identify states that, after accounting for the explanatory variables, have an elevated level of risk.

