

Differential Expression Meta-Analysis with DExMA package

Juan Antonio Villatoro-García^{1,2} and Pedro Carmona-Sáez*

¹Department of Statistics and Operational Research. University of Granada

²Bioinformatics Unit. GENYO, Centre for Genomics and Oncological Research

*pedro.carmona@genyo.es

April 28, 2026

Abstract

DExMA (Differential Expression Meta-Analysis) performs all the necessary steps of differential expression meta-analysis considering the possible existence of missing genes. It controls the existence of unmeasured genes by two approaches: considering the genes that are contained in a proportion of datasets (set by the users) or imputing the expression values of the unmeasured genes using the *Knn method* in the space of samples. In addition, it allows to apply quality controls, download GEO datasets and show a graphical representation of the results.

packageVersionDExMA 1.20.0

Contents

1	Introduction	3
2	Previous steps: Meta-analysis object	3
2.1	Meta-analysis object creation (objectMA)	4
2.2	Adding a new dataset to the meta-analysis object	7
3	Performing Meta-analysis	9
3.1	Gene annotation and quality controls	9
3.1.1	Setting all the datasets in the same annotation	9
3.1.2	Logarithm transformation	11
3.1.3	Heterogeneity study	12
3.2	Imputing missing genes expression values (optional).	13
3.3	Performing meta-analysis: <i>metaAnalysisDE()</i>	14
3.3.1	Effects size combination results	16
3.3.2	P-value combination results	17
3.4	Visualization of the results: heatmap	18

- 4 Additional information 23
 - 4.1 GEO microarray data download 23
 - 4.2 Using RNA-Seq data 24
 - 4.3 Removing Batch Effects 24
 - 4.4 Calculating Effects size 26
 - 4.5 Calculating Individual P-values 27
- 5 Session info 27

1 Introduction

DExMA is a package designed to perform gene expression meta-analysis. Gene expression meta-analysis comprises a set of methods that combine the results of several differential expression studies into a single common result [1]. Furthermore, this package has the advantage that it allows to take into account those genes that are not measured in a study. The first approach to control these missing genes, **DExMA** offers two options: apply the k-nearest neighbors method (knn method) in the space of samples to impute the expression values of these missing genes [2]. The second option is to take into account those genes that are contained in at least a certain proportion (set by the user) of datasets. The use of only the genes common to all the datasets could lead to the loss of some genes that are not measured in a single study, which would lead to the loss of information. Due to this fact, **DExMA** package is very useful to work with Microarray data, because this type of data is what usually produces the existence of uncommon genes between datasets with different annotations. However, previously normalized RNA-Seq data can also be used, as well as, both Microarray data and normalized RNA-Seq data at the same time.

DExMA package has implemented methods from the three main types of gene expression meta-analysis [1] meta-analysis based on effects sizes combination and meta-analysis based on p-values combination. Once one of the methods has been applied, **DExMA** package provides a specific and adapted results table. Moreover, this package contains some functions that allows the user to carry out a previous quality control in order to results obtained are more reliable. Finally, this package provides some additional function that, for example, help the user to download public Microarray data from NCBI GEO public database [3] or to visualize the significant genes in a heatmap.

This document gives a tutorial-style introduction to all the steps that must be carry out in order to properly perform gene expression meta-analysis by making use of **DExMA** package.

2 Previous steps: Meta-analysis object

DExMA uses a specific object as input, which is a list of nested lists where each nested list corresponds to a study. This object can be created directly by the users or they can use `createObjectMA()` function to create it.

For the examples that are going to be shown, synthetic data will be used. We load the sample data into our R session.

```
> library(DExMA)
> data("DExMAExampleData")
```

- `listMatrixEX`: a list of four expression arrays
- `listPhenodatas`: a list of the four phenodata corresponding to the four expression arrays
- `listExpression`: a list of four `ExpressionSets` object. It contains the same information as `listMatrixEX` and `listPhenodatas`
- `ExpressionSetStudy5`: an `ExpressionSet` object similar to the `ExpressionSets` objects of `listExpression`.
- `maObjectDif`: the meta-analysis object created from the `listMatrixEX` and `listPhenodatas` objects.

- `maObject`: the meta-analysis object after setting all the studies in Official Gene Symbol annotation

2.1 Meta-analysis object creation (`objectMA`)

As previously stated, the meta-analysis input in DExMA is a list of nested lists. Each nested list contains two elements:

- A gene expression matrix with genes in rows and samples in columns
- A vector of 0 and 1 indicating the group of each sample. 0 represents reference group (usually controls) and 1 represents experimental group (usually cases).

This object can be created directly by the user or we can make use of `createObjectMA()` function, which creates the `*objectMA*` after indicating how the reference and experimental groups are identified.

`createObjectMA()` function allows to create the object needed to perform meta-analysis. In this case, it is necessary to indicate as input of the function the variables that contain the experimental and reference groups:

- `listEX`: a list of dataframes or matrix (genes in rows and samples in columns). A list of `ExpressionSets` can be used too:

```
> #List of expression matrices
> data("DExMAExampleData")
> ls(listMatrixEX)

[1] "Study1" "Study2" "Study3" "Study4"

> head(listMatrixEX$Study1)

      Sample1 Sample2 Sample3 Sample4
100859927 5.439524 6.253319 2.926444 4.4304023
8086      5.769823 5.971453 1.831349 4.0466288
8212      7.558708 5.957130 2.365252 3.4352889
65985     6.070508 7.368602 2.971158 3.7151784
729522    6.129288 5.774229 3.670696 3.9171749
13        7.715065 7.516471 1.349453 0.3390772

> #List of ExpressionSets
> ls(listExpressionSets)

[1] "Study1" "Study2" "Study3" "Study4"

> listExpressionSets$Study1

ExpressionSet (storageMode: lockedEnvironment)
assayData: 200 features, 4 samples
  element names: exprs
protocolData: none
phenoData
  rowNames: Sample1 Sample2 Sample3 Sample4
  varLabels: condition gender organism race
  varMetadata: labelDescription
featureData: none
```

```
experimentData: use 'experimentData(object)'
Annotation:
```

- `listPheno`: a list of phenodatas (samples in rows and covariables in columns). If the object `listEX` is a list of `ExpressionSets` this element can be null.

```
> data("DExMAExampleData")
> #Example of a phenodata object
> ls(listPhenodatas)

[1] "Study1" "Study2" "Study3" "Study4"

> listPhenodatas$Study1

      condition gender    organism race
Sample1  Diseased Female Homo Sapiens  AA
Sample2  Diseased Female Homo Sapiens  AA
Sample3   Healthy Female Homo Sapiens  AA
Sample4   Healthy Female Homo Sapiens   H
```

- `namePheno`: a list or vector of the different column names or column positions from the pheno used for performing the comparison between groups. Each element of `namePheno` correspond to its equivalent element in the `listPheno`. (default a vector of 1, all the first columns of each elements of `listPheno` are selected)
- `expGroups`: a list or vector of the group names or positions from `namePheno` variable used as experimental group (cases) to perform the comparison (default a vector of 1, all the first groups are selected).
- `refGroups`: a list or vector of the group names or positions from `namePheno` variable used as reference group (controls) to perform the comparison (default a vector of 2, all the second groups are selected).

It is important to note that if any element does not belong to the experimental or the reference group, that sample is not taken into account in the creation of meta-analysis object.

Here, we have included an example to show how exactly the function is used:

Since this function can be a bit complicated if there are many datasets, we recommend creating a vector to keep the column names of the phenodatas that contains the variable that identifies the groups to compare (*namePheno* argument). Moreover, we should create two others lists to indicate how to identify experimental (cases) and reference (controls) groups in these variables (*expGroups* and *refGroups* arguments).

If we look at the example phenodatas list we have the following four objects:

```
> listPhenodatas$Study1

      condition gender    organism race
Sample1  Diseased Female Homo Sapiens  AA
Sample2  Diseased Female Homo Sapiens  AA
Sample3   Healthy Female Homo Sapiens  AA
Sample4   Healthy Female Homo Sapiens   H
```

In the "Study1" phenoData, the groups variable is "condition". Experimental group is named as "Diseased" and reference group as "Healthy".

DExMA package

```
> listPhenodatas$Study2
```

	condition	gender	organism	race
Sample5	Diseased	Female	Homo Sapiens	AA
Sample6	Diseased	Female	Homo Sapiens	AA
Sample7	ill	Male	Homo Sapiens	C
Sample8	Healthy	Female	Homo Sapiens	H
Sample9	control	Female	Homo Sapiens	H
Sample10	control	Male	Homo Sapiens	H

In the "Study2" phenoData, the groups variable is "condition". Experimental group is named as "Diseased" or "ill" and reference group as "Healthy" or "control"

```
> listPhenodatas$Study3
```

	state	gender	organism	race
Sample11	Diseased	Female	Homo Sapiens	AA
Sample12	Diseased	Female	Homo Sapiens	AA
Sample13	Healthy	Female	Homo Sapiens	AA
Sample14	Healthy	Female	Homo Sapiens	H

In the "Study3" phenoData, the groups variable is "state". Experimental group is named as "Diseased" and reference group as "Healthy".

```
> listPhenodatas$Study4
```

	state	gender	organism	race
Sample15	ill	Female	Homo Sapiens	AA
Sample16	ill	Female	Homo Sapiens	AA
Sample17	ill	Male	Homo Sapiens	AA
Sample18	control	Female	Homo Sapiens	C
Sample19	control	Female	Homo Sapiens	H
Sample20	control	Male	Homo Sapiens	H

In this phenoData, the groups variable is "state". Experimental group is named as "ill" and reference group as "control".

We all this information we can create the vector for *namePheno* argument and the two list for *expGroups* and *refGroups*:

```
> phenoGroups = c("condition", "condition", "state", "state")
> phenoCases = list(Study1 = "Diseased", Study2 = c("Diseased", "ill"),
+                   Study3 = "Diseased", Study4 = "ill")
> phenoControls = list(Study1 = "Healthy", Study2 = c("Healthy", "control"),
+                      Study3 = "Healthy", Study4 = "control")
```

Then, we can apply more easily *createObjectMA()* function:

```
> newObjectMA <- createObjectMA(listEX=listMatrixEX,
+                               listPheno = listPhenodatas,
+                               namePheno=phenoGroups,
+                               expGroups=phenoCases,
+                               refGroups = phenoControls)
> #Study 1
```

```

> head(newObjectMA[[1]][[1]])
      Sample1 Sample2 Sample3 Sample4
100859927 5.439524 6.253319 2.926444 4.4304023
8086      5.769823 5.971453 1.831349 4.0466288
8212      7.558708 5.957130 2.365252 3.4352889
65985     6.070508 7.368602 2.971158 3.7151784
729522    6.129288 5.774229 3.670696 3.9171749
13        7.715065 7.516471 1.349453 0.3390772

> newObjectMA[[1]][[2]]
[1] 1 1 0 0

> #Study 2
> head(newObjectMA[[2]][[1]])
      Sample5 Sample6 Sample7 Sample8 Sample9 Sample10
100859927 4.367690 6.648252 5.786897 2.820073 2.688687 3.331881
8086      5.937004 5.914434 7.124650 3.234439 3.228559 4.521235
8212      5.294553 4.092231 5.194514 3.368330 2.750491 5.369983
65985     5.685822 5.900363 5.539580 2.153294 3.166798 3.162232
729522    5.733054 5.553966 6.592146 3.275042 2.659061 2.452524
13        6.153159 5.435909 6.991360 4.860632 3.751608 3.121853

> newObjectMA[[2]][[2]]
[1] 1 1 1 0 0 0

```

The result obtained is the proper object to perform meta-analysis (objectMA).

2.2 Adding a new dataset to the meta-analysis object

It may happen that once the meta-analysis object is created we want to add a new dataset before doing the meta-analysis. **DExMA** provides the `elementObjectMA()` function, which allows the creation of an element of the meta-analysis object. This function contains the following arguments:

- `expressionMatrix`: a dataframe or matrix that containing genes in rows and samples in columns. An `ExpressionSet` object can be used too.
- `pheno`: a data frame or a matrix containing samples in rows and covariates in columns. If `NULL` (default), `pheno` is extracted from the `ExpressionSet` object
- `groupPheno`: the column name or position from `pheno` where experimental group (cases) and reference group (control) are identified
- `expGroups`: a vector of the names or positions from `groupPheno` variable used as experimental group (cases). By default the first group (character) is taken
- `refGroups`: a vector of the names or positions from `groupPheno` variable used as reference group (control). By default the second group (character) is taken.

As with the `createObjectMA()` function, if any element does not belong to the experimental or the reference group, that sample is not included in the creation of the object.

Here we provided an example of the use of this function, in which we create an element of the meta-analysis object from the information of the "Study 2" of the `listExpressionSets` object:

DExMA package

```
> data("DExMAExampleData")
> ExpressionSetStudy5

ExpressionSet (storageMode: lockedEnvironment)
assayData: 200 features, 6 samples
  element names: exprs
protocolData: none
phenoData
  rowNames: newSample1 newSample2 ... newSample6 (6 total)
  varLabels: condition gender organism race
  varMetadata: labelDescription
featureData: none
experimentData: use 'experimentData(object)'
Annotation:

> library(Biobase)
> pData(ExpressionSetStudy5)

      condition gender  organism race
newSample1  Diseased Female Homo Sapiens  AA
newSample2  Diseased Female Homo Sapiens  AA
newSample3    ill    Male Homo Sapiens    C
newSample4  Healthy Female Homo Sapiens    H
newSample5  control Female Homo Sapiens    H
newSample6  control  Male Homo Sapiens    H
```

We had to load [Biobase](#) package in order to the information in the *ExpressionSet*. In the *phenoData* we can observe that groups variable is "condition". In addition, Experimental group is name as "Diseased" or "ill" and reference group as "Healthy" or "control"

```
> newElem <- elementObjectMA(expressionMatrix = ExpressionSetStudy5,
+                             groupPheno = "condition",
+                             expGroup = c("Diseased", "ill"),
+                             refGroup = c("Healthy", "control"))
> head(newElem[[1]])

      newSample1 newSample2 newSample3 newSample4 newSample5 newSample6
100859927  5.867690  8.148252  7.286897  4.320073  4.188687  4.831881
8086      7.437004  7.414434  8.624650  4.734439  4.728559  6.021235
8212      6.794553  5.592231  6.694514  4.868330  4.250491  6.869983
65985     7.185822  7.400363  7.039580  3.653294  4.666798  4.662232
729522    7.233054  7.053966  8.092146  4.775042  4.159061  3.952524
13        7.653159  6.935909  8.491360  6.360632  5.251608  4.621853

> head(newElem[[2]])

[1] 1 1 1 0 0 0
```

As we can see, we obtain a list that has the same structure as the elements of the meta-analysis object (*objectMA*). This new element can be added to a *objectMA* that has been created previously:

```
> newObjectMA2 <- newObjectMA
> newObjectMA2[[5]] <- newElem
```



```
> head(newObjectMA2[[5]][[1]])
```

	newSample1	newSample2	newSample3	newSample4	newSample5	newSample6
100859927	5.867690	8.148252	7.286897	4.320073	4.188687	4.831881
8086	7.437004	7.414434	8.624650	4.734439	4.728559	6.021235
8212	6.794553	5.592231	6.694514	4.868330	4.250491	6.869983
65985	7.185822	7.400363	7.039580	3.653294	4.666798	4.662232
729522	7.233054	7.053966	8.092146	4.775042	4.159061	3.952524
13	7.653159	6.935909	8.491360	6.360632	5.251608	4.621853

```
> newObjectMA2[[5]][[2]]
```

```
[1] 1 1 1 0 0 0
```

Moreover, an advantage of this function is that it can be used to create one by one all the elements and finally join all of them to create the meta-analysis object.

3 Performing Meta-analysis

DExMA package contains the main gene expression meta-analysis methods:

- Meta-analysis based on effect size combination: Fixed Effects Model (FEM) and Random Effects Model (REM).
- Meta-analysis based on P-value combination: Fisher's method, Stouffer's method, Wilkonson's method (maxP), Tippet's method (minP) and Aggregated Cauchy Association Test method (ACAT).

These methods can be applied directly, but it is advisable to apply some previous **DExMA** function to ensure the results are accurate.

3.1 Gene annotation and quality controls

Before performing the meta-analysis, all the genes must be in the same annotation and a quality control should be done in order to obtain reliable results [1]. **DExMA** provides some useful functions to help the user to do it.

3.1.1 Setting all the datasets in the same annotation

All genes must be in the same annotation in order to perform the meta-analysis successfully and avoid incorrect interpretations of the results. In cases where all datasets do not have the same gene ID, **DExMA** contains a function called *allSameID()* that allows to annotated all the datasets with the same gene ID. Specifically, the function allows to annotate those datasets that use the Official Gene Symbol, Entrez or Ensembl. The inputs of this function are:

- *objectMA*: the meta-analysis object of **DExMA** package. The result obtained by *createObjectMA()* function should be used.
- *finalID*: a character that indicates the final gene ID that all the studies will have.
- *organism*: a character that indicates the organism that the datasets belong to. Use *availableOrganism* function to see the organism admitted.

DExMA package

Here we include an example of how to use this function. In this example we have used "newObjectMA" that has been created before. The first two expression arrays are annotated in "entrez", the third expression matrix in "Official Gene Symbol" and the last one in "Official Gene Symbol" with some synonyms:

```
> rownames(newObjectMA$Study1$mExpres)[1:20]

[1] "100859927" "8086"      "8212"      "65985"     "729522"    "13"
[7] "344752"    "126767"    "343066"    "51166"     "79719"     "22848"
[13] "14"        "15"        "16"        "57505"     "80755"     "132949"
[19] "60496"     "10157"

> rownames(newObjectMA$Study2$mExpres)[1:20]

[1] "100859927" "8086"      "8212"      "65985"     "729522"    "13"
[7] "344752"    "126767"    "343066"    "51166"     "79719"     "22848"
[13] "14"        "15"        "16"        "57505"     "80755"     "132949"
[19] "60496"     "10157"

> rownames(newObjectMA$Study3$mExpres)[1:20]

[1] "AAA4"      "AAAS"      "AABT"      "AACS"      "AACSP1"    "AADAC"
[7] "AADACL2"   "AADACL3"   "AADACL4"   "AADAT"     "AAGAB"     "AAK1"
[13] "AAMP"      "AANAT"     "AARS1"     "AARS2"     "AARSD1"    "AASDH"
[19] "AASDHPPT" "AASS"

> rownames(newObjectMA$Study4$mExpres)[1:20]

[1] "AAA4"      "ADRACALIN" "AABT"      "ACSF1"     "AACSP1"    "AADAC"
[7] "AADACL2"   "AADACL3"   "AADACL4"   "AADAT"     "PPKP1"     "AAK1"
[13] "AAMP"      "AANAT"     "AARS1"     "AARS2"     "AARSD1"    "AASDH"
[19] "AASDHPPT" "AASS"
```

We can use *availableIDs* and *availableOrganism* in order to know how to write in the function the *finalID* and the *organism* arguments:

```
> head(availableIDs)

[1] "Entrez"      "Ensembl"    "GeneSymbol"

> availableOrganism

[1] "Bos taurus"          "Caenorhabditis elegans"
[3] "Canis familiaris"    "Danio rerio"
[5] "Drosophila melanogaster" "Gallus gallus"
[7] "Homo sapiens"        "Mus musculus"
[9] "Rattus norvegicus"   "Arabidopsis thaliana"
[11] "Saccharomyces cerevisiae" "Escherichia coli"
```

Then, we are ready to run the *allSameID()* function. We are going to annotated all the datasets in Official Gene Symbol (*finalID*="GeneSymbol"):

```
> newObjectMA <- allSameID(newObjectMA, finalID="GeneSymbol", organism = "Homo sapiens")

|
|
|
|=====| 25%
```

```

|
|=====| 50%
|
|=====| 75%
|
|=====| 100%

> rownames(newObjectMA$Study1$mExpres)[1:20]

[1] "AAA4" "AAAS" "AABT" "AACS" "AACSP1" "AADAC"
[7] "AADACL2" "AADACL3" "AADACL4" "AADAT" "AAGAB" "AAK1"
[13] "AAMP" "AANAT" "AARS1" "AARS2" "AARSD1" "AASDH"
[19] "AASDHPPT" "AASS"

> rownames(newObjectMA$Study2$mExpres)[1:20]

[1] "AAA4" "AAAS" "AABT" "AACS" "AACSP1" "AADAC"
[7] "AADACL2" "AADACL3" "AADACL4" "AADAT" "AAGAB" "AAK1"
[13] "AAMP" "AANAT" "AARS1" "AARS2" "AARSD1" "AASDH"
[19] "AASDHPPT" "AASS"

> rownames(newObjectMA$Study3$mExpres)[1:20]

[1] "AARS1" "AATF" "ABCC2" "ABCD1P4" "ABCD1P3" "ABCD1P2" "ACAD8"
[8] "AAVS1" "ACAD9" "ABT1" "VSX1" "ABHD5" "AADAT" "ABI3"
[15] "VRK3" "VPS54" "ABAT" "VSI10" "VRTN" "VPS53"

> rownames(newObjectMA$Study4$mExpres)[1:20]

[1] "AARS1" "AATF" "ABCC2" "ABCD1P4" "ABCD1P3" "ABCD1P2" "ACAD8"
[8] "AAVS1" "ACAD9" "ABT1" "VSX1" "ABHD5" "AADAT" "ABI3"
[15] "VRK3" "VPS54" "ABAT" "VSI10" "VRTN" "VPS53"

```

As it can be seen, all the studies are now annotated in Official Gene Symbol.

3.1.2 Logarithm transformation

To avoid problems with the returned fold-change by the meta-analysis, \log_2 should be applied to the gene expression values. We can make use of `dataLog()` function to check if each dataset expression values have the \log_2 applied already. If not, the function will make the transformation:

```

> newObjectMA <- dataLog(newObjectMA)
> head(newObjectMA[[1]][[1]])

      Sample1 Sample2 Sample3 Sample4
AAA4  5.439524 6.253319 2.926444 4.4304023
AAAS  5.769823 5.971453 1.831349 4.0466288
AABT  7.558708 5.957130 2.365252 3.4352889
AACS  6.070508 7.368602 2.971158 3.7151784
AACSP1 6.129288 5.774229 3.670696 3.9171749
AADAC  7.715065 7.516471 1.349453 0.3390772

```

3.1.3 Heterogeneity study

Some heterogeneity between studies may lead to some methods, such as the Fixed Effects Model, not providing reliable results. Therefore, it is advisable to carry out a study of heterogeneity to correctly choose the meta-analysis method [1]. The *heterogeneityTest()* function shows two ways of measuring heterogeneity.

On the one hand, it returns a QQ-plot of the Cochran's test [4]. In this plot, if most of the values are close to the central line, that is, most of the Cochran's test values are close to the expected distribution (chi-squared distribution), it can be said that there is homogeneity. In the case that these values deviate greatly from the expected distribution, it must be assumed that there is heterogeneity.

On the other hand, I^2 measures the percentage of variation across studies due to heterogeneity [5]. In the case of gene expression data, an I^2 for each gene across datasets would have to be calculated. As in the case of many genes, it can be difficult to observe all the I^2 values obtained, the *heterogeneityTest()* function returns the quantiles of the different I^2 values calculated. I^2 values equal to 0 indicate homogeneity and values less than 0.25 are usually categorized as low heterogeneity [5]. Therefore, to assume homogeneity in the gene expression meta-analysis, almost all I^2 values must be 0 or at least less than 0.25.

In the example shown below, it is observed that in the QQ-plot of the Cochran's test, Q-values deviate considerably from the expected distribution and approximately 10% of the I^2 values are greater than 0.25, therefore homogeneity could not be assumed.

```
> heterogeneityTest(newObjectMA)

[1] "I^2 Quantiles"
      0%      25%      50%      75%
0.0000000 0.0000000 0.4484812 0.6865470
```



3.2 Imputing missing genes expression values (optional)

Optionally, the `missGenesImput()` function allows to impute the expression values of the unmeasured genes. To perform this imputation, the function apply the knn method in the space of samples (*sampleKnn*) [2]. Once the function have been applied, it returns the same *objectMA* with all the datasets imputed and some imputation indicators:

```
> nrow(newObjectMA$Study1[[1]])
[1] 144
> nrow(newObjectMA$Study2[[1]])
[1] 144
> imputation <- missGenesImput(maObject, k = 7)
[1] "Number of values imputed 310 (0.089 %)"
[1] "Number of genes imputed in Study1: 31 of 175 (17.7%)"
[1] "Number of genes imputed in Study2: 31 of 175 (17.7%)"
```

```
[1] "Number of genes imputed in Study3: 0 of 175 (0%)"
[1] "Number of genes imputed in Study4: 0 of 175 (0%)"
> maObject_imput <- imputation$objectMA
```

If we observe the number of genes in Study1, once the imputation is performed, we can observe that it goes from having 144 genes to 175.

```
> nrow(maObject_imput$Study1[[1]])
[1] 175
> nrow(maObject_imput$Study2[[1]])
[1] 175
```

In addition, the function returns some indicators to evaluate the reliability of the imputation:

- `imputValuesSample`: Number of missing values imputed per sample
- `imputPercentageSample`: Percentage of missing values imputed per sample
- `imputValuesGene`: Number of missing values imputed per gene
- `imputPercentageGene`: Percentage of missing values imputed per gen

```
> head(imputation$imputIndicators$imputValuesSample)
Sample1 Sample2 Sample3 Sample4 Sample5 Sample6
      31      31      31      31      31      31
> head(imputation$imputIndicators$imputPercentageSample)
Sample1 Sample2 Sample3 Sample4 Sample5 Sample6
17.71429 17.71429 17.71429 17.71429 17.71429 17.71429
> head(imputation$imputIndicators$imputValuesGene)
AAA4  AAAS  AABT  AACS AACSP1  AADAC
    0     0     0     0     0     0
> head(imputation$imputIndicators$imputPercentageGene)
AAA4  AAAS  AABT  AACS AACSP1  AADAC
    0     0     0     0     0     0
```

3.3 Performing meta-analysis: *metaAnalysisDE()*

The *metaAnalysisDE()* function allows to perform a meta-analysis in only one step, needing only the meta-analysis object created previously.

This function has as input:

- `objectMA`: The meta-analysis object of DExMA package. The result obtained by *createObjectMA* function should be used.
- `effectS`: A list of three elements. The first element is a dataframe with genes in rows and studies in columns. Each component of the dataframe is the effect of a gene in a study. The second element of the list is also a dataframe with the same structure, but in this case each component of the dataframe represent the

variance of the effect of a gene in a study. The result of the `calculateES` function can be used too. The third element of the list is also a dataframe with the same structure, but in this case each component of the dataframe represent the log fold change of a gene in a study. This argument should be only used in the case that `objectMA` argument is null.

- `pvalues`: A list of two elements. The first element is a dataframe with genes in rows and studies in columns. Each component of the dataframe is the p-value of a gene in a study. The second element of the list is also a dataframe with the same structure, but in this case each component of the dataframe represent the log fold change of a gene in a study. This argument should be only used in the case that `objectMA` argument is null.
- `weight`: A vector of the weights of each dataset. This argument should only be included in case `objectMA` is null and you want to use "Stouffer" or "ACAT" method.
- `typeMethod`: a character that indicates the method to be performed:
 - "FEM": Fixed Effects model.
 - "REM": Random Effects model.
 - "Fisher": Fisher's method (sum of logarithms of p-values)
 - "Stouffer": Stouffer's method (sum of z-scores)
 - "maxP": Wilkinson's method (maximun of p-values)
 - "minP": Tippet's method (minimun of p-values)
 - "ACAT": Aggregated Cauchy Association Test method
- `missAllow`: a number between 0 and 1 that indicates the maximum proportion of missing values allows in a sample. If the sample has more proportion of missing values, the sample will be eliminated. In the other case, the missing values will be imputed by using the K-NN algorithm included in [impute](#) package [6]. In case the `objectMA` has been previously imputed, this element is not necessary.
- `proportionData`: a number between 0 and 1 that indicates the minimum proportion of datasets in which a gene must be contained to be included. In case the `objectMA` has been previously imputed, this element is not necessary.
- Adjusted p-value from which a gene is considered significant. Default 0.05. This value only takes you into account in the results generated in rank combination methods.

In the following example, we have applied a Random Effect model to the `DExMA` object ("newObjectMA") we have been working with so far. In addition we have allowed a 0.3 proportion of missing values in a sample and a gene must have been contained in at least the 50% of studies.

```
> resultsMA <- metaAnalysisDE(newObjectMA, typeMethod="REM",  
+                               missAllow=0.3, proportionData=0.50)  
[1] "Performing Random Effects Model"
```

The output of this function is a dataframe with the results of the meta-analysis where rows are the genes and columns are the different variables provided by the meta-analysis:

```
> head(resultsMA)
```

	Com.ES	ES.var	Qval	Qpval	tau2	Zval	Pval
AAA4	2.170422	0.5769960	7.390096	0.06045064	1.2841381	2.857312	4.272453e-03
AAAS	2.164285	0.2161334	1.348614	0.71762287	0.0000000	4.655362	3.234118e-06
AABT	2.502180	0.8198233	7.905694	0.04800146	1.9149389	2.763493	5.718633e-03
AACS	2.256773	0.4634824	5.424099	0.14324946	0.8093607	3.314906	9.167395e-04
AACSP1	3.685009	0.4675196	2.393007	0.49493773	0.0000000	5.389378	7.070214e-08
AADAC	2.641814	0.9830839	9.279712	0.02579411	2.3971453	2.664446	7.711514e-03

	FDR	AveFC	propDataset
AAA4	1.967577e-02	2.734153	1
AAAS	9.432844e-05	3.163741	1
AABT	2.357305e-02	2.746359	1
AACS	7.292246e-03	3.009776	1
AACSP1	1.237287e-05	3.088231	1
AADAC	2.998922e-02	2.897440	1

Moreover, this function can also be applied to the imputed *objectMA*, but in this case, it is not necessary to indicate the *missAllow* and *proportionData* elements:

```
> resultsMA_imput <- metaAnalysisDE(maObject_imput, typeMethod="REM",)

[1] "Performing Random Effects Model"

> head(resultsMA_imput)
```

	Com.ES	ES.var	Qval	Qpval	tau2	Zval	Pval
AAA4	2.170422	0.5769960	7.390096	0.06045064	1.2841381	2.857312	4.272453e-03
AAAS	2.164285	0.2161334	1.348614	0.71762287	0.0000000	4.655362	3.234118e-06
AABT	2.502180	0.8198233	7.905694	0.04800146	1.9149389	2.763493	5.718633e-03
AACS	2.256773	0.4634824	5.424099	0.14324946	0.8093607	3.314906	9.167395e-04
AACSP1	3.685009	0.4675196	2.393007	0.49493773	0.0000000	5.389378	7.070214e-08
AADAC	2.641814	0.9830839	9.279712	0.02579411	2.3971453	2.664446	7.711514e-03

	FDR	AveFC	propDataset
AAA4	3.367599e-02	2.734153	1
AAAS	9.432844e-05	3.163741	1
AABT	3.706521e-02	2.746359	1
AACS	1.234072e-02	3.009776	1
AACSP1	1.237287e-05	3.088231	1
AADAC	4.641638e-02	2.897440	1

The variables of the dataframe change from one type of meta-analysis to another. A more detailed explanation of these results will be addressed in the following sections.

3.3.1 Effects size combination results

The "FEM" and "REM" methods provide a dataframe with the variables:

- Com.ES: combined effect of the gene.
- ES.var: variance of the combined effect of the gene.
- Qval: total variance of the gene.
- Qpval:p-value for the total variance of the gene.

- tau2: between-study variance of the gene.
- zval: combined effect value for a standard normal. I can be use in order to find out if the gene is overexpressed (positive value) or underexpressed (negative value).
- Pval: P-value of the meta-analysis for the gene.
- FDR: P-value adjusted of the meta-analysis for the gene.
- AveFC: Weighted average of log Fold-Change values for the gene used in order to find out if the gene is overexpressed (positive value) or underexpressed (negative value).
- Prop.dataset: Proportion of the datasets in which the gene is included.

```
> resultsES <- metaAnalysisDE(newObjectMA, typeMethod="REM", proportionData=0.5)
[1] "Performing Random Effects Model"
> head(resultsES)
```

	Com.ES	ES.var	Qval	Qpval	tau2	Zval	Pval
AAA4	2.170422	0.5769960	7.390096	0.06045064	1.2841381	2.857312	4.272453e-03
AAAS	2.164285	0.2161334	1.348614	0.71762287	0.0000000	4.655362	3.234118e-06
AABT	2.502180	0.8198233	7.905694	0.04800146	1.9149389	2.763493	5.718633e-03
AACS	2.256773	0.4634824	5.424099	0.14324946	0.8093607	3.314906	9.167395e-04
AACSP1	3.685009	0.4675196	2.393007	0.49493773	0.0000000	5.389378	7.070214e-08
AADAC	2.641814	0.9830839	9.279712	0.02579411	2.3971453	2.664446	7.711514e-03
	FDR	AveFC	propDataset				
AAA4	1.967577e-02	2.734153	1				
AAAS	9.432844e-05	3.163741	1				
AABT	2.357305e-02	2.746359	1				
AACS	7.292246e-03	3.009776	1				
AACSP1	1.237287e-05	3.088231	1				
AADAC	2.998922e-02	2.897440	1				

3.3.2 P-value combination results

The "Fisher", "Stouffer", "minP", "maxP" and "ACAT" methods provide a dataframe with the following variables:

- Stat: Statistical calculated in the method
- Pval: P-value of the meta-analysis for the gene.
- FDR: P-value adjusted of the meta-analysis for the gene.
- AveFC: Average of log Fold-Change values for the gene used in order to find out if the gene is overexpressed (positive value) or underexpressed (negative value).
- Prop.dataset: Proportion of the datasets in which the gene is included.

Here we present an example making use of "maxP" method:

```
> resultsPV <- metaAnalysisDE(newObjectMA, typeMethod="maxP", proportionData=0.5)
[1] "Performing MaxP's method"
> head(resultsPV)
```

	Stat	Pval	FDR	AveFC	propDataset
AAA4	0.13754548	3.579194e-04	3.296626e-03	2.831428	1
AAAS	0.07824187	3.747632e-05	6.811215e-04	3.180932	1
AABT	0.27659986	5.853395e-03	2.768497e-02	2.935515	1
AACS	0.11561455	1.786693e-04	1.737063e-03	3.060848	1
AACSP1	0.01862270	1.202736e-07	1.052394e-05	2.953126	1
AADAC	0.22367842	2.503204e-03	1.510554e-02	3.629187	1

3.4 Visualization of the results: heatmap

Finally, we can represent in a heatmap the significant genes in order to observe how they are expressed in each of the studies. In `makeHeatmap()` function we have to include both the object that has been used in the meta-analysis, the result of it and the applied method. In addition, this package offers three different scaling approaches (*scaling*) in order to compare properly the gene expression of the studies in the heatmap:

- "zscor": It calculates a z-score value for each gene, that is, the mean gene expression from each gene is subtracted from each gene expression value and then it is divided by the standard deviation.
- "swr": Scaling relative to reference dataset approach [7].
- "rscale": It uses the `rescale` function of the `scales` package to scale the gene expression [8].
- "none": no scaling approach is applied.

Moreover, in *regulation* argument, we can choose if we want to represent the overexpressed or underexpressed genes:

- "up": only up-expressed genes are represented.
- "down": only down-expressed genes are represented
- "all": up-expressed and down-expressed genes are represented.

We can choose the number of significant genes (*numSig*) that we want to be shown on the graph and the adjusted p-value from which a gene is considered as significant (*fdrSig*). In addition, the genes that are not presented in one sample are represented in gray.

Here we present an example of the heatmap which have been obtained from the result of applying a random effects model to the object "newObjectMA" and making use of a "zscor" scaling approach.

```
> makeHeatmap(objectMA=newObjectMA, resMA=resultsMA, scaling = "zscor",
+             regulation = "all", numSig=40,
+             fdrSig = 0.05, logFCSig = 1.5, show_rownames = TRUE)

[1] "scaling using z-score..."
      Sample1.Study1 Sample2.Study1 Sample3.Study1 Sample4.Study1
AARS2           0.9682999       0.50830923      -1.3357281      -0.1408811
AADACL3         0.1962999       1.27309503      -0.3733564      -1.0960385
AARS1           0.9972642       0.69944418      -0.6523539      -1.0443545
AAAS            0.7106395       0.81561074      -1.3397753      -0.1864749
AACSP1          0.9985590       0.71637534      -0.9554120      -0.7595223
```

DEMA package

AAGAB	1.0769772	0.61814883	-0.9507280	-0.7443980
AACS	0.5086833	1.14412657	-1.0085114	-0.6442985
AASDHPPT	0.8071383	0.91551061	-0.7459808	-0.9766682
AADAC	0.8863152	0.83580877	-0.7325829	-0.9895411
AARSD1	0.8645997	0.83776414	-1.0765674	-0.6257964
AABT	1.1556595	0.47758562	-1.0431379	-0.5901072
AAA4	0.4724367	1.04024800	-1.2810233	-0.2316614
AADACL2	1.3011499	0.02106791	-1.1261009	-0.1961170
AAMP	1.1004181	0.59274116	-0.7999765	-0.8931828
AANAT	1.1659982	0.49625758	-0.7516622	-0.9105936
AASDH	-0.4675273	1.32468488	-1.0027167	0.1455591
AAK1	1.0835176	0.59586852	-0.6653927	-1.0139935
AADACL4	0.5993446	1.07065287	-1.0586534	-0.6113440
AADAT	0.6561996	1.04082847	-0.6966487	-1.0003793
ACAD11	-1.1108356	-0.58004971	0.8409657	0.8499196
VRK1	NA	NA	NA	NA
VSTM4	NA	NA	NA	NA
VPS9D1-AS1	NA	NA	NA	NA
VSNL1	NA	NA	NA	NA
ACAP1	-0.5392450	-1.13710621	0.9127669	0.7635843
VSTM5	NA	NA	NA	NA
VSIG10L2	NA	NA	NA	NA
VSIG10	NA	NA	NA	NA
VSTM1	NA	NA	NA	NA
VRK2	NA	NA	NA	NA
VSTM2L	NA	NA	NA	NA
VSIR	NA	NA	NA	NA
VSTM2B	NA	NA	NA	NA
VSIG4	NA	NA	NA	NA
VRK3	NA	NA	NA	NA
VSIG1	NA	NA	NA	NA
VSIG10L	NA	NA	NA	NA
VSTM2A	NA	NA	NA	NA
VPS9D1	NA	NA	NA	NA
VSX1	NA	NA	NA	NA
	Sample5.Study2	Sample6.Study2	Sample7.Study2	Sample8.Study2
AARS2	1.29003246	-0.1493767	1.1881662	-0.6257615
AADACL3	1.18101154	0.9235015	0.5709817	-0.7336264
AARS1	0.32261014	1.3600903	0.7049359	-1.2920142
AAAS	0.59178756	0.5776332	1.3366176	-1.1031207
AACSP1	0.75769816	0.6575860	1.2379418	-0.6163623
AAGAB	0.66979718	0.5913302	1.3541761	-0.8630518
AACS	0.87262982	1.0046754	0.7826212	-1.3015653
AASDHPPT	1.03567093	0.9364763	0.5862831	-1.4423486
AADAC	0.75691392	0.2637028	1.3332952	-0.1318817
AARSD1	0.88971101	0.2844332	1.3444491	-1.1891260
AABT	0.85037008	-0.2263859	0.7607782	-0.8746852
AAA4	0.05717809	1.4476910	0.9225027	-0.8864413
AADACL2	1.40453080	0.2067201	0.7295036	-0.4744804
AAMP	0.61483783	1.0780356	0.7377432	-1.1325895
AANAT	0.42906396	0.8208962	1.3123105	-0.7528130

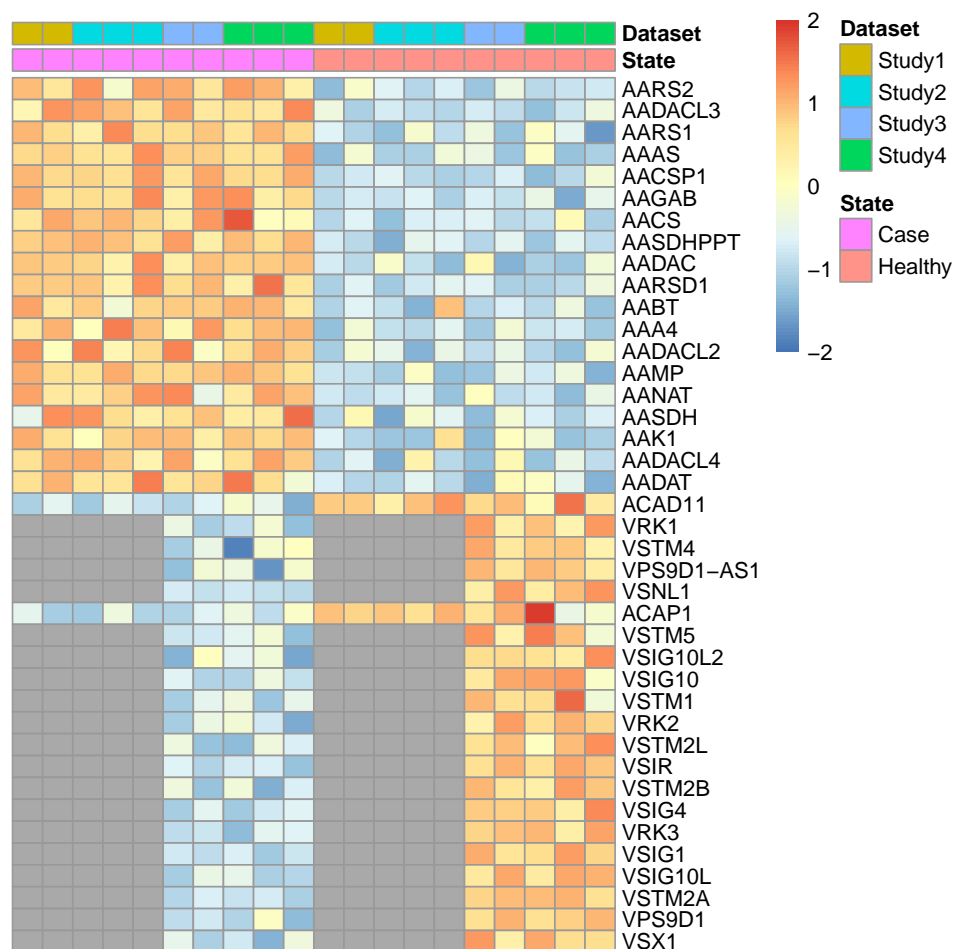
DExMA package

AASDH	1.28122621	0.6882994	0.3313130	-1.5532911
AAK1	0.03726305	0.7816216	0.9843186	-1.2179787
AADACL4	1.09122261	0.8224126	0.2253374	-1.4373346
AADAT	0.55307845	0.5457234	1.4479847	-1.0401099
ACAD11	-1.16714015	-0.5365638	-0.8363435	0.3055333
VRK1	NA	NA	NA	NA
VSTM4	NA	NA	NA	NA
VPS9D1-AS1	NA	NA	NA	NA
VSNL1	NA	NA	NA	NA
ACAP1	-1.17329107	-0.3798016	-1.0565983	0.8999814
VSTM5	NA	NA	NA	NA
VSIG10L2	NA	NA	NA	NA
VSIG10	NA	NA	NA	NA
VSTM1	NA	NA	NA	NA
VRK2	NA	NA	NA	NA
VSTM2L	NA	NA	NA	NA
VSIR	NA	NA	NA	NA
VSTM2B	NA	NA	NA	NA
VSIG4	NA	NA	NA	NA
VRK3	NA	NA	NA	NA
VSIG1	NA	NA	NA	NA
VSIG10L	NA	NA	NA	NA
VSTM2A	NA	NA	NA	NA
VPS9D1	NA	NA	NA	NA
VSX1	NA	NA	NA	NA
	Sample9.Study2	Sample10.Study2	Sample11.Study3	Sample12.Study3
AARS2	-1.02470491	-0.6783556	1.0765064	0.511536842
AADACL3	-0.93417370	-1.0076946	1.1791022	0.472019038
AARS1	-0.17388068	-0.9217414	0.7049468	0.900420939
AAAS	-1.10680832	-0.2961094	0.8055885	0.825278291
AACSP1	-0.96070349	-1.0761601	0.5469671	1.121549627
AAGAB	-0.66653569	-1.0857160	0.3155386	1.272230600
AACS	-0.67777542	-0.6805857	0.3448361	1.245414822
AASDHPPT	-0.51632041	-0.5997614	1.2322820	0.353638199
AADAC	-0.89449227	-1.3275379	0.3168962	0.927391951
AARSD1	-0.77010577	-0.5593615	0.6718124	1.010801637
AABT	-1.42799917	0.9179220	0.8656974	0.849364871
AAA4	-0.96655055	-0.5743799	0.1665761	1.238929034
AADACL2	-1.42596732	-0.4403068	1.3999718	-0.037794528
AAMP	-0.02038311	-1.2776440	0.7285864	0.893817981
AANAT	-0.57603553	-1.2334222	1.3902037	-0.460590455
AASDH	-0.17029375	-0.5772538	0.6129201	0.910089484
AAK1	-1.23178212	0.6465576	0.9894857	0.371213632
AADACL4	0.26967066	-0.9713088	1.1607444	-0.034476495
AADAT	-0.54991192	-0.9567647	0.5509839	0.777842738
ACAD11	0.94328094	1.2912332	-1.0427073	-0.659717646
VRK1	NA	NA	-0.3991328	-1.132808807
VSTM4	NA	NA	-1.1480953	-0.443804656
VPS9D1-AS1	NA	NA	-1.2947008	-0.222935977
VSNL1	NA	NA	-0.7343571	-0.877211096
ACAP1	0.64129122	1.0684184	-1.0551468	-0.606268006

VSTM5	NA	NA	-0.8247090	-0.759672559
VSIG10L2	NA	NA	-1.4155409	0.004719214
VSIG10	NA	NA	-0.5943645	-1.046173579
VSTM1	NA	NA	-1.1241648	-0.538460201
VRK2	NA	NA	-1.1208364	-0.395656835
VSTM2L	NA	NA	-0.3575971	-1.237439226
VSIR	NA	NA	-0.6372153	-1.047895072
VSTM2B	NA	NA	-0.3130491	-1.253243977
VSIG4	NA	NA	-1.1285994	-0.555042433
VRK3	NA	NA	-0.9173044	-0.808749557
VSIG1	NA	NA	-0.7653212	-0.923883092
VSIG10L	NA	NA	-1.1438809	-0.453636753
VSTM2A	NA	NA	-1.0084318	-0.704574832
VPS9D1	NA	NA	-0.9493557	-0.755147481
VSX1	NA	NA	-0.4791305	-1.072203520
	Sample13.Study3	Sample14.Study3	Sample15.Study4	Sample16.Study4
AARS2	-1.193097649	-0.39494560	1.2047919	1.07007824
AADACL3	-0.716438370	-0.93468287	0.6082930	0.47422102
AARS1	-0.353412568	-1.25195516	0.5683881	0.99032956
AAAS	-0.403074596	-1.22779222	0.5868315	0.60634862
AACSP1	-0.993913293	-0.67460343	0.7438120	0.70095867
AAGAB	-0.690542772	-0.89722647	1.3253285	0.34768914
AACS	-0.614044000	-0.97620692	1.7246362	0.01073741
AASDHPPT	-1.015277040	-0.57064316	0.9699261	0.67584429
AADAC	0.173278628	-1.41756675	0.8166646	0.83547046
AARSD1	-0.605348342	-1.07726574	0.3208675	1.54083274
AABT	-1.028452101	-0.68661020	1.0582122	1.01834320
AAA4	-1.176875375	-0.22862980	0.6756384	0.98541331
AADACL2	-0.916553115	-0.44562416	0.6482716	1.08819763
AAMP	-1.232006994	-0.39039735	1.0541916	0.87742615
AANAT	-0.004861139	-0.92475211	0.4283357	1.18895237
AASDH	-1.325546900	-0.19746268	0.3936066	0.46554875
AAK1	-1.372219271	0.01151991	0.8914460	0.71351002
AADACL4	-1.276311403	0.15004352	0.6127288	1.15154039
AADAT	-1.441744415	0.11291774	1.4996941	0.68539206
ACAD11	0.732030611	0.97039437	-0.1564368	-0.49161788
VRK1	1.204157191	0.32778445	-0.9103028	-0.22005511
VSTM4	1.125981608	0.46591839	-1.8484634	-0.17431453
VPS9D1-AS1	1.006243659	0.51139316	-0.3766663	-1.71358002
VSNL1	0.362701612	1.24886662	-0.7663241	-0.88793936
ACAP1	0.567121587	1.09429317	-0.3501057	-0.91685493
VSTM5	1.285962031	0.29841958	-0.5761074	-0.25540357
VSIG10L2	0.682909288	0.72791243	-0.5669168	-0.27272316
VSIG10	0.498889938	1.14164815	-1.0549086	-0.36552545
VSTM1	1.010411751	0.65221327	-0.3253520	-1.22683282
VRK2	0.290643408	1.22584984	-0.2475666	-0.75203655
VSTM2L	0.612048448	0.98298788	-1.3242282	-0.30928837
VSIR	0.647524480	1.03758593	-0.7307610	-0.66790259
VSTM2B	1.011994595	0.55429844	-0.2874520	-1.45039745
VSIG4	0.862633571	0.82100827	-1.1848497	-0.76428048
VRK3	0.776871260	0.94918271	-1.3310580	-0.56329900

DExMA package

VSIG1	1.103462931	0.58574140	-0.6755072	-1.18598708
VSIG10L	0.474938718	1.12257892	-0.4928980	-1.07710114
VSTM2A	0.757903621	0.95510300	-0.8669201	-0.72768843
VPS9D1	0.657559123	1.04694410	-1.0661534	-0.04791842
VSX1	1.232555953	0.31877803	-0.7673579	-1.42403271
	Sample17.Study4	Sample18.Study4	Sample19.Study4	Sample20.Study4
AARS2	0.33548921	-0.978168251	-0.8478691	-0.78432197
AADACL3	1.38821537	-1.296333970	-0.8194346	-0.35496081
AARS1	0.74282169	-0.047906853	-0.5778748	-1.67575772
AAAS	1.21907318	-0.053486390	-1.2717939	-1.08697301
AACSP1	1.08356763	-1.350536645	-0.9520291	-0.22577249
AAGAB	0.71612384	-0.436915497	-1.4827170	-0.46950899
AACS	0.12387329	-0.900151433	0.1287143	-1.08780978
AASDHPPT	1.01563000	-1.194586455	-0.5487922	-0.91802170
AADAC	0.93597171	-1.091781144	-1.2211976	-0.27512806
AARSD1	0.52121887	-1.087312823	-0.9866706	-0.30893567
AABT	0.48724030	-0.977729098	-0.3503127	-1.23575392
AAA4	1.02608096	-0.790252699	-0.7292882	-1.16759179
AADACL2	0.79068870	-1.024505524	-1.2952874	-0.20736496
AAMP	0.61802876	-0.782743132	-0.3527996	-1.41410376
AANAT	0.92838867	-0.766724415	-1.3200648	-0.45888756
AASDH	1.58695523	-0.670733809	-1.0926837	-0.68269299
AAK1	0.96135491	-0.243104099	-1.2461008	-1.07710606
AADACL4	0.84468751	-1.244625253	-0.4527872	-0.91154424
AADAT	-0.24171822	-0.035805706	-0.5054236	-1.40213857
ACAD11	-1.46496299	0.124966311	1.5398884	0.44816300
VRK1	-1.27953065	0.944625717	0.2227733	1.24248951
VSTM4	0.01327709	0.837086850	0.8731125	0.29930145
VPS9D1-AS1	-0.17021280	1.012295278	0.8351452	0.41301863
VSNL1	-0.96501288	0.388287090	0.9517297	1.27925953
ACAP1	-0.06453924	1.948688579	-0.4582433	-0.15894545
VSTM5	-1.28372603	1.437254202	0.9131807	-0.23519795
VSIG10L2	-1.51565527	0.608446024	0.4200656	1.32678363
VSIG10	-0.90635013	1.163428345	1.2531779	-0.08982205
VSTM1	-0.47355971	0.701930653	1.6094881	-0.28567422
VRK2	-1.48970178	0.651594331	1.0616402	0.77607049
VSTM2L	-0.66769822	0.004892528	0.9632726	1.33304965
VSIR	-1.25441145	0.664738911	1.1119140	0.87642214
VSTM2B	-0.68584179	0.331013623	1.1948646	0.89781307
VSIG4	-0.59129239	0.830268073	0.3497454	1.36040909
VRK3	-0.64249280	0.994620857	0.3713171	1.17091190
VSIG1	-0.79024789	0.686231980	1.2173862	0.74812392
VSIG10L	-1.02829017	0.429639636	1.1256359	1.04301371
VSTM2A	-1.09845817	0.971509565	1.0688633	0.65269384
VPS9D1	-1.34019192	0.658610706	0.8004777	0.99517526
VSX1	-0.33785216	1.144663968	0.6788051	0.70577373



4 Additional information

DExMA provides some functions which may be useful for the user, although they are not essential to perform meta-analysis.

4.1 GEO microarray data download

In addition to using own user data, **DExMA** package allows to make use of public microarray data from the NCBI GEO public database [3]. For doing that, we can make use of `downloadGEOData()` function. This function uses internally `GEOquery` package in order to download some files at the same time. This function has an input a character vector (`GEOobject`) with the GEO ID of the different datasets that we want to download and a character (`directory`) that indicates the directory where GSE Series Matrix files [9] are going to be stored.

```
> GEOobjects<- c("GSE4588", "GSE10325")
> dataGEO<-downloadGEOData(GEOobjects)
```

Once the download process is completed, we get a list of ExpressionSets. This list can be used as input of `createObjectMA()` function, although it is advisable to homogenize gene annotation, in case genes IDs are not Entrez, Official Gene Symbol or Ensembl.

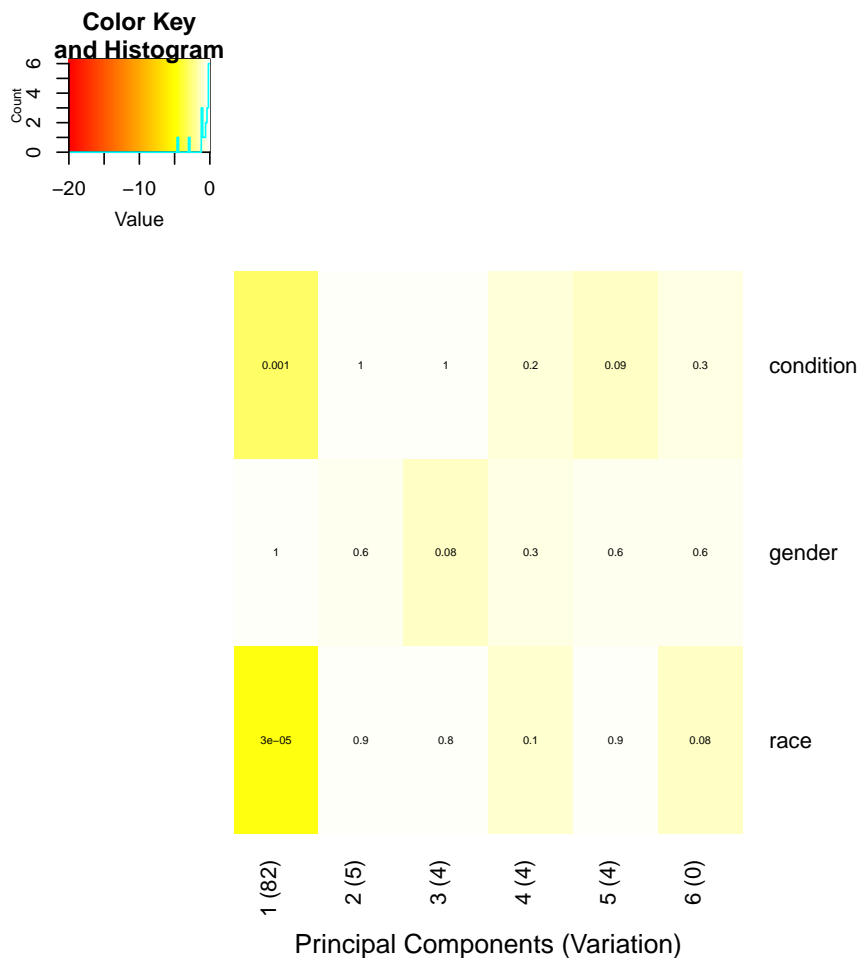
4.2 Using RNA-Seq data

DExMA internally uses [limma](#) package in order to assess differential expression. Therefore, RNA-Seq data must be previously normalized by the user in order to be able to include correctly these data in the gene-expression meta-analysis. Since [limma](#) is used internally, we recommend to apply the steps described in the limma user's guide for the RNA-Seq data normalization [10], although the users can use the type of normalization they prefer.

4.3 Removing Batch Effects

Before the creation of the `objectMA`, a batch effect correction can be applied in order to reduce the effect of covariates that may be affecting to gene expression [1]. Firstly, with functions `prince()` and `prince.plot()` function of the [swamp](#) package [11], we can obtain a visualization of the p-values of each principal component associated with the categorical covariates

```
> library(swamp)
> pheno <- listPhenodatas$Study2
> pheno <- pheno[,apply(pheno,2,function(x) length(table(x)))>1]
> # Character variables must be converted in numeric
> pheno <- data.frame(apply(pheno, 2, factor), stringsAsFactors = TRUE)
> res_prince <- prince(listMatrixEX$Study2, pheno,top=ncol(listMatrixEX$Study2))
> prince.plot(res_prince,note=TRUE , notecex=0.5)
```

The categorical variables that may be causing a batch effect can be corrected by using the `removeBatch()` function. The input of this function is the expression matrix and the phenodata. In addition, we also have to add a formula with the variables for which we want to correct the gene expression and the name of the variable that contains the cases and controls groups. Finally, if there is a covariate inside the formula that we want to give greater importance, we will have the option to indicate in the function (`mainCov()`). Here we show an example in which we have corrected the gene expression of the previous study by two of their covariates:

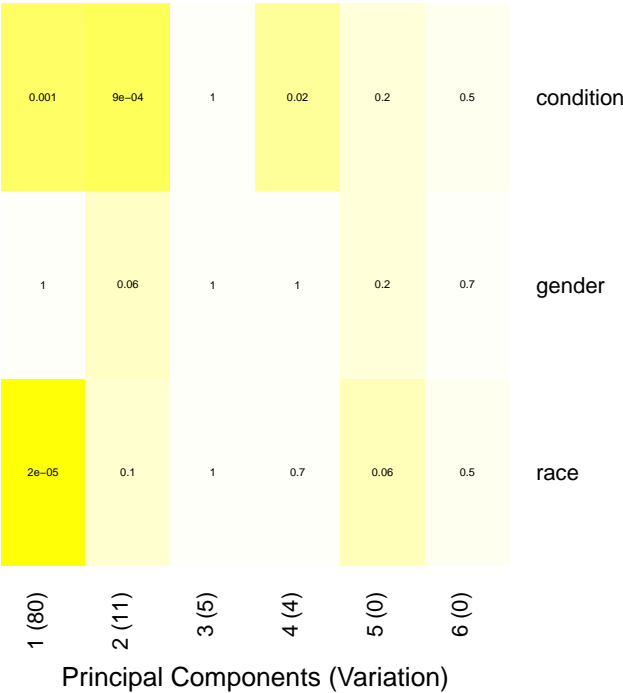
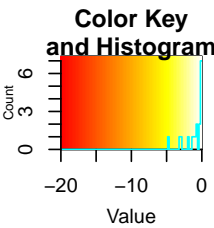
```
> listMatrixEX$Study2 <- batchRemove(listMatrixEX$Study2, listPhenodatas$Study2,
+                                     formula=~gender+race,
+                                     mainCov = "race", nameGroup="condition")
```

Coefficients not estimable: batch1 batch2 (Intercept) raceC raceH

```
> head(listMatrixEX$Study2)
```

	Sample5	Sample6	Sample7	Sample8	Sample9	Sample10
100859927	4.582088	6.862649	5.358102	3.034471	2.903085	2.903085
8086	6.367896	6.345326	6.262866	3.665331	3.659451	3.659451
8212	6.167717	4.965395	3.448186	4.241494	3.623655	3.623655
65985	5.684300	5.898841	5.542624	2.151772	3.165276	3.165276
729522	5.664208	5.485121	6.729837	3.206196	2.590215	2.590215

13	5.943241	5.225990	7.411197	4.650713	3.541689	3.541689
----	----------	----------	----------	----------	----------	----------



4.4 Calculating Effects size

The `calculateES()` function returns the effects size in each of the studies. Moreover, it calculates the variance of each of the effects and the \log_2 Fold Change. The effects size are calculated by making use of the *Hedges' g estimator* [1].

```
> effects <- calculateES(newObjectMA)
> head(effects$ES)

      Study1  Study2  Study3  Study4
AAA4  1.448942 2.5008385 1.1221262 6.642233
AAAS  1.506204 2.9643045 2.2593392 2.430575
AABT  2.288894 0.7656633 5.7273196 3.478798
AACS  2.579019 5.3315818 1.8723418 1.207949
AACSP1 5.705553 5.1083967 2.9008711 3.138604
AADAC 7.515582 2.1970319 0.8345501 3.776154

> head(effects$Var)
```

```

      Study1   Study2   Study3   Study4
AAA4  0.5889596 0.9478494 0.4839265 4.1032711
AAAS  0.6101119 1.1589251 0.9646073 0.9189745
AABT  0.9814100 0.4755200 4.4268043 1.4351698
AACS  1.1579482 2.7954803 0.7647386 0.5482618
AACSP1 4.3956973 2.6013097 1.3784123 1.2475697
AADAC 7.3870275 0.8289124 0.4135899 1.6149449

```

```
> head(effects$logFC)
```

```

      Study1   Study2   Study3   Study4
AAA4  2.167998 2.654066 3.118286 3.385361
AAAS  2.931649 2.663952 4.321073 2.807056
AABT  3.857649 1.030831 3.259410 3.594168
AACS  3.376387 2.881147 4.252159 1.733698
AACSP1 2.157823 3.164180 3.998437 2.492064
AADAC 6.771502 2.282112 1.747278 3.715857

```

4.5 Calculating Individual P-values

Similar to the calculation of effects sizes, the individual p-values of each of the studies and the \log_2 fold change of each one can also be calculated by applying *pvalueIndAnalysis()*. P-value are obtained by assessing differential expression with *limma* package.

```

> pvalues <- pvalueIndAnalysis(newObjectMA)
> head(pvalues$p)

```

```

      Study1   Study2   Study3   Study4
AAA4  0.084586213 0.0124328637 0.137545477 0.0002392007
AAAS  0.078241874 0.0065691298 0.032924671 0.0139623839
AABT  0.031889060 0.2765998575 0.003696266 0.0035646909
AACS  0.024335148 0.0005986238 0.049748561 0.1156145550
AACSP1 0.003718102 0.0007176307 0.018622696 0.0053471186
AADAC 0.001900842 0.0197199607 0.223678418 0.0025641765

```

```
> head(pvalues$logFC)
```

```

      Study1   Study2   Study3   Study4
AAA4  2.167998 2.654066 3.118286 3.385361
AAAS  2.931649 2.663952 4.321073 2.807056
AABT  3.857649 1.030831 3.259410 3.594168
AACS  3.376387 2.881147 4.252159 1.733698
AACSP1 2.157823 3.164180 3.998437 2.492064
AADAC 6.771502 2.282112 1.747278 3.715857

```

5 Session info

```

R version 4.6.0 Patched (2026-04-24 r89963)
Platform: aarch64-apple-darwin23
Running under: macOS Tahoe 26.3.1

```

DExMA package

```
Matrix products: default
BLAS:   /Library/Frameworks/R.framework/Versions/4.6/Resources/lib/libRblas.0.dylib
LAPACK: /Library/Frameworks/R.framework/Versions/4.6/Resources/lib/libRlapack.dylib; LAPACK version 3.

locale:
[1] C/en_US.UTF-8/en_US.UTF-8/C/en_US.UTF-8/en_US.UTF-8

time zone: America/New_York
tzcode source: internal

attached base packages:
[1] stats      graphics  grDevices  utils      datasets  methods    base

other attached packages:
[1] swamp_1.5.1      MASS_7.3-65      gplots_3.3.0
[4] amap_0.8-20      impute_1.86.0    Biobase_2.72.0
[7] BiocGenerics_0.58.0 generics_0.1.4    DExMA_1.20.0
[10] DExMAdata_1.19.0

loaded via a namespace (and not attached):
[1] tidyselect_1.2.1      dplyr_1.2.1
[3] farver_2.1.2          blob_1.3.0
[5] Biostrings_2.80.0     bitops_1.0-9
[7] fastmap_1.2.0         GEOquery_2.80.0
[9] XML_3.99-0.23         digest_0.6.39
[11] lifecycle_1.0.5       bnstruct_1.0.15
[13] survival_3.8-6        statmod_1.5.1
[15] KEGGREST_1.52.0       RSQLite_2.4.6
[17] magrittr_2.0.5        genefilter_1.94.0
[19] compiler_4.6.0        rlang_1.2.0
[21] tools_4.6.0           igraph_2.3.0
[23] yaml_2.3.12           data.table_1.18.2.1
[25] knitr_1.51            S4Arrays_1.12.0
[27] bit_4.6.0             DelayedArray_0.38.0
[29] plyr_1.8.9            xml2_1.5.2
[31] RColorBrewer_1.1-3    abind_1.4-8
[33] BiocParallel_1.46.0   KernSmooth_2.23-26
[35] purrr_1.2.2           grid_4.6.0
[37] stats4_4.6.0          caTools_1.18.3
[39] xtable_1.8-8          edgeR_4.10.0
[41] scales_1.4.0          gtools_3.9.5
[43] dichromat_2.0-0.1     SummarizedExperiment_1.42.0
[45] cli_3.6.6             rmarkdown_2.31
[47] crayon_1.5.3          otel_0.2.0
[49] tzdb_0.5.0            httr_1.4.8
[51] DBI_1.3.0             cachem_1.1.0
[53] splines_4.6.0         parallel_4.6.0
[55] AnnotationDbi_1.74.0  BiocManager_1.30.27
[57] XVector_0.52.0        matrixStats_1.5.0
[59] vctrs_0.7.3           Matrix_1.7-5
[61] jsonlite_2.0.0        sva_3.60.0
```

[63] hms_1.1.4	IRanges_2.46.0
[65] S4Vectors_0.50.0	bit64_4.8.0
[67] locfit_1.5-9.12	limma_3.68.0
[69] tidyr_1.3.2	snpStats_1.62.0
[71] annotate_1.90.0	glue_1.8.1
[73] codetools_0.2-20	gtable_0.3.6
[75] GenomicRanges_1.64.0	tibble_3.3.1
[77] pillar_1.11.1	htmltools_0.5.9
[79] Seqinfo_1.2.0	R6_2.6.1
[81] evaluate_1.0.5	lattice_0.22-9
[83] readr_2.2.0	rentrez_1.2.4
[85] png_0.1-9	pheatmap_1.0.13
[87] memoise_2.0.1	BiocStyle_2.40.0
[89] Rcpp_1.1.1-1.1	SparseArray_1.12.0
[91] nlme_3.1-169	mgcv_1.9-4
[93] xfun_0.57	MatrixGenerics_1.24.0
[95] pkgconfig_2.0.3	

References

- [1] Toro-Domínguez D., Villatoro-García J.A., Martorell-Marugán J., and et al. A survey of gene expression meta-analysis: methods and applications. *Briefings in Bioinformatics*, pages 1–12, 2020. doi:<https://doi.org/10.1093/bib/bbaa019>.
- [2] Christopher A. Mancuso, Jacob L. Canfield, Deepak Singla, and Arjun Krishnan. A flexible, interpretable, and accurate approach for imputing the expression of unmeasured genes. *Nucleic Acids Research*, 48(21):e125–e125, 2020. URL: <https://doi.org/10.1093/nar/gkaa881>.
- [3] Barret T., Wilhite S., Ledoux P., and et al. Ncbi geo: archive for functional genomics data sets–update. *Nucleic Acids Research*, pages 991–995, 2020. doi:<https://doi.org/10.1093/nar/gks1193>.
- [4] Higgins J. and Thompson S. Quantifying heterogeneity in a meta-analysis. *Statistics in Medicine*, pages 1539–1558, 2002. doi:[10.1002/sim.1186](https://doi.org/10.1002/sim.1186).
- [5] Higgins J., Thompson S., Deeks J., and Altman D. Measuring inconsistency in meta-analyses. *BMJ*, pages 557–560, 2003. doi:[10.1136/bmj.327.7414.557](https://doi.org/10.1136/bmj.327.7414.557).
- [6] Hastie T., Tibshirani R., Narasimhan B., and Chu G. impute: Imputation for microarray data. 2019.
- [7] Lazar C., Meganck S., Taminau J., and et al. Batch effect removal methods for microarray gene expression data integration: a survey. *Briefings in Bioinformatics*, pages 469–490, 2013. doi:[10.1093/bib/bbs037](https://doi.org/10.1093/bib/bbs037).
- [8] Wickham H. and Siedel D. Scale functions for visualization. 2020.
- [9] Davis S. and Meltzer P.S. Geoquery: a bridge between the gene expression omnibus (geo) and bioconductor. *Bioinformatics*, pages 1846–1847, 2007. doi:[10.1093/bioinformatics/btm254](https://doi.org/10.1093/bioinformatics/btm254).
- [10] Smyth G. K., Ritchie M., Thorne N., Yifang Hu, and et al. Linear models for microarray and rna-seq data user's guide. 2020.

- [11] Lauss M. Visualization, analysis and adjustment of high-dimensional data in respect to sample annotations. 2019.