

# Package ‘OncoScore’

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**Depends** R (>= 4.1.0),

**Imports** biomaRt, grDevices, graphics, utils, methods,

**Suggests** BiocGenerics, BiocStyle, knitr, testthat,

**Description** OncoScore is a tool to measure the association of genes to cancer based on citation frequencies in biomedical literature. The score is evaluated from PubMed literature by dynamically updatable web queries.

**Encoding** UTF-8

**License** file LICENSE

**URL** <https://github.com/danro9685/OncoScore>

**BugReports** <https://github.com/danro9685/OncoScore>

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---

combine.query.results *combine.query.results*

---

### Description

Merge a set of genes in a unique one in order to account for possible aliases

### Usage

```
combine.query.results(query, genes, new.name)
```

### Arguments

|          |  |
|----------|--|
| query    | The result of perform.query, perform.query.timeseries or perform.query.from.region |
| genes    | A list of genes to be merged   |
| new.name | A string containing the new name to be used for the new genes                      |

### Value

The frequencies of the genes in the cancer related documents and in all the documents retrieved on PubMed

### Examples

```
data(query)
combine.query.results(query, c('IDH1', 'IDH2'), 'new_gene')
```

---

`combine.single.matrix` *combine.single.matrix*

---

### **Description**

Perform merge procedure on a matrix

### **Usage**

```
combine.single.matrix(query, genes, new.name)
```

### **Arguments**

|                       |  |
|-----------------------|--|
| <code>query</code>    | The result of <code>perform.query</code> , <code>perform.query.timeseries</code> or <code>perform.query.from.region</code> |
| <code>genes</code>    | A list of genes to be merged   |
| <code>new.name</code> | A string containing the new name to be used for the new genes  |

### **Value**

a merged matrix

---

`compute.frequencies.scores`  
*compute.frequencies.scores*

---

### **Description**

compute the logarithmic scores based on the frequencies of the genes

### **Usage**

```
compute.frequencies.scores(data, filter.threshold = 1, analysis.mode = "Log2")
```

### **Arguments**

|                               |   |
|-------------------------------|---|
| <code>data</code>             | input data as result of the function <code>perform.query</code>           |
| <code>filter.threshold</code> | threshold to filter for a minimum number of citations for the genes       |
| <code>analysis.mode</code>    | logarithmic scores to be computed, i.e., log10, log2, natural log or log5 |

### **Value**

the computed scores

---

`compute.oncoscore`      *compute.oncoscore*

---

### Description

compute the OncoScore for a list of genes

### Usage

```
compute.oncoscore(
  data,
  filter.threshold = 0,
  analysis.mode = "Log2",
  cutoff.threshold = 21.09,
  file = NULL,
  filter.invalid = TRUE
)
```

### Arguments

`data`                    input data as result of the function `perform.query`

`filter.threshold`                    threshold to filter for a minimum number of citations for the genes

`analysis.mode`    logarithmic scores to be computed, i.e., log10, log2, natural log or log5

`cutoff.threshold`                    threshold to be used to asses the oncogenes

`file`                    should I save the results to text files?

`filter.invalid`    auto-remove genes with invalid count

### Value

the computed OncoScores and the clusters for the genes

### Examples

```
data(query)
compute.oncoscore(query)
```

---

`compute.oncoscore.from.region`  
*compute.oncoscore.from.region*

---

### Description

Perform OncoScore analysis on a given chromosomal region

**Usage**

```
compute.oncoscore.from.region(  
  chromosome,  
  start = NA,  
  end = NA,  
  gene.num.limit = 100,  
  filter.threshold = NA,  
  analysis.mode = "Log2",  
  cutoff.threshold = 21.09,  
  file = NULL  
)
```

**Arguments**

|                  |  |
|------------------|--|
| chromosome       | chromosome to be retrieved   |
| start            | initial position to be used  |
| end              | final position to be used  |
| gene.num.limit   | A limit to the genes to be considered in the analysis; this is done to limit the number of queries to PubMed |
| filter.threshold | threshold to filter for a minimum number of citations for the genes  |
| analysis.mode    | logarithmic scores to be computed, i.e., log10, log2, natural log or log5                                    |
| cutoff.threshold | threshold to be used to assess the oncogenes   |
| file             | should I save the results to text files?   |

**Value**

the computed scores

**Examples**

```
chromosome = 15  
start = 200000  
end = 300000
```

---

```
compute.oncoscore.timeseries  
compute.oncoscore.timeseries
```

---

**Description**

perform the OncoScore time series analysis for a list of genes and data times

**Usage**

```
compute.oncoscore.timeseries(
  data,
  filter.threshold = 0,
  analysis.mode = "Log2",
  cutoff.threshold = 21.09,
  file = NULL
)
```

**Arguments**

`data` input data as result of the function `perform.query.timeseries`

`filter.threshold` threshold to filter for a minimum number of citations for the genes

`analysis.mode` logarithmic scores to be computed, i.e., log10, log2, natural log or log5

`cutoff.threshold` threshold to be used to asses the oncogenes

`file` should I save the results to text files?

**Value**

the performed OncoScores time series analysis

**Examples**

```
data(query.timepoints)
compute.oncoscore.timeseries(query.timepoints)
```

---

|                                 |                           |
|---------------------------------|---------------------------|
| <code>estimate.oncogenes</code> | <i>estimate.oncogenes</i> |
|---------------------------------|---------------------------|

---

**Description**

estimate the oncoscore for the genes

**Usage**

```
estimate.oncogenes(data, cutoff.threshold = 21.09)
```

**Arguments**

`data` input data as result of the function `compute.frequencies.scores`

`cutoff.threshold` threshold to be used to asses the oncogenes

**Value**

the computed scores and oncogenes

---

genes

*A list of genes*

---

**Description**

This dataset contains a list of genes to be used in the analysis as an example

**Usage**

```
data(genes)
```

**Format**

rdata

**Value**

list of 5 elements

**Source**

example data

---

```
get.genes.from.biomart
```

*get.genes.from.biomart*

---

**Description**

Get a gene list from biomart

**Usage**

```
get.genes.from.biomart(chromosome, start = NA, end = NA)
```

**Arguments**

|            |                             |
|------------|-----------------------------|
| chromosome | chromosome to be retrieved  |
| start      | initial position to be used |
| end        | final position to be used   |

**Value**

A list of genes

**Examples**

```
chromosome = 15  
start = 200000  
end = 300000
```

---

`get.list.from.xml`      *get.list.from.xml*

---

**Description**

process the result of the query

**Usage**

`get.list.from.xml(webget)`

**Arguments**

`webget`              The result from the query to PubMed

**Value**

Processed result obtained from the query to PubMed

---

`get.pubmed.driver.analysis`  
*get.pubmed.driver.analysis*

---

**Description**

query PubMed for a list of genes

**Usage**

`get.pubmed.driver.analysis(keywords, gene)`

**Arguments**

`keywords`              The set of keywords to be used for the query to PubMed

`gene`                    The name of a gene to be used for the query to PubMed

**Value**

The frequency for the current gene retrieved with the query on the provided set of keywords

---

`perform.query`      *perform.query*

---

### **Description**

perform the query to PubMed

### **Usage**

```
perform.query(list.of.genes, gene.num.limit = 100, custom.search = NA)
```

### **Arguments**

- `list.of.genes`    The list of genes to be used in the queries to PubMed
- `gene.num.limit`    A limit to the genes to be considered in the analysis; this is done to limit the number of queries to PubMed
- `custom.search`    A custom set of keywords to be used when querying PubMed

### **Value**

The frequencies of the genes in the cancer related documents and in all the documents retrieved on PubMed

### **Examples**

```
data(genes)
```

---

`perform.query.from.region`  
*perform.query.from.region*

---

### **Description**

Perform the query to PubMed on a given chromosomal region

### **Usage**

```
perform.query.from.region(  
  chromosome,  
  start = NA,  
  end = NA,  
  gene.num.limit = 100  
)
```

**Arguments**

|                |  |
|----------------|--|
| chromosome     | chromosome to be retrieved   |
| start          | initial position to be used  |
| end            | final position to be used  |
| gene.num.limit | A limit to the genes to be considered in the analysis; this is done to limit the number of queries to PubMed |

**Value**

The frequencies of the genes in the cancer related documents and in all the documents retrieved on PubMed

**Examples**

```
chromosome = 15
start = 200000
end = 300000
```

---

```
perform.query.timeseries
```

```
perform.query.timeseries
```

---

**Description**

perform the query to PubMed for the time series analysis

**Usage**

```
perform.query.timeseries(
  list.of.genes,
  list.of.datatimes,
  gene.num.limit = 100,
  timepoints.limit = 10,
  custom.search = NA
)
```

**Arguments**

|                   |  |
|-------------------|--|
| list.of.genes     | The list of genes to be used in the queries to PubMed  |
| list.of.datatimes | The list of time points to be used in the queries to PubMed  |
| gene.num.limit    | A limit to the genes to be considered in the analysis; this is done to limit the number of queries to PubMed       |
| timepoints.limit  | A limit to the time points to be considered in the analysis; this is done to limit the number of queries to PubMed |
| custom.search     | A custom set of keywords to be used when querying PubMed   |

**Value**

The frequencies of the genes in the cancer related documents and in all the documents retrieved on PubMed at the specified time points

**Examples**

```
data(genes)
data(timepoints)
```

---

|                |                       |
|----------------|-----------------------|
| plot.oncoscore | <i>plot.oncoscore</i> |
|----------------|-----------------------|

---

**Description**

plot the OncoScore for a list of genes

**Usage**

```
## S3 method for class 'oncoscore'
plot(
  x,
  gene.number = 5,
  main = "OncoScore",
  xlab = "score",
  ylab = "genes",
  file = NA,
  ...
)
```

**Arguments**

|             |  |
|-------------|--|
| x           | input data as result of the function compute.OncoScore |
| gene.number | number of genes to print                               |
| main        | the title  |
| xlab        | description of x asix (default score)                  |
| ylab        | description of y asix (default genes)                  |
| file        | where to save the plot                                 |
| ...         | additional parameter to pass to the barplot function   |

**Value**

A plot

**Examples**

```
data(query)
result = compute.oncoscore(query)
plot.oncoscore(result)
```

---

```
plot.oncoscore.timeseries
      plot.oncoscore.timeseries
```

---

### Description

plot the OncoScore for a list of genes

### Usage

```
## S3 method for class 'oncoscore.timeseries'
plot(
  x,
  gene.number = 5,
  incremental = FALSE,
  relative = FALSE,
  main = "OncoScore",
  xlab = "timepoints",
  ylab = "score",
  legend.pos = "top",
  file = NA,
  ...
)
```

### Arguments

|             |  |
|-------------|--|
| x           | input data as result of the function compute.OncoScore |
| gene.number | number of genes to print                               |
| incremental | display the OncoScore increment                        |
| relative    | display the incrementa as relative value               |
| main        | the title  |
| xlab        | description of x asix (default score)                  |
| ylab        | description of y asix (default genes)                  |
| legend.pos  | Position of the legend                                 |
| file        | where to save the plot                                 |
| ...         | additional parameter to pass to the lines function     |

### Value

A plot

### Examples

```
data(query.timepoints)
result = compute.oncoscore.timeseries(query.timepoints)
plot.oncoscore.timeseries(result)
```

---

query

*The result of perform.web.query on genes*

---

**Description**

This dataset contains the result of perform.web.query on genes

**Usage**

```
data(query)
```

**Format**

rdata

**Value**

matrix 5 x 2

**Source**

example data

---

query.timepoints

*The result of perform.time.series.query on genes and timepoints*

---

**Description**

This dataset contains the result of perform.time.series.query on genes and timepoints

**Usage**

```
data(query.timepoints)
```

**Format**

rdata

**Value**

list of 5 matrix 5 x 2

**Source**

example data

---

|            |                             |
|------------|-----------------------------|
| timepoints | <i>A list of timepoints</i> |
|------------|-----------------------------|

---

**Description**

This dataset contains a list of time points to be used in the analysis as an example

**Usage**

```
data(timepoints)
```

**Format**

rdata

**Value**

list of 5 elements

**Source**

example data

---

|          |                 |
|----------|-----------------|
| try.scan | <i>try.scan</i> |
|----------|-----------------|

---

**Description**

try to query the given URL

**Usage**

```
try.scan(getURL)
```

**Arguments**

getURL            The given URL

**Value**

Result obtained from PubMed

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