

# Normalization Example using spike-in control as a standard

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## Introduction

This is a normalisation using linear regression to CTCF of MCF7 cells chip treameant etc.

## Load convience functions

These functions facilitate the normalisation of data.

```
source('../package/brundle.R')
```

## Apply settings

```
dbaSummits          <- 200
jg.controlMinOverlap <- 5
jg.controlSampleSheet <- "samplesheet/samplesheet_SLX8047_dm.csv"
jg.experimentSampleSheet <- "samplesheet/samplesheet_SLX8047_hs.csv"
jg.treatedCondition  = "Fulvestrant"
jg.untreatedCondition = "none"
```

## Load control and experimental DiffBind object

To keep file size down these are provided as a Rdata File rather than as raw counts.

```
filename<-"Rdata/example_003_SLX-8047_dba_HsDm.rda"
if(!file.exists(filename)){
  dbaExperiment <- jg.getDbA(jg.experimentSampleSheet, bRemoveDuplicates=TRUE)
  dbaControl    <- jg.getDbA(jg.controlSampleSheet, bRemoveDuplicates=TRUE)
  save(dbaExperiment,dbaControl,file=filename)
} else {
  load(filename)
}
```

```
#Load Sample Ids from control sample sheet.
```

```
jg.sampleIds <- jg.getSampleIds(jg.controlSampleSheet)
```

```
## Extract Peak set from DiffBind
```

```
jg.experimentPeakset <- jg.dbaGetPeakset(dbaExperiment)
```

```
jg.controlPeakset    <- jg.dbaGetPeakset(dbaControl)
```

```
#Get counts for each condition
```

```
jg.controlCountsTreated<-jg.getControlCounts(jg.controlPeakset,
                                              jg.controlSampleSheet,
                                              jg.treatedCondition)
```

```
jg.controlCountsUntreated<-jg.getControlCounts(jg.controlPeakset,
                                                jg.controlSampleSheet,
                                                jg.untreatedCondition)
```

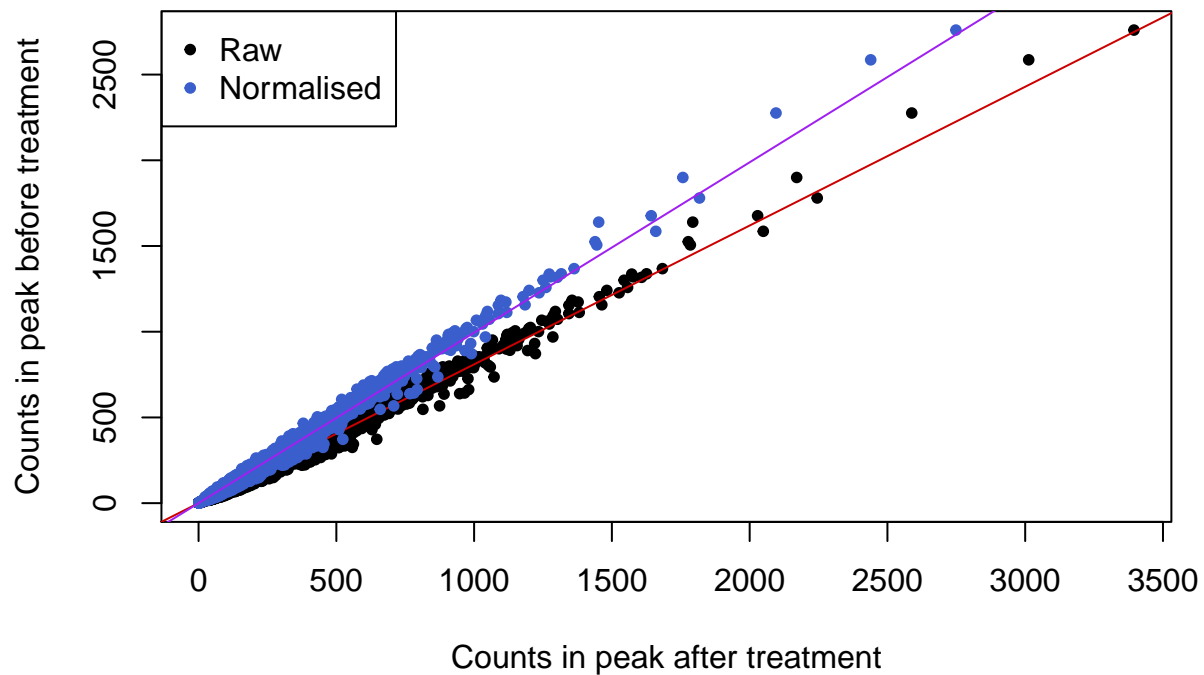
```
#Get sample names for conditions
```

```
jg.untreatedNames <- names(jg.controlCountsUntreated)
jg.treatedNames   <- names(jg.controlCountsTreated)
```

```
##Plot showing normalization calculation (Optional)
```

```
jg.plotNormalization(jg.controlCountsTreated,
                     jg.controlCountsUntreated)
```

## Comparison of Counts in peaks



```
## rowMeans(jg.controlCountsTreated)
##          0.8096594
```

```
##Get Normalization Coefficient
```

```
jg.coefficient<-jg.getNormalizationCoefficient(jg.controlCountsTreated,
                                                jg.controlCountsUntreated)
jg.correctionFactor<-jg.getCorrectionFactor(jg.experimentSampleSheet,
                                             jg.treatedNames,
                                             jg.untreatedNames)
```

```
#Apply coefficient and control factor
```

```
jg.experimentPeaksetNormalised<-jg.applyNormalisation(jg.experimentPeakset,
                                                        jg.coefficient,
                                                        jg.correctionFactor,
                                                        jg.treatedNames)
```

```
#Return values to Diffbind and plot normalised result.
```

```
jg.dba <- DiffBind::pv.resetCounts(dbaExperiment,
                                   jg.experimentPeaksetNormalised)
```

```
jg.dba_analysis<-dba.analyze(jg.dba)
```

```
## converting counts to integer mode
```

```
## [1] "ANH"
```

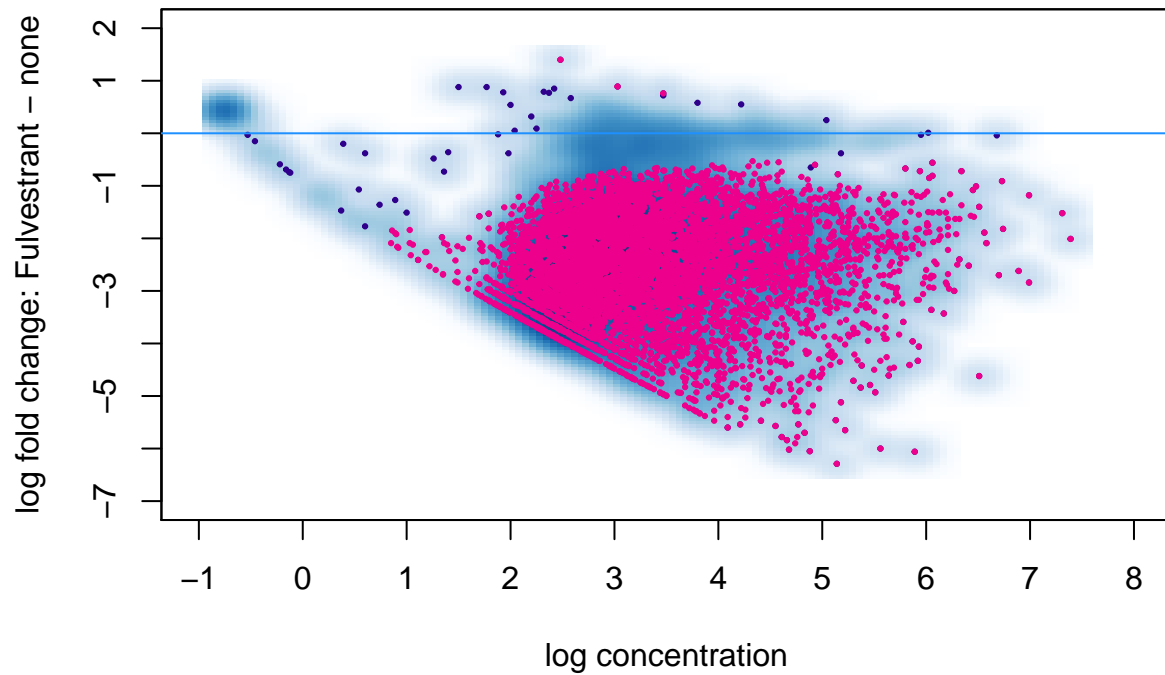
```
## gene-wise dispersion estimates
```

```
## mean-dispersion relationship
```

```
## final dispersion estimates
```

```
dba.plotMA(jg.dba_analysis)
```

### Binding Affinity: Fulvestrant vs. none (5901 FDR < 0.050)



Save results

```
write.csv(dba.report(jg.dba_analysis),file="results/Example_003.csv")
```