

# Further Changepoint Analysis

Rebecca Killick(r.killick@lancs.ac.uk) NHS Workshop 2021

## Workshop Plan



- Recap of changepoints
- Checking assumptions
- Autocorrelation
- Multivariate changepoints
- Influence

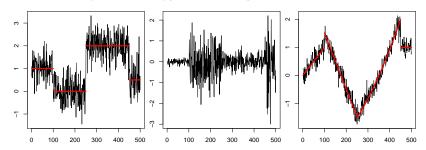
There will be tasks throughout the sections.

## Recall: changepoints



For data  $y_1, \ldots, y_n$ , if a changepoint exists at  $\tau$ , then  $y_1, \ldots, y_{\tau}$  differ from  $y_{\tau+1}, \ldots, y_n$  in some way.

There are many different types of change.



## **Packages**



Today we will use the following packages

library(changepoint)

library(EnvCpt)

library(changepoint.influence)

library(changepoint.geo)

Other notable R packages are available for changepoint analysis including

- ecp for univariate and multivariate energy test statistics
- InspectChangepoint for multivariate Inspect projection direction mean only change
- hdbinseg for multivariate double CUSUM test statistic
- BayesProject for multivariate changepoints

## **Checking Assumptions**



The main assumptions for a Normal likelihood ratio test for a change in mean are:

- Independent data points;
- Normal distributed points pre and post change;
- Constant variance across the data.

How can we check these?

### How to check



Check the residuals

```
set.seed(1)
m1=c(rnorm(100,0,1),rnorm(100,5,1))
m1.amoc=cpt.mean(m1)
means=param.est(m1.amoc)$mean
m1.resid=m1-rep(means, seg.len(m1.amoc))
shapiro.test(m1.resid)
##
##
    Shapiro-Wilk normality test
##
  data: m1.resid
## W = 0.99228, p-value = 0.3721
```

### **Residual Check**



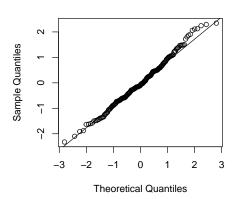
```
ks.test(m1.resid,pnorm,mean=mean(m1.resid),sd=sd(m1.resid))
##
## One-sample Kolmogorov-Smirnov test
##
## data: m1.resid
## D = 0.045812, p-value = 0.7953
## alternative hypothesis: two-sided
```

### **Residual Check**



qqnorm(m1.resid)
qqline(m1.resid)

#### Normal Q-Q Plot

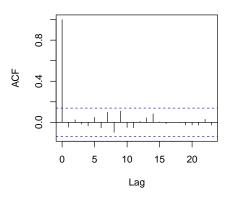


### **Residual Check**



acf(m1.resid)

Series m1.resid

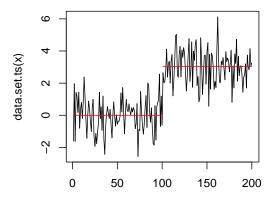


### **Autocorrelation**



What effect does autocorrelation have on our analysis?

```
set.seed(879123)
x=c(rnorm(100),rnorm(100,3))
plot(cpt.meanvar(x,method='PELT'))
```

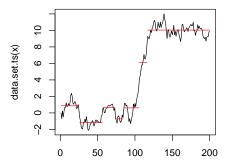


### Autocorrelation +ve



What effect does autocorrelation have on our analysis?

```
source('sim.cpt.AR1.R')
set.seed(879123)
x=sim.cpt.AR1(cpts=c(0,100,200),X=cbind(rep(1,200)),init=0,
    beta=rbind(c(0,0.9),c(1,0.9)),sig2=(1-0.9^2),nsim=1)
plot(cpt.meanvar(x,method='PELT'))
```

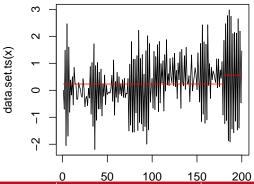


### Autocorrelation -ve



What effect does autocorrelation have on our analysis?

```
set.seed(879123)
x=sim.cpt.AR1(cpts=c(0,100,200),X=cbind(rep(1,200)),init=0,
    beta=rbind(c(0,-0.9),c(1,-0.9)),sig2=(1-0.9^2),nsim=1)
plot(cpt.meanvar(x,method='PELT'))
```



### **Exercise**



Take a look at the "Lai2005fig4" data in the changepoint package. Fit changes in mean as below, then check the residuals.

Are the assumptions of our model reasonable?

```
data("Lai2005fig4")
out=cpt.mean(Lai2005fig4$GBM29,method='PELT')
```

Don't forget to look at the data!

# **EnvCpt**



EnvCpt automatically fits 12 different models to your data:

- Flat mean (+AR1, +AR2, +Change, +AR1+Change, +AR2+Change)
- Trend mean (+AR1, +AR2, +Change, +AR1+Change, +AR2+Change)

AR1= autoregressive of order 1 = current data point is strongly related to the last data point.

BONUS: Can see which model is best

**PITFALL**: Might be best to use another model which isn't checked - always look at the fit!

### **EnvCpt: Example**

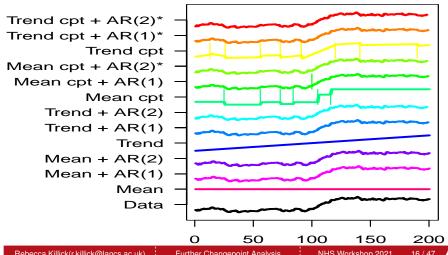


```
set.seed(879123)
x=sim.cpt.AR1(cpts=c(0,100,200),X=cbind(rep(1,200)),init=0,
    beta=rbind(c(0,0.9),c(1,0.9)),sig2=(1-0.9^2),nsim=1)
out=envcpt(x)
## Fitting 12 models
##
which.min(BIC(out))
  meanar1cpt
##
```

## EnvCpt: Example



plot(out)



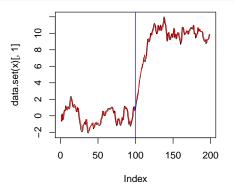
## EnvCpt: Example



cpts(out\$meanar1cpt)

```
## [1] 100
```

```
plot(out[[which.min(BIC(out))+1]])
abline(v=cpts(out$meanar1cpt),col='blue')
```

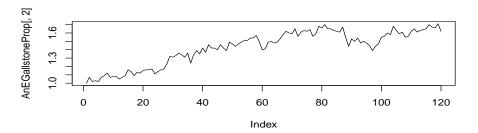


## EnvCpt: A&E Gallstone



HES Data on monthly proportion of A&E admissions for gallstone disease from Jan 2010 - Dec 2019.

```
load('AnEGallstoneProp.Rdata')
plot(AnEGallstoneProp[,2], type='1')
```



Use EnvCpt to see if there is evidence for changes in the monthly proportion of A&E admissions for gallstone disease.

### Gallstone Solution



```
out=envcpt(AnEGallstoneProp[,2])

## Fitting 12 models

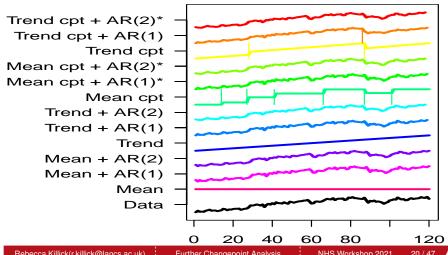
## |
which.min(BIC(out))

## trendar1cpt
## 11
```

### Gallstone Solution



plot(out)

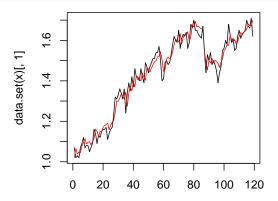


### Gallstone Solution



AnEGallstoneProp\$Date[cpts(out\$trendar1cpt)]

plot(out\$trendar1cpt)



### **Exercise**



Go back to the "Lai2005fig4" data. Use envcpt() to identify the best model. Does this fit with what we observed previously?

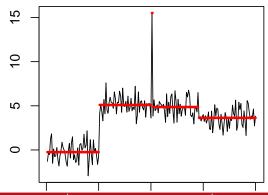
### Influence



- Which data points are influential for obtaining the segmentation?
  - Changepoints versus Outliers
  - How to measure influence?
- How stable is the obtained segmentation?

## Influence: Example

```
set.seed(30)
x=c(rnorm(50),rnorm(50,mean=5),rnorm(1,mean=15),
    rnorm(49,mean=5),rnorm(50,mean=4))
xcpt=cpt.mean(x,method='PELT')
plot(xcpt,cpt.width=3,ylab='')
```



### How to measure?



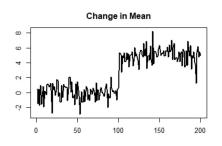
#### Sources of Inspiration:

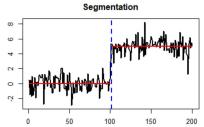
- Regression Analysis: Measures of Influence (e.g., Cook's distance)
- Robust Statistics: Influence Functions

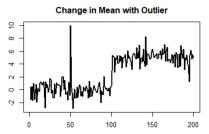
#### Two routes:

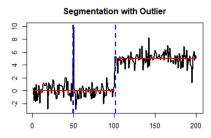
- Modifying an observation
- Leaving out an observation

# Modify









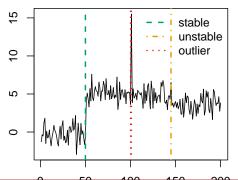
## Stability Dashboard: Out





```
x.inf.out=influence(xcpt,method='outlier')
out.Stability=StabilityOverview(x,cpts(xcpt),x.inf.out,
    legend.args=list(display=TRUE,x="topright",y=NULL,cex=1,
    horiz=FALSE,xpd=FALSE,bty='n'))
```

#### Stability Dashboard: Outlier method

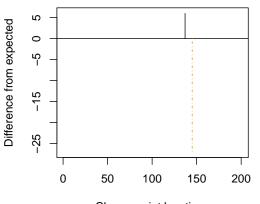


# Location Stability: Out



out.location=LocationStability(cpts(xcpt),x.inf.out,
 type='Difference')

#### **Location Stability: Outlier method**

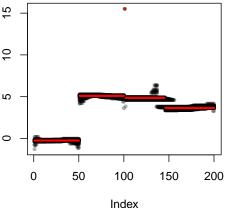


## Parameter Stability: Out



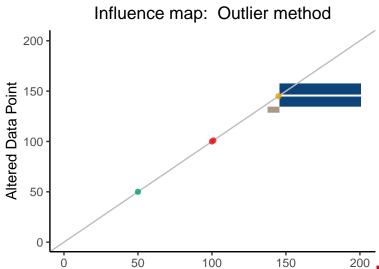
```
ParameterStability(x.inf.out,original.mean=rep(
 param.est(xcpt)$mean,times=diff(c(0,xcpt@cpts))))
```

#### Parameter Stability: Outlier method



# Influence Map: Out

out.map=InfluenceMap(cpts(xcpt),x.inf.out)



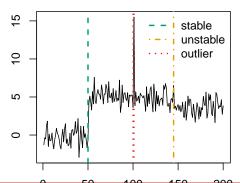
indev

## Stability Dashboard: Del



```
x.inf.del=influence(xcpt,method='delete')
del.Stability=StabilityOverview(x,cpts(xcpt),x.inf.del,
    legend.args=list(display=TRUE,x="topright",y=NULL,cex=1,
    horiz=FALSE,xpd=FALSE,bty='n'))
```

#### Stability Dashboard: Deletion method

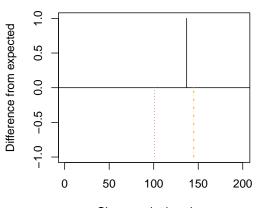


## Location Stability: Del



del.location=LocationStability(cpts(xcpt),x.inf.del,
 type='Difference')

#### Location Stability: Deletion method

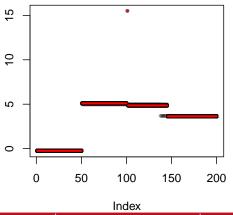


## Parameter Stability: Del



```
ParameterStability(x.inf.del,original.mean=rep(
  param.est(xcpt)$mean,times=diff(c(0,xcpt@cpts))))
```

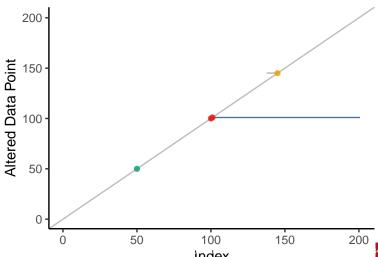
#### Parameter Stability: Deletion method



## Influence Map: Del

del.map=InfluenceMap(cpts(xcpt),x.inf.del)





## Multivariate changes



In moving to the multivariate setting a number of different scenarios could arise.

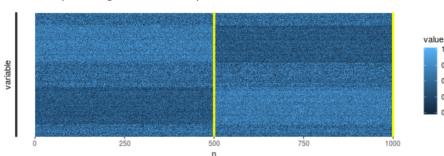
- The process in each channel could be unconnected to the rest (i.e. repeated use of univariate cpt methods might be appropriate);
- There may be some shared structure across channels. For example
  - Changes occur at the same time in all channels;
  - Changes occur in a subset of channels at the same time.
- The nature of the change could vary from one channel to another;
- ... and doubtless many more scenarios!

## Multivariate changes



In the multivariate setting we encounter new challenges:

- Computational expense.
- Sparsity of changepoints.
- Incorporating multivariate power.



1.00

0.75

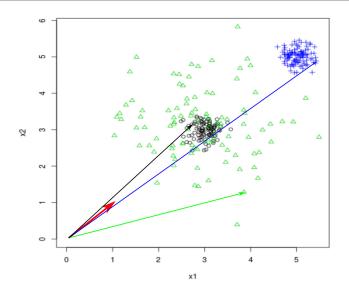
## Many methods



Some well known multivariate changepoint approaches include:

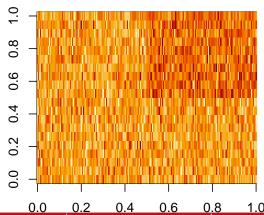
- ecp: James, Matteson (2015)
- Inspect: Wang, Samworth (2017)
- DoubleCUSUM: Cho (2016)

### **GeomCP Intuition**



### GeomCP Mean Ex

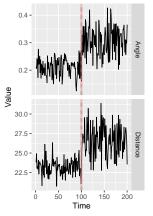




### GeomCP: Mean Ex



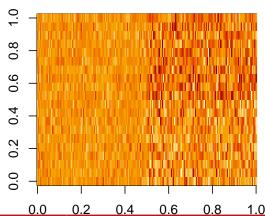
res <- geomcp(Y)
plot(res)</pre>





### GeomCP MeanVar Ex

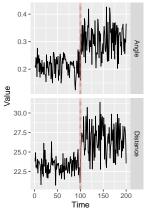




### GeomCP: MeanVar Ex



res <- geomcp(Y)
plot(res)</pre>



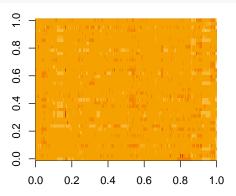


### Task: Genetics



Analyse the ACGH Bladder Tumor data from the ecp package. It is 2215x43 with 43 patients. How many changes do you find?

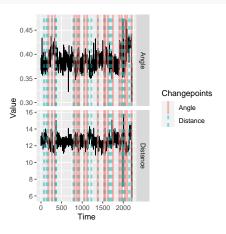
data(ACGH)
image(ACGH\$data)



### Solution: Genetics



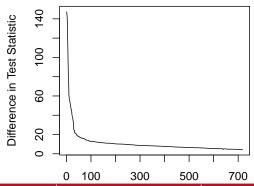
outACGH <- geomcp(ACGH\$data)
plot(outACGH)</pre>



### Solution: Genetics



Can also take the distance and angle vectors and analyse using CROPS



## Summary



- Multivariate is interesting but still lots of challenges in the univariate space
- Lots of interesting research in the changepoint space
- Always looking for interesting problems to work on
- Reach out if you want help / guidance

### References



PELT: Killick, Fearnhead, Eckley (2012)

EnvCpt: Beaulieu, Killick (2018)

geomCP: Grundy, Killick (2020)

Influence: Wilms, Killick, Matteson (2021+)