Bio284 Computer Lab 6 Spatial Cluster Detection for Binary Data

1 Goals for this lab:

- 1. Independent Binary Data
 - (a) Spatial Scan Statistic
 - (b) Cumulative Residual test
 - (c) Apply to North Humberside Dataset
- 2. Matched Binary Data
 - (a) Conditional Cumulative Residual test
 - (b) Apply to Taiwan Petrochemical Study
- 3. ArcGIS to plot results



North Humberside

2 Independent Binary Data

1. Dataset

(Included within the SatScan Package)

Childhood Leukemia and Lymphoma Incidence in North Humberside

This data set describes the spatial location of 62 cases of childhood leukemia and lymphoma in North Humberside between 1974 and 1986, as well as 141 controls.

-Copy all NHumberside data on G:/shared into a directory

2. Spatial Scan Statistic

We will run a cluster detection analysis on the North Humberside dataset using the Spatial Scan Statistic and the SatScan software.

- (a) Download the SatScan Software from www.satscan.us
 (Note: Documentation and downloads for SatScan at www.satscan.us)
 -Open SatScan from Start→ All Programs→ SatScan
 -Click on create a new session.
- (b) Conduct Analysis
 - Input Tab

Click \cdots by Case File space and go to where you saved the NHumberside files. You will automatically be looking for files of type ".cas". What this file type means is a text file with a column with ID of subject and a column with binary indicator of case (1) or control (0) and no headers.

Click \cdots by Control File space. You will automatically be looking for files of type ".ctl". What this file type means is a text file with a column with ID of subject and a column with binary indicator of case (0) or control (1) and no headers.

Under time precision select, none, since we have no time component in our analysis.

Click \cdots by Coordinates File space. You will automatically be looking for files of type ".geo". What this file type means is a text file with a column with ID of subject, a column with x coordinates and a column with y coordinates, no header.

Under coordinates select Cartesian.

- Analysis Tab Under Probability model select Bernoulli.
- Output Tab

Select all outputs of type dbf since ArcGIS handles this readily. Select a file name for the results

• Execute Analysis

Click the yellow lightning rod to execute the code and perform the analysis.

(c) **Results**

SaTScan v5.0

Program run on: Wed Dec 15 19:50:20 2004 Purely Spatial analysis scanning for clusters with high rates using the Bernoulli model. _____ SUMMARY OF DATA Study period.....: 2000/1/1 - 2000/12/31 Number of locations...: 191 Total population.....: 203 Total cases....: 62 _____ MOST LIKELY CLUSTER 1.Location IDs included.: 19, 18, 14, 26 Coordinates / radius..: (5026,4301) / 4.47 Population....: 4 Number of cases....: 4 Expected cases....: 1.22 Observed / expected...: 3.274 Log likelihood ratio..: 4.836516 Monte Carlo rank....: 674/1000

Is there any clustering?

Where is the highest cluster located?

P-value....: 0.674

3. Cumulative Geographic Residual

For the cumulative geographic residual test we will be using the R package to conduct the analysis.

We will be using the file NHnumberside.txt for the dataset. It has a header with the variables ID, Y (indicator of case(1) and control(0)), x, and y.

```
NH<-read.table("C:/NHumberside.txt",header=T)
```

```
# Binary Outcome Variable
Y<-NH$Y</pre>
```

```
# Matrix of locations (x,y)
loc<-NH[,3:4]</pre>
```

The program is in a txt file called cumres-BinR.txt. It contains the main function, CumRes.Bern, and two corresponding plot function, plot.CR and plot.CRs.

```
source("C:/cumres-BinR.txt")
```

Lets run the first analysis. There is one main thing to think about when running the cumulative residual test.

Window Size : (b_1, b_2)

We will use (50,50) for now, but when running this method one should choose several window sizes and conduct a sensitivity analysis.

```
# Range of Study area
xrange<-c(min(loc$x),max(loc$x))
yrange<-c(min(loc$y),4710)</pre>
```

```
CRtest1<-CumRes.Bern(Y,loc,X=NULL,xarea=xrange,yarea=yrange,
b=c(50,50),jumps=c(25,25),1000)
```

RESULTS:

CRtest1[11:16] \$Sloc [1] 0.2398025

\$pval Z D [1,] 0.609 0.639

\$maxloc					
x y [1,] 5127 4325					
\$crit					
		Z		D	
90%	0.41	28042	0.411	6202	
95%	0.46	80560	0.469	6235	
97.5%	0.51	.21604	0.532	5975	
\$meanSlochat					
		Z		D	
[1,] ().282	9770	0.2848	552	
\$quantSlochat					
		Z		D	
2.5%	0.13	848415	0.133	0336	
5%	0.15	34783	0.149	4689	
10%	0.17	16653	0.175	5866	
50%	0.26	54225	0.268	4124	
90%	0.41	28042	0.411	6202	
95%	0.46	80560	0.469	6235	
97.5%	0.51	21604	0.532	5975	

- (a) Is there significant clustering?
- (b) Where is the highest cluster?

Plot the Results

plot.CR(CRtest1)





For input into ArcGIS:

```
NHarc<-data.frame(NH)
names(NHarc)<-c("ID","Y","xcoord","ycoord")
write.table(NHarc,"C:/NHarc.txt",row.names=F,sep=",")</pre>
```

```
# create a cluster area dataset maxloc<-CRtest1$maxloc
b<-CRtest1$b</pre>
```

```
x<-c(maxloc[1]-b[1],maxloc[1]-b[1],maxloc[1],maxloc[1])
y<-c(maxloc[2],maxloc[2]-b[2],maxloc[2]-b[2],maxloc[2])</pre>
```

write.table(cbind(x,y),"C:/NHcr.txt",row.names=F,sep=",")

3 Matched Binary Data

1. Dataset

Childhood Leukemia and Petrochemical Exposure in Kaohsiung, Taiwan (P.I. David Christiani)

This data set describes the spatial location of 121 cases of childhood leukemia in Kaohsiung as well as 287 controls.

-Copy all petro data on G:/shared into a directory

2. Analysis

We will be using the file petro.txt for the dataset. It has a header with the variables stratum, CASE, X.coord, Y.coord, SMOKE, and SUBSMOKE.

petro<-read.table("C:/petro.txt",header=T)</pre>

```
Y<-petro$CASE
x<-petro$X.coord
y<-petro$Y.coord
loc<-cbind(x,y)
stratum<-petro$stratum
X<-cbind(petro$SMOKE,petro$SUBSMOKE)</pre>
```

The program is in a txt file called cumres-MBinR.txt. It contains the main function CumRes.MBern.

source("C:/cumres-MBinR.txt")

We will use window size of (4800,9400) for now, but when running this method one should choose several window sizes and conduct a sensitivity analysis.

(a) WITHOUT COVARIATES

b<-c(4800,9400) jumps<-c(1000,1000)

```
CRpetro1<-CumRes.MBern(Y=Y,loc=loc,X=NULL,b=b,stratum=stratum,jumps=jumps,
nsims=1000,type="mixed")
```

RESULTS:

CRpetro1[12:17] \$Sloc [1] 0.939616

\$pval Z D [1,] 0.028 0.045

\$maxloc

x y [1,] 179187 2511273

\$crit

Z D 90% 0.7967008 0.8179731 95% 0.8761764 0.9090721 97.5% 0.9703929 1.0148213

meanSlochat

Z D [1,] 0.526095 0.5274363 \$quantSlochat

	Z	D
2.5%	0.2136219	0.1985431
5%	0.2425349	0.2396608
10%	0.2843336	0.2856456
50%	0.4982981	0.4992677
90%	0.7967008	0.8179731
95%	0.8761764	0.9090721
97.5%	0.9703929	1.0148213

- i. Is there significant clustering?
- ii. Where is the highest cluster?

Plot the Results

plot.CR(CRpetro1)



plot.CRs(CRpetro1)
(b) WITH SMOKING COVARIATES

b<-c(4800,9400) jumps<-c(1000,1000)

CRpetro2<-CumRes.MBern(Y=Y,loc=loc,X=X,b=b,stratum=stratum,jumps=jumps, nsims=1000,type="mixed")

RESULTS:

CRpetro2[12:17] \$Sloc [1] 0.9390835

\$pval

Z D



[1,] 0.148 0.139 \$maxloc х у [1,] 179187 2511273 \$crit Ζ D 90% 1.017588 1.010507 95% 1.106199 1.138549 97.5% 1.213654 1.257334 \$meanSlochat D Ζ [1,] 0.695705 0.69668 \$quantSlochat Ζ D 0.3344124 0.3375272 2.5% 0.3746685 0.3760654 5% 10% 0.4202907 0.4424850 0.6664539 0.6676163 50% 90% 1.0175876 1.0105074 1.1061986 1.1385487 95% 97.5% 1.2136542 1.2573343

i. Is there significant clustering?

ii. Where is the highest cluster?

Plot the Results

plot.CR(CRpetro2)



plot.CRs(CRpetro2)



4 ArcGIS

We will plot the NHumberside data results for both the spatial scan statistic and the cumulative geographic residual test under the independence assumption.

1. Data Manipulation

To get the data from SatScan into ArcGIS you need to open the SSresults.col.dbf in excel. The x,y, and radius columns are in text form instead of number format. Reformat these columns into number format by just selected the columns without the header and clicking on the question mark box. Also remove any ", marks in the parts of the column names since ArcGIS cannot handle that type of naming. For our example delete all of the rows except for the first entry since we will only be looking at the first cluster.

2. Open ArcGIS and import data

Addxy data NHarc.txt, SSResults.col.dbf, and NHcr.txt onto your layout.

You will have a plot of all the cases and controls. You will also have a point for the center point of the circle for the Spatial Scan and will have four points depicting the highest square area from the cumulative residual results.

3. Spatial Scan buffer

You need to add the buffer wizard to draw the circle. Go to Tools, customize, and click on the Commands tab. Scroll to tools and you will see the Buffer Wizard in the commands window. Drag and click it to the main map ArcGIS toolbar. Close the customize window.

We need to put map units before using the buffer tool. Right click in map area and select Properties and click on the General tab. In the units section, click on kilometers and push okay.

Open the attributes table of SS results.col and highlight the first row. We need to select this row to create a buffer around this point.

Now double click on the Buffer Wizard in the toolbar. Scroll to SSresults.col and you should see 1 for both number of features and number of features selected. Click next.

At the specified distance click 4 kilometers since this is the radius of the file.

Click next and rename your new file for a buffer. Click finish and a new shape file with your buffer will appear on the display. It is very small since it only had included four subjects.

4. Cumulative Residual

Open attributes table of NHcr.txt. We will be using these points to draw a square. Click on the Editors tool and say start editing. (If the Editor tool is not displayed, right click in toolbar area, and select Editor).

Click on the pencil icon. Then right-click in the map area and select Absolute X,Y. Enter the coordinates in the first line of NHcr.txt and press return. Right-click again, select Absolute X,Y, and enter coordinates in second line and press return. Continue for all four lines. Afterward, Right-click again and now click on Finish Sketch.

Now you have both areas outlined for both methods. Click on Editor toolbar and say stop editing and save your results.

Here is one way to create maps of these results. You can use all of the tools from previous labs to make things look better then what you have now.