PowerExplorer Manual

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Abstract

This vignette demonstrates R package PowerExplorer as a power and sample size estimation tool for RNA-Seq and quantitative proteomics data.

PowerExplorer contains the following main features:

- Estimation of power based on the current data
- Prediction of power corresponding to the increased sample sizes
- Result visualizations

Introduction

Power and sample size estimation is one of the important principles in designing next-generation sequencing experiments to discover differential expressions. PowerExplorer is a power estimation and prediction tool currently applicable to RNA-Seq and quantitative proteomics experiments.

The calculation procedure starts with estimating the distribution parameters of each gene or protein (following referred as feature for simplicity). With the obtained prior distribution of each feature, a specified amount of simulations are executed to generate data (read counts for RNA-Seq and protein abundance for proteomics) repetitively for each entry based on null and alternative hypotheses. Furthermore, the corresponding statistical tests (t-test or Wald-test) are performed and the test statistics are collected. Eventually the statistics will be summarized to calculate the statistical power.

Prepare Input Data

For both RNA-Seq (gene expression levels) and quantitative proteomics (protein abundance levels) datasets, the data matrix should be arranged as features in rows and samples in columns, for example:

```
library(PowerExplorer)
data("exampleRNASeqData")
head(exampleRNASeqData$dataMatrix[,1:6])
#>
          Sample\_A\_1 Sample\_A\_2 Sample\_A\_3 Sample\_A\_4 Sample\_A\_5 Sample\_B\_1
#> Gene_1
                  469
                              324
                                          38
                                                     1059
                                                                   64
                                                                              737
#> Gene 2
                   84
                              276
                                          263
                                                      182
                                                                  181
#> Gene 3
                  293
                              173
                                          272
                                                      123
                                                                  475
                                                                              169
                                                                             1064
                  310
#> Gene_4
                              209
                                          550
                                                      212
                                                                  394
                                                                              293
#> Gene_5
                   82
                              141
                                          216
                                                      202
                                                                  494
                               98
#> Gene_6
                  583
                                          137
                                                      179
                                                                              884
                                                                  214
```

A grouping vector indicating the sample groups to which all the samples belong should also be created, for example:

The sample groups corresponding to the data:

```
colnames(exampleProteomicsData$dataMatrix)
#> [1] "Sample_A_1" "Sample_A_2" "Sample_A_3" "Sample_A_4" "Sample_A_5"
#> [6] "Sample_B_1" "Sample_B_2" "Sample_B_3" "Sample_B_4" "Sample_B_5"
#> [11] "Sample_C_1" "Sample_C_2" "Sample_C_3" "Sample_C_4" "Sample_C_5"
```

Note that the grouping vector length should be equal to the column number of the data matrix.

Run Estimation

Here we use a randomly generated RNASeq dataset exampleRNASeqData as an example to estimate the current power of the dataset. The input dataset is named as dataMatrix and the grouping vector as groupVec.

To run the estimation, apart from the input, we still need to specify the following parameters:

- isLogTransformed: FALSE; the input data is not log-transformed.
- dataType: "RNA-Seq"; the datatype can be declared as "Proteomics" or "RNA-Seq".
- minlfc: 0.5; the threshold of Log2 Fold Change, proteins with lower LFC will be discarded.
- alpha: 0.05; the controlled false positive (Type I Error) rate.
- ST: 50; the simulation of each gene will be run 50 times (ST>50 is recommended).
- seed: 345; optional, a seed value for the random number generator to maintain the reproducibility.
- showProcess: FALSE; no detailed processes will be shown, set to TRUE if debug is needed.
- saveSimulatedData: FALSE; if TRUE, save the simulated data in ./savedData directory.

The results will be summaried in barplot, boxplot and summary table.

```
library(PowerExplorer)
data("exampleRNASeqData")
res <- estimatePower(inputObject = exampleRNASeqData$dataMatrix,</pre>
                    groupVec = exampleRNASeqData$groupVec,
                    isLogTransformed = FALSE,
                     dataType = "RNASeq",
                    minLFC = 0.5,
                     alpha = 0.05,
                    ST = 50,
                    seed = 345)
#> ##---- Mon Feb 19 13:31:44 2018 ----##
#> Num. of groups:
#> Num. of replicates:
#> Num. of simulations: 50
#> Min. Log Fold Change:
                             0.5
#> False Postive Rate:
                        0.05
#> Transformed: FALSE
#> ##---- Mon Feb 19 13:31:44 2018 ----##
#> 0 of 110 entries are filtered due to excessive zero counts
#> Estimating distribution parameters...
#> Estimating NB parameters by DESeq2...
#>
#> [A.vs.B] 14 of 110 genes are over minLFC threshold 0.5:
#>
#> [A.vs.B] Log2 Fold Change Quantiles:
#> 0% 10% 20% 30% 40% 50% 60% 70% 80% 90% 100%
#> 0.00 0.05 0.08 0.13 0.16 0.20 0.26 0.31 0.39 0.58 1.80
#>
#> [A.vs.C] 17 of 110 genes are over minLFC threshold 0.5:
#> [A.vs.C] Log2 Fold Change Quantiles:
#> 0% 10% 20% 30% 40% 50% 60% 70% 80% 90% 100%
#> 0.01 0.04 0.08 0.12 0.18 0.23 0.27 0.33 0.46 0.89 2.19
#> [B.vs.C] 16 of 110 genes are over minLFC threshold 0.5:
```

```
#> [B.vs.C] Log2 Fold Change Quantiles:

#> 0% 10% 20% 30% 40% 50% 60% 70% 80% 90% 100%

#> 0.00 0.03 0.08 0.12 0.15 0.23 0.27 0.33 0.41 0.56 2.18

#>

** Simulation in process, it may take a few minutes...

#> Power Estimation between groups A.vs.B:

#>

#> OVERALL ESTIMATED POWER: 0.4986

#>

*> Simulation in process, it may take a few minutes...

#>

#> Power Estimation between groups A.vs.C:

#>

#> OVERALL ESTIMATED POWER: 0.5894

#>

*> Simulation in process, it may take a few minutes...

#>

#> Power Estimation between groups B.vs.C:

#>

#> Power Estimation between groups B.vs.C:

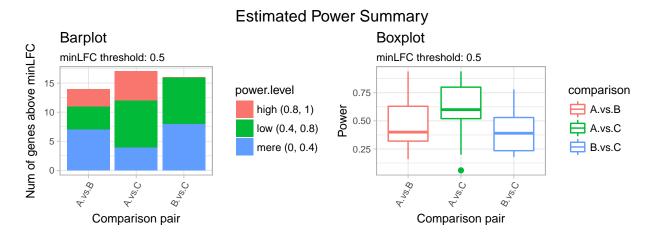
#>

#> OVERALL ESTIMATED POWER: 0.4275
```

Visualization

The estimated results can be summarized using plotEstPwr, the only input needed is the estimatedPower, which should be the estimated power object returned from estimatePower.

plotEstPwr(res)



Comp.	Gene Num.	Avg. Power	H (0.8, 1)	L (0.4, 0.8)	M (0, 0.4)
A.vs.B	14	0.50	3 (21%)	4 (29%)	7 (50%)
A.vs.C	17	0.59	5 (29%)	8 (47%)	4 (24%)
B.vs.C	16	0.43	0 (0%)	8 (50%)	8 (50%)

The graph contains 3 plots, the barplot vertically shows the number of genes/proteins above the minLFC threshold, columns indicates the comparison pairs, each column presents the proportions of three power levels in three colours as indicated in the legend power.level; The boxplot shows the overall power distribution of each comparsion; And the summary table summarize the power in a numerical way with the same information shown in the previous two plots.

Run Predictions

With the same dataset, to run a prediction, a different parameter is needed:

• rangeSimNumRep: the power of replicate number 5 to 20 will be predicted.

Similar to the estimation process, however, the simulations will be excuted with each sample size specified in rangeSimNumRep. (Note: the term sample size in this vignette refers to the replicate number of each group/case)

It is possible to append the prediction results within the same object by using the same result object as an input.

```
data("exampleRNASeqData")
res <- predictPower(inputObject = res,</pre>
                   groupVec = exampleRNASeqData$groupVec,
                   isLogTransformed = FALSE,
                   dataType = "RNASeq",
                   rangeSimNumRep = c(5, 10, 15, 20),
                   minLFC = 1,
                   alpha = 0.05,
                   ST = 50,
                   seed = 345)
#> ##---- Mon Feb 19 13:32:08 2018 -----##
#> Num. of groups:
                    3
#> Num. of replicates:
                        5, 10, 15, 20
#> Num. of simulations: 50
#> Min. Log Fold Change:
#> False Postive Rate:
                        0.05
#> Transformed: FALSE
#>
#> ##---- Mon Feb 19 13:32:08 2018 -----##
#> 0 of 110 entries are filtered due to excessive zero counts
#> Estimating distribution parameters...
#> Estimating NB parameters by DESeg2...
#>
#> [A.vs.B] 4 of 110 genes are over minLFC threshold 1:
#>
#> [A.vs.B] Log2 Fold Change Quantiles:
#> 0% 10% 20% 30% 40% 50% 60% 70% 80% 90% 100%
#> 0.00 0.05 0.08 0.13 0.16 0.20 0.26 0.31 0.39 0.58 1.80
#> [A.vs.C] 10 of 110 genes are over minLFC threshold 1:
#> [A.vs.C] Log2 Fold Change Quantiles:
   0% 10% 20% 30% 40% 50% 60% 70% 80% 90% 100%
#> 0.01 0.04 0.08 0.12 0.18 0.23 0.27 0.33 0.46 0.89 2.19
#> [B.vs.C] 5 of 110 genes are over minLFC threshold 1:
#> [B.vs.C] Log2 Fold Change Quantiles:
#> 0% 10% 20% 30% 40% 50% 60% 70% 80% 90% 100%
#> 0.00 0.03 0.08 0.12 0.15 0.23 0.27 0.33 0.41 0.56 2.18
```

```
#> ##--Simulation with 5 replicates per group--##
#>
#> [repNum:5] Simulation in process, it may take a few minutes...
#>
#> [repNum:5] Power Estimation between groups A.vs.B:
#>
#> OVERALL ESTIMATED POWER: 0.73
#>
#>
#> [repNum:5] Simulation in process, it may take a few minutes...
#>
#> [repNum:5] Power Estimation between groups A.vs.C:
#>
#> OVERALL ESTIMATED POWER: 0.65
#>
#>
#> [repNum:5] Simulation in process, it may take a few minutes...
#> [repNum:5] Power Estimation between groups B.vs.C:
#>
#> OVERALL ESTIMATED POWER: 0.66
#>
#>
#> ##--Simulation with 10 replicates per group--##
#> [repNum:10] Simulation in process, it may take a few minutes...
#>
#> [repNum:10] Power Estimation between groups A.vs.B:
#>
#> OVERALL ESTIMATED POWER: 0.96
#>
#>
#> [repNum:10] Simulation in process, it may take a few minutes...
#>
#> [repNum:10] Power Estimation between groups A.vs.C:
#>
#> OVERALL ESTIMATED POWER: 0.882
#>
#>
#> [repNum:10] Simulation in process, it may take a few minutes...
#>
#> [repNum:10] Power Estimation between groups B.vs.C:
#>
#> OVERALL ESTIMATED POWER: 0.872
#>
#>
#> ##--Simulation with 15 replicates per group--##
#> [repNum:15] Simulation in process, it may take a few minutes...
#> [repNum:15] Power Estimation between groups A.vs.B:
#> OVERALL ESTIMATED POWER: 0.995
```

```
#>
#>
#> [repNum:15] Simulation in process, it may take a few minutes...
#> [repNum:15] Power Estimation between groups A.vs.C:
#>
#> OVERALL ESTIMATED POWER: 0.966
#>
#>
#> [repNum:15] Simulation in process, it may take a few minutes...
#> [repNum:15] Power Estimation between groups B.vs.C:
#>
#> OVERALL ESTIMATED POWER: 0.964
#>
#>
#> ##--Simulation with 20 replicates per group--##
#> [repNum:20] Simulation in process, it may take a few minutes...
#>
#> [repNum:20] Power Estimation between groups A.vs.B:
#>
#> OVERALL ESTIMATED POWER: 1
#>
#>
#> [repNum:20] Simulation in process, it may take a few minutes...
#>
#> [repNum:20] Power Estimation between groups A.vs.C:
#>
#> OVERALL ESTIMATED POWER: 0.99
#>
#>
#> [repNum:20] Simulation in process, it may take a few minutes...
#>
#> [repNum:20] Power Estimation between groups B.vs.C:
#>
#> OVERALL ESTIMATED POWER: 0.984
```

Visualization

The predicted results can be summaried using plotPredPwr. The input should be the predicted power object returned from predictPower, the summary can be optionally visualized by setting the following parameters:

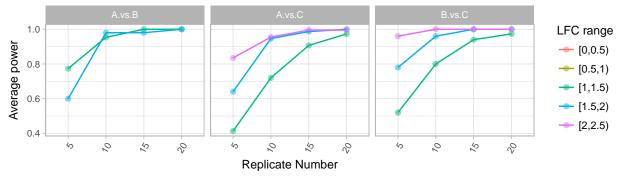
- PEObject: A PEObject returned from PowerExplorer as input
- minLFC and maxLFC: to observe power in a specific range of LFC
- LFCscale: to determine the LFC scale of the observation

Lineplot (LFCscale = 0.5):

plotPredPwr(res, LFCscale = 0.5)

Average Predicted Power within LFC ranges

segmented by every 0.5 Log2FoldChange (minLFC: 0, maxLFC: 2)



	repNum:5	repNum:10	repNum:15	repNum:20
A.vs.B	0.69	0.97	0.99	1
A.vs.C	0.63	0.87	0.96	0.99
B.vs.C	0.75	0.92	0.98	0.99

Lineplot is one of the optional outputs of plotPredPwr, the output contains a lineplot and a summary table. For each comparison, the lineplot shows the power tendency across every Log2 Fold Change segment resulted from a complete LFC list divided by a specified LFCscale. Each dot on the lines stands for the average power (y-axis) of the genes within the LFC range (x-axis), and each colour indicates the average power of a certain sample size (as shown in the legend besides the plot). In addition, a summary table below displays the average power of each comparison across the sample sizes.

For instance, the line plot here shows the average power of four sample sizes (5 to 30, with increment of 5) in LFCscale of 0.5. The LFC ranges from 0 to 5, and within each LFC segment, the graph shows the average power of the features. Here, the higher LFC shows higher power, the average power of each LFC range increases with the larger sample sizes, as expected.