

Easy Test

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```
library(dplyr)
library(changepoint)
library(fpop)
data(neuroblastoma, package = "neuroblastoma")
```

Get relevant data

```
# Desired id and chromosome to examine.
id <- "4"
chr <- "2"

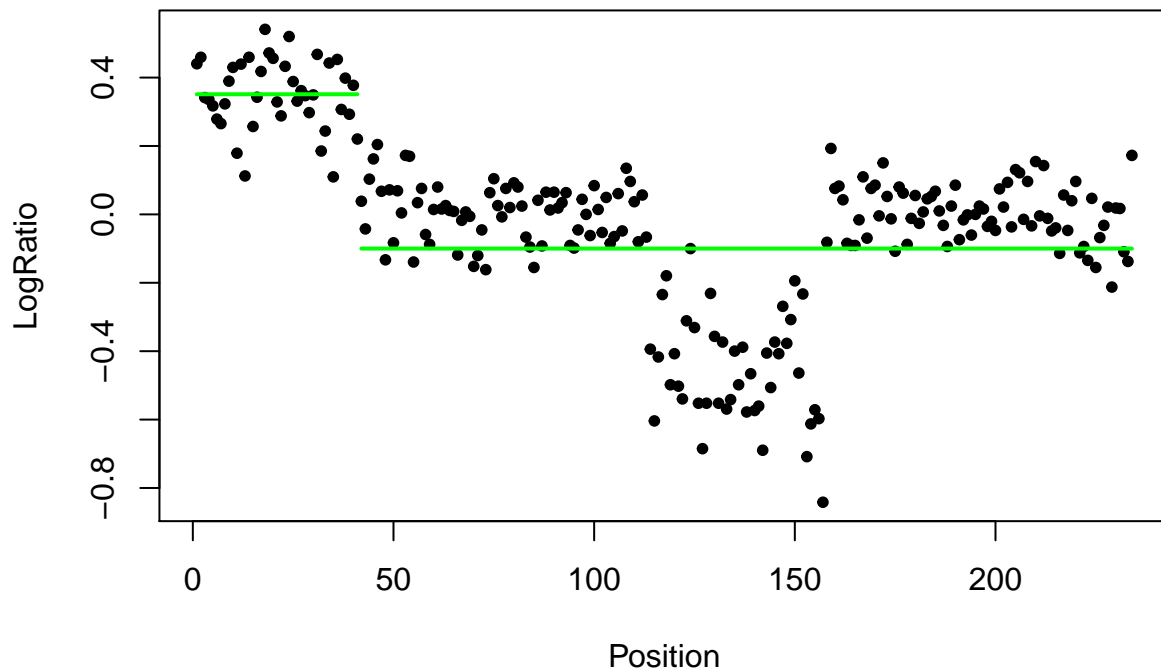
# Filter data for only specified id and chromosome.
profile <- filter(neuroblastoma$profiles,
                  profile.id == id,
                  chromosome == chr)
```

cpt.mean()

```
# Calculate changepoints using changepoint library.
cpt_profile <- cpt.mean(profile$logratio,
                       method="PELT",
                       penalty="Manual",
                       pen.value="log(n)")

plot(cpt_profile,
     type = "p", pch=20, col="black", # Data point params.
     cpt.col="green", cpt.width=2, # Changepoint line params.
     main="Changepoint intervals using cpt.mean()",
     xlab="Position", ylab="LogRatio")
```

Changepoint intervals using cpt.mean()



Fpop()

```
# Calculate changepoints using fpop library.
fpop_profile <- Fpop(profile$logratio, lambda=1)

# Get the starts and ends of each changepoint interval.
seg_end <- fpop_profile$t.est
seg_start <- c(1, seg_end[1:length(seg_end)-1])

# Get mean over each changepoint interval.
seg_mean <- vector(length=length(seg_start))
for(i in 1:length(seg_start)) {
  seg_mean[i] <-
    profile$logratio[seg_start[i]:seg_end[i]] %>%
    mean()
}

# We use NA to avoid jumps in plot.
fpop_segs <- c(rep(seg_mean, times=(seg_end-seg_start)), NA)
fpop_segs[seg_start] <- NA

# Plot data and changepoint segments.
plot(profile$position, profile$logratio,
      type = "p", pch=20, col="black",
      main="Changepoint intervals using Fpop()",
```

```
xlab="Position", ylab="LogRatio")  
  
par(new=TRUE)  
  
lines(profile$position, fpop_segs,  
      col="green", lwd=2)
```

Changepoint intervals using Fpop()

