

Evaluation and statistics of expression data using NormalyzerDE

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Abstract

Technical biases reduces the ability to see the desired biological changes when performing omics experiments. There are numerous normalization techniques available to counter the biases, but to find the optimal normalization is often a non-trivial task. Furthermore there are limited tools available to counter biases such as retention-time based biases caused by fluctuating electrospray intensities. NormalyzerDE helps this process by performing a wide range of normalization techniques including a general and openly available approach to countering retention-time based biases. More importantly it calculates and visualizes a number of performance measures to guide the users selection of normalization technique. Furthermore, NormalyzerDE provides means to easily perform differential expression statistics using either the empirical Bayes Limma approach or ANOVA. Evaluation visualizations are available for both normalization performance measures and as P-value histograms for the subsequent differential expression analysis comparisons. NormalyzerDE package version: 0.99.8

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1 Installation

Installation is preferably performed using BiocManager (requires R version ≥ 3.5):

```
install.packages("BiocManager")
BiocManager::install("NormalyzerDE")
```

2 Default use

2.1 Citing

Willforss, J., Chawade, A., Levander, F. Submitted article.

2.2 Input format

NormalyzerDE expects a raw data file. Columns can contain annotation information or sample data. Each column should start with a header.

pepseq	s1	s2	s3	s4
ATAAGG	20.0	21.2	19.4	18.5
AWAG	23.3	24.1	23.5	17.3
ACATGM	22.1	22.3	22.5	23.2

This data should be provided with a design matrix where all data samples should be represented. One column (default header "sample") should match the columns containing samples in the raw data. Another column (default header "group") should contain condition levels which could be used for group-based evaluations.

sample	group
s1	condA
s2	condA
s3	condB
s4	condB

Alternatively the data can be provided as an instance of a `SummarizedExperiment` S4 class. This class is available in the `SummarizedExperiment` Bioconductor package.

2.3 Running NormalyzerDE evaluation

The evaluation step can be performed with one command, `normalyzer`. This command expects a path to the data file, a name for the run-job, a path to a design matrix and finally a path to an output directory.

Alternatively the `designPath` and `dataPath` arguments can be replaced with the `experimentObj` argument where the first assay should contain the data matrix of interest, the `colData` attribute the design matrix and the `rowData` attribute the annotation columns.

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```
library(NormalyzerDE)
outDir <- tempdir()
designFp <- system.file(package="NormalyzerDE", "extdata", "design.tsv")
dataFp <- system.file(package="NormalyzerDE", "extdata", "data.tsv")
normalyzer(jobName="vignette_run", designPath=designFp, dataPath=dataFp,
           outputDir=outDir)
## [Step 1/5] Load data and verify input
## Input data checked. All fields are valid.
## Sample check: More than one sample group found
## Sample replication check: All samples have replicates
## RT annotation column found (5)
## [Step 1/5] Input verified, job directory prepared at:/tmp/RtmpLLiGuf/vignette_run
## [Step 2/5] Performing normalizations
## [Step 2/5] Done!
## [Step 3/5] Generating evaluation measures...
## [Step 3/5] Done!
## [Step 4/5] Writing matrices to file
## [Step 4/5] Matrices successfully written
## [Step 5/5] Generating plots...
## [Step 5/5] Plots successfully generated
## All done! Results are stored in: /tmp/RtmpLLiGuf/vignette_run, processing time was 0.7 minutes
```

2.4 Running NormalyzerDE statistical comparisons

When you after performing the evaluation and having evaluated the report have decided for which normalization approach seems to work best you can continue to the statistical step.

Here, expected parameters are the path to the target normalization matrix, the sample design matrix as in the previous step, a job name, the path to an output directory and a list of the pairwise comparisons for which you want to calculate contrasts. They are provided as a character vector with conditions to compare separated by a dash ("-").

Similarly as for the normalization step the `designPath` and `dataPath` arguments can be replaced with an instance of `SummarizedExperiment` sent to the `experimentObj` argument.

```
normMatrixPath <- paste(outDir, "vignette_run/CycLoess-normalized.txt", sep="/")
normalyzerDE("vignette_run",
             designFp,
             normMatrixPath,
             outputDir=outDir,
             comparisons=c("1-2", "1-3"),
             condCol="group")
## [1] "Setting up statistics object"
## [1] "Calculating statistical contrasts..."
## [1] "Contrast calculations done!"
## [1] "Writing 100 annotated rows to /tmp/RtmpLLiGuf/vignette_run/vignette_run_stats.tsv"
## [1] "Writing statis\006tics report"
## [1] "All done! Results are stored in: /tmp/RtmpLLiGuf/vignette_run, processing time was 0 minutes"
```

3 Retention time normalization

Retention time based normalization can be performed with an arbitrary normalization matrix.

3.1 Basic usage

There are two points of access for the higher order normalization. Either by calling `getRTNormalizedMatrix` which applies the target normalization approach stepwise over the matrix based on retention times, or by calling `getSmoothedRTNormalizedMatrix` which generates multiple layered matrices and combines them. To use them you need your raw data loaded into a matrix, a list containing retention times and a normalization matrix able to take a raw matrix and return a normalized in similar format.

```
fullDf <- read.csv(dataFp, sep="\t")
designDf <- read.csv(designFp, sep="\t")
head(fullDf, 1)
##      Cluster.ID Peptide.Sequence External.IDs Charge Average.RT Average.m.z
## 1 1493882053114      AAAAEINVKD      P38156      2   20.25051   501.268
## s_12500amol_1 s_12500amol_2 s_12500amol_3 s_125amol_1 s_125amol_2
## 1   115597000   109302000   100314000   98182352   87241776
## s_125amol_3 s_25000amol_1 s_25000amol_2 s_25000amol_3 s_2500amol_1
## 1   98702800   99315200   185658000   161331008   124908000
## s_2500amol_2 s_2500amol_3 s_250amol_1 s_250amol_2 s_250amol_3
## 1   116218000   112467000   90528896   82128304   90611000
## s_50000amol_1 s_50000amol_2 s_50000amol_3 s_5000amol_1 s_5000amol_2
## 1   155924000   132508000   139174000   110084000   107755000
## s_5000amol_3 s_500amol_1 s_500amol_2 s_500amol_3 s_50amol_1 s_50amol_2
## 1   108817000   98170000   96898704   104109000   88484096   92873168
## s_50amol_3
## 1   97952512
head(designDf, 1)
##      sample group batch
## 1 s_50amol_1      1      1
```

At this point we have loaded the full data into dataframes. Next, we use the sample names present in the design matrix to extract sample columns from the raw data. Be careful that the sample names is a character vector. If it is a factor it will extract wrong columns.

Make sure that sample names extracted from design matrix are in right format. We expect it to be in 'character' format.

```
sampleNames <- as.character(designDf$sample)
typeof(sampleNames)
## [1] "character"
```

Now we are ready to extract the data matrix from the full matrix. We also need to get the retention time column from the full matrix.

```
dataMat <- as.matrix(fullDf[, sampleNames])
retentionTimes <- fullDf$Average.RT
```

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```
head(dataMat, 1)
##      s_50amol_1 s_50amol_2 s_50amol_3 s_125amol_1 s_125amol_2 s_125amol_3
## [1,] 88484096  92873168  97952512   98182352   87241776   98702800
##      s_250amol_1 s_250amol_2 s_250amol_3 s_500amol_1 s_500amol_2 s_500amol_3
## [1,] 90528896   82128304   90611000   98170000   96898704   104109000
##      s_2500amol_1 s_2500amol_2 s_2500amol_3 s_5000amol_1 s_5000amol_2
## [1,] 124908000   116218000   112467000   110084000   107755000
##      s_5000amol_3 s_12500amol_1 s_12500amol_2 s_12500amol_3 s_25000amol_1
## [1,] 108817000   115597000   109302000   100314000   99315200
##      s_25000amol_2 s_25000amol_3 s_50000amol_1 s_50000amol_2 s_50000amol_3
## [1,] 185658000   161331008   155924000   132508000   139174000
```

If everything is fine the data matrix should be `double`, and have the same number of rows as the number of retention time values we have.

```
typeof(dataMat)
## [1] "double"

print("Rows and columns of data")
## [1] "Rows and columns of data"
dim(dataMat)
## [1] 100 27

print("Number of retention times")
## [1] "Number of retention times"
length(retentionTimes)
## [1] 100
```

The normalization function is expected to take a raw intensity matrix and return log transformed values. We borrow the wrapper function for Loess normalization from NormalyzerDE. It can be replaced with any custom function as long as the wrapper has the same input/output format.

```
performCyclicLoessNormalization <- function(rawMatrix) {
  log2Matrix <- log2(rawMatrix)
  normMatrix <- limma::normalizeCyclicLoess(log2Matrix, method="fast")
  colnames(normMatrix) <- colnames(rawMatrix)
  normMatrix
}
```

We are ready to perform the normalization.

```
rtNormMat <- getRTNormalizedMatrix(dataMat,
                                  retentionTimes,
                                  performCyclicLoessNormalization,
                                  stepSizeMinutes=1,
                                  windowMinCount=100)
```

Let's double check the results. We expect a matrix in the same format and shape as before. Furthermore, we expect similar but not the exact same values as if we'd applied the normalization globally.

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```
globalNormMat <- performCyclicLoessNormalization(dataMat)
dim(rtNormMat)
## [1] 100 27
dim(globalNormMat)
## [1] 100 27
head(rtNormMat, 1)
## s_50amol_1 s_50amol_2 s_50amol_3 s_125amol_1 s_125amol_2 s_125amol_3
## 26.4195 26.49648 26.60364 26.53692 26.31423 26.54111
## s_250amol_1 s_250amol_2 s_250amol_3 s_500amol_1 s_500amol_2 s_500amol_3
## 26.3765 26.32032 26.41595 26.44011 26.43208 26.59369
## s_2500amol_1 s_2500amol_2 s_2500amol_3 s_5000amol_1 s_5000amol_2
## 26.94727 26.85807 26.76096 26.69457 26.70806
## s_5000amol_3 s_12500amol_1 s_12500amol_2 s_12500amol_3 s_25000amol_1
## 26.68261 26.76658 26.70266 26.59615 26.71226
## s_25000amol_2 s_25000amol_3 s_50000amol_1 s_50000amol_2 s_50000amol_3
## 27.24251 27.0744 27.00559 26.78887 26.85155
head(globalNormMat, 1)
## s_50amol_1 s_50amol_2 s_50amol_3 s_125amol_1 s_125amol_2 s_125amol_3
## [1,] 26.4195 26.49648 26.60364 26.53692 26.31423 26.54111
## s_250amol_1 s_250amol_2 s_250amol_3 s_500amol_1 s_500amol_2 s_500amol_3
## [1,] 26.3765 26.32032 26.41595 26.44011 26.43208 26.59369
## s_2500amol_1 s_2500amol_2 s_2500amol_3 s_5000amol_1 s_5000amol_2
## [1,] 26.94727 26.85807 26.76096 26.69457 26.70806
## s_5000amol_3 s_12500amol_1 s_12500amol_2 s_12500amol_3 s_25000amol_1
## [1,] 26.68261 26.76658 26.70266 26.59615 26.71226
## s_25000amol_2 s_25000amol_3 s_50000amol_1 s_50000amol_2 s_50000amol_3
## [1,] 27.24251 27.0744 27.00559 26.78887 26.85155
```

3.2 Performing layered normalization

We have everything set up to perform the layered normalization. The result here is expected to be overall similar to the normal retention time approach.

```
layeredRtNormMat <- getSmoothedRTNormalizedMatrix(
  dataMat,
  retentionTimes,
  performCyclicLoessNormalization,
  stepSizeMinutes=1,
  windowMinCount=100,
  windowShifts=3,
  mergeMethod="mean")

dim(layeredRtNormMat)
## [1] 100 27
head(layeredRtNormMat, 1)
## s_50amol_1 s_50amol_2 s_50amol_3 s_125amol_1 s_125amol_2 s_125amol_3
## [1,] 26.4195 26.49648 26.60364 26.53692 26.31423 26.54111
## s_250amol_1 s_250amol_2 s_250amol_3 s_500amol_1 s_500amol_2 s_500amol_3
## [1,] 26.3765 26.32032 26.41595 26.44011 26.43208 26.59369
```

```
##      s_2500amol_1 s_2500amol_2 s_2500amol_3 s_5000amol_1 s_5000amol_2
## [1,]      26.94727      26.85807      26.76096      26.69457      26.70806
##      s_5000amol_3 s_12500amol_1 s_12500amol_2 s_12500amol_3 s_25000amol_1
## [1,]      26.68261      26.76658      26.70266      26.59615      26.71226
##      s_25000amol_2 s_25000amol_3 s_50000amol_1 s_50000amol_2 s_50000amol_3
## [1,]      27.24251      27.0744      27.00559      26.78887      26.85155
```

4 Stepwise processing (normalization part)

NormalyzerDE consists of a set of steps. The workflow can be run as a whole, or step by step.

4.1 Step 1: Loading data

This step performs input validation of the data, and generates an object of the class `NormalyzerDataset`.

```
jobName <- "vignette_run"
experimentObj <- setupRawDataObject(dataFp, designFp, "default", TRUE, "sample", "group")
normObj <- getVerifiedNormalyzerObject(jobName, experimentObj)
## Input data checked. All fields are valid.
## Sample check: More than one sample group found
## Sample replication check: All samples have replicates
## RT annotation column found (5)
```

The function `setupRawDataObject` returns a `SummarizedExperiment` object. This object can be prepared directly and should in that case contain the raw data as the default assay, the design matrix as `colData` and annotation rows as `rowData`.

4.2 Step 2: Generate normalizations

Here, normalizations are performed. This generates a `NormalyzerResults` object containing both a reference to its original dataset object, but also generated normalization matrices.

```
normResults <- normMethods(normObj)
```

4.3 Step 3: Generate performance measures

Performance measures are calculated for normalizations. These are stored in an object `NormalizationEvaluationResults`. A `NormalyzerResults` object similar to the one sent in is returned, but with this field added.

```
normResultsWithEval <- analyzeNormalizations(normResults)
```


4.4 Step 4: Output matrices to file

Generated normalization matrices are written to the provided folder.

```
jobDir <- setupJobDir("vignette_run", tempdir())  
writeNormalizedDatasets(normResultsWithEval, jobDir)
```

4.5 Step 5: Generate evaluation plots

Performance measures are used to generate evaluation figures which is written in an evaluation report.

```
generatePlots(normResultsWithEval, jobDir)  
## pdf  
## 2
```

After this evaluation of normalizations and progression to statistics follows as described previously in this report.

5 Stepwise processing (differential expression analysis part)

5.1 Step 1: Setup folders and data matrices

For continued processing you select the matrix containing the normalized data from the best performing normalization. The design matrix is the same as for the normalization step.

```
bestNormMatPath <- paste(jobDir, "RT-Loess-normalized.txt", sep="/")  
experimentObj <- setupRawContrastObject(bestNormMatPath, designFp, "sample")  
nst <- NormalyzerStatistics(experimentObj,  
                           logTrans=FALSE,  
                           leastRepCount=2,  
                           conditionCol="group")
```

Similarly as to for the normalization evaluation step the `experimentObj` above can be prepared directly as a `SummarizedExperiment` object.

5.2 Step 2: Calculate statistics

Now we are ready to perform the contrasts. Contrasts are provided as a vector in the format `c("condA-condB", "condB-condC")`, where `condX` is the group levels.

```
comparisons <- c("1-2", "2-3")  
nst <- calculateContrasts(nst, comparisons, condCol="group")
```

5.3 Step 3: Generate final matrix and output

Finally we generate a table containing the statistics results for each feature and write it to file together with an evaluation report containing P-value histograms for each comparison.

```
annotDf <- generateAnnotatedMatrix(nst)
utils::write.table(annotDf, file=paste(jobDir, "stat_table.tsv", sep="/"))
generateStatsReport(nst, "Vignette stats", jobDir)
## pdf
## 2
```

6 Code organization

NormalyzerDE consists of a number of scripts and classes. They are focused around two separate workflows. One is for normalizing and evaluating the normalizations. The second is for performing differential expression analysis. Classes are contained in scripts with the same name.

Code organization:

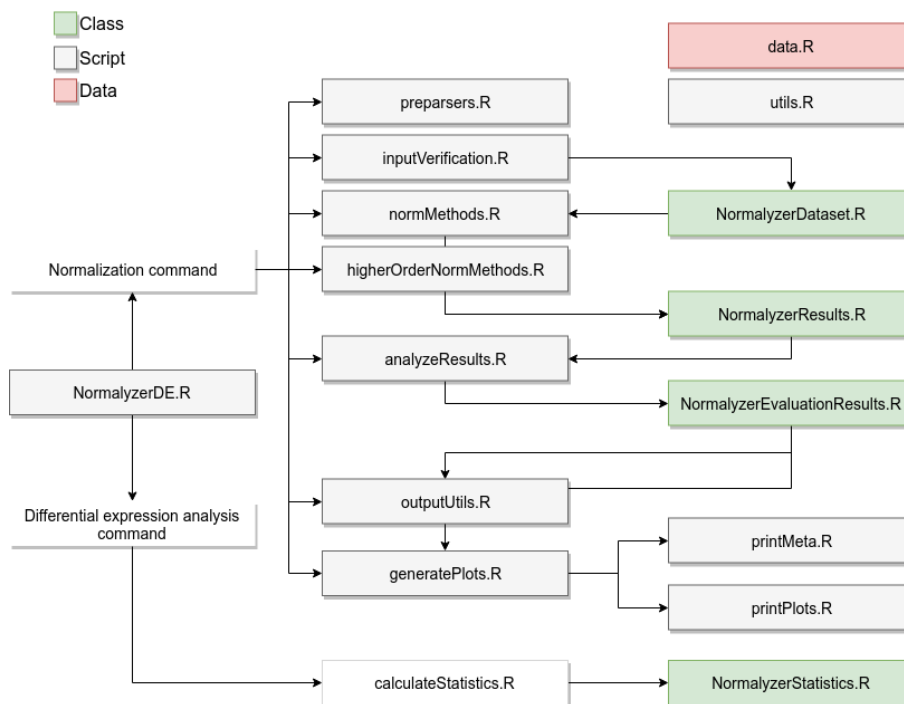


Figure 1: NormalyzerDE schematics

The standard workflow for the normalization is the following:

- The `normalyzer` function in the `NormalyzerDE.R` script is called, starting the process.
- If applicable (that is, input is in Proteois or MaxQuant format), the dataset is preprocessed into the standard format using code in `preparers.R`.

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- The input is verified to capture standard errors early on using code in `inputVerification.R`. This results in an instance of the `NormalyzerDataset` class.
- The data is normalized using several normalization methods present in `normMethods.R`. This yields an instance of `NormalyzerResults` which links to the original `NormalyzerDataset` instance and also contains all the resulting normalized datasets.
- If specified (and if a column with retention time values is present) retention-time segmented approaches are performed by applying normalizations from `normMethods.R` over retention time using functions present in `higherOrderNormMethods.R`.
- The results are analyzed using functions present in `analyzeResults.R`. This yields an instance of `NormalyzerEvaluationResults` containing the evaluation results. This instance is attached to the `NormalyzerResults` object.
- The final results are sent to `outputUtils.R` where the normalizations are written to an output directory, and to `generatePlots.R` which contains visualizations for the performance measures. It also uses code in `printMeta.R` and `printPlots.R` to output the results in a desired format.

When a normalized matrix is selected the analysis proceeds to the statistical analysis.

- The `normalyzerde` function in the `NormalyzerDE.R` script is called starting the differential expression analysis pipeline.
- An instance of `NormalyzerStatistics` is prepared containing the input data.
- Code in the `calculateStatistics.R` script is used to calculate the statistical contrasts. The results are attached to the `NormalyzerStatistics` object.
- The resulting statistics are used to generate a report and an annotated output matrix where key statistical measures are attached to the original matrix.