

# Package ‘cogeqc’

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**Title** Systematic quality checks on comparative genomics analyses

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**Description** cogeqc aims to facilitate systematic quality checks on standard comparative genomics analyses to help researchers detect issues and select the most suitable parameters for each data set. cogeqc can be used to assess: i. genome assembly and annotation quality with BUSCOs and comparisons of statistics with publicly available genomes on the NCBI; ii. orthogroup inference using a protein domain-based approach and; iii. synteny detection using synteny network properties. There are also data visualization functions to explore QC summary statistics.

**License** GPL-3

**URL** <https://github.com/almeidasilvaf/cogeqc>

**BugReports** <https://support.bioconductor.org/t/cogeqc>

**biocViews** Software, GenomeAssembly, ComparativeGenomics, FunctionalGenomics, Phylogenetics, QualityControl, Network

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**RoxygenNote** 7.2.3

**SystemRequirements** BUSCO ( $\geq 5.1.3$ ) [<https://busco.ezlab.org/>](https://busco.ezlab.org/)

**Imports** utils, graphics, stats, methods, reshape2, ggplot2, scales, ggtree, patchwork, igraph, rlang, ggbeeswarm, jsonlite, Biostrings

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**Author** Fabrício Almeida-Silva [aut, cre] (ORCID:

<<https://orcid.org/0000-0002-5314-2964>>),

Yves Van de Peer [aut] (ORCID: <<https://orcid.org/0000-0003-4327-3730>>)

**Maintainer** Fabrício Almeida-Silva <[fabricao\\_almeidasilva@hotmail.com](mailto:fabricao_almeidasilva@hotmail.com)>

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`cogeqc-package`*cogeqc: Systematic quality checks on comparative genomics analyses*

---

## Description

cogeqc aims to facilitate systematic quality checks on standard comparative genomics analyses to help researchers detect issues and select the most suitable parameters for each data set. cogeqc can be used to asses: i. genome assembly and annotation quality with BUSCOs and comparisons of statistics with publicly available genomes on the NCBI; ii. orthogroup inference using a protein domain-based approach and; iii. synteny detection using synteny network properties. There are also data visualization functions to explore QC summary statistics.

## Author(s)

**Maintainer:** Fabrício Almeida-Silva <fabricio\_almeidasilva@hotmail.com> ([ORCID](#))

Authors:

- Yves Van de Peer <yves.vandeppeer@psb.vib-ugent.be> ([ORCID](#))

## See Also

Useful links:

- <https://github.com/almeidasilvaf/cogeqc>
- Report bugs at <https://support.bioconductor.org/t/cogeqc>

---

`assess_orthogroups`*Assess orthogroup inference based on functional annotation*

---

## Description

Assess orthogroup inference based on functional annotation

## Usage

```
assess_orthogroups(  
  orthogroups = NULL,  
  annotation = NULL,  
  correct_overclustering = TRUE  
)
```

**Arguments**

|                        |   |
|------------------------|---|
| orthogroups            | A 3-column data frame with columns <b>Orthogroup</b> , <b>Species</b> , and <b>Gene</b> . This data frame can be created from the 'Orthogroups.tsv' file generated by OrthoFinder with the function <code>read_orthogroups()</code> .   |
| annotation             | A list of 2-column data frames with columns <b>Gene</b> (gene ID) and <b>Annotation</b> (annotation ID). The names of list elements must correspond to species names as in the second column of <i>orthogroups</i> . For instance, if there are two species in the <i>orthogroups</i> data frame named "SpeciesA" and "SpeciesB", <i>annotation</i> must be a list of 2 data frames, and each list element must be named "SpeciesA" and "SpeciesB". |
| correct_overclustering | Logical indicating whether to correct for overclustering in orthogroups. Default: TRUE.   |

**Value**

A data frame.

**Examples**

```
data(og)
data(interpro_ath)
data(interpro_bol)
# Subsetting annotation for demonstration purposes.
annotation <- list(Ath = interpro_ath[1:1000,], Bol = interpro_bol[1:1000,])
assess <- assess_orthogroups(og, annotation)
```

---

|               |   |
|---------------|---|
| assess_synnet | <i>Assess synteny network based on graph properties</i> |
|---------------|---|

---

**Description**

Assess synteny network based on graph properties

**Usage**

```
assess_synnet(synnet = NULL, cc_type = "average")
```

**Arguments**

|         |   |
|---------|---|
| synnet  | Edge list for the synteny network in a 2-column data frame, with columns 1 and 2 representing names of loci in anchor 1 and anchor 2, respectively. |
| cc_type | Type of clustering coefficient to be calculated. One of 'global' or 'average'. Default: 'average'.  |

**Details**

Network score is the product of the network's clustering coefficient, node count, and R squared for the scale-free topology fit.

**Value**

A data frame with the following variables:

**CC** Numeric representing clustering coefficient.

**Node\_count** Numeric representing number of nodes in the network.

**Rsquared** Numeric indicating the coefficient of determination for the scale-free topology fit.

**Score** Numeric representing network score, which is the product of 'CC' and 'Node\_number'.

**Examples**

```
data(synnet)
assess_synnet(synnet)
```

---

assess\_synnet\_list      *Assess list of synteny networks as in assess\_synnet*

---

**Description**

Assess list of synteny networks as in assess\_synnet

**Usage**

```
assess_synnet_list(synnet_list = NULL, cc_type = "average")
```

**Arguments**

**synnet\_list**      A list of networks, each network being an edge list as a 2-column data frame, with columns 1 and 2 representing names of loci in anchor 1 and anchor 2, respectively.

**cc\_type**          Type of clustering coefficient to be calculated. One of 'global' or 'average'. Default: 'average'.

**Value**

A data frame with the following variables:

**CC** Numeric representing clustering coefficient.

**Node\_count** Numeric representing number of nodes in the network.

**Rsquared** Numeric indicating the coefficient of determination for the scale-free topology fit.

**Score** Numeric representing network score, which is the product of 'CC' and 'Node\_number'.

**Network** Character of network name.

### Examples

```
set.seed(123)
data(synnet)
net1 <- synnet
net2 <- synnet[-sample(1:10000, 500), ]
net3 <- synnet[-sample(1:10000, 1000), ]
synnet_list <- list(net1 = net1, net2 = net2, net3 = net3)
assess_synnet_list(synnet_list)
```

---

batch\_summary

*BUSCO summary output for batch mode*

---

### Description

This object was created with the function `read_busco()` using a batch run of BUSCO on the genomes of *Herbaspirillum seropedicae* SmR1 and *Herbaspirillum rubrisubalbicans* M1.

### Usage

```
data(batch_summary)
```

### Format

A 2-column data frame with the following variables:

**Class** Factor of BUSCO classes

**Frequency** Numeric with the percentage of BUSCOs in each class.

**Lineage** Character with the lineage dataset used.

**File** Character with the name of the FASTA file used.

### Examples

```
data(batch_summary)
```

---

busco\_is\_installed

*Check if BUSCO is installed*

---

### Description

Check if BUSCO is installed

### Usage

```
busco_is_installed()
```

**Value**

Logical indicating whether BUSCO is installed or not.

**Examples**

```
busco_is_installed()
```

---

calculate\_H

*Calculate homogeneity scores for orthogroups*

---

**Description**

Calculate homogeneity scores for orthogroups

**Usage**

```
calculate_H(  
  orthogroup_df,  
  correct_overclustering = TRUE,  
  max_size = 200,  
  update_score = TRUE  
)
```

**Arguments**

- orthogroup\_df** Data frame with orthogroups and their associated genes and annotation. The columns **Gene**, **Orthogroup**, and **Annotation** are mandatory, and they must represent Gene ID, Orthogroup ID, and Annotation ID (e.g., Interpro/PFAM), respectively.
- correct\_overclustering** Logical indicating whether to correct for overclustering in orthogroups. Default: TRUE.
- max\_size** Numeric indicating the maximum orthogroup size to consider. If orthogroups are too large, calculating Sorensen-Dice indices for all pairwise combinations could take a long time, so setting a limit prevents that. Default: 200.
- update\_score** Logical indicating whether to replace scores with corrected scores or not. If FALSE, the dispersal term and corrected scores are returned as separate variables in the output data frame.

**Details**

Homogeneity is calculated based on pairwise Sorensen-Dice similarity indices between gene pairs in an orthogroup, and they range from 0 to 1. Thus, if all genes in an orthogroup share the same domain, the orthogroup will have a homogeneity score of 1. On the other hand, if genes in an orthogroup do not have any domain in common, the orthogroup will have a homogeneity score of 0. The percentage of orthogroups with size greater than **max\_size** will be subtracted from the

homogeneity scores, since too large orthogroups typically have very low scores. Additionally, users can correct for overclustering by penalizing protein domains that appear in multiple orthogroups (default).

### Value

A 2-column data frame with the variables **Orthogroup** and **Score**, corresponding to orthogroup ID and orthogroup score, respectively. If **update\_score = FALSE**, additional columns named **Dispersal** and **Score\_c** are added, which correspond to the dispersal term and corrected scores, respectively.

### Examples

```
data(og)
data(interpro_ath)
orthogroup_df <- merge(og[og$Species == "Ath", ], interpro_ath)
# Filter data to reduce run time
orthogroup_df <- orthogroup_df[1:10000, ]
H <- calculate_H(orthogroup_df)
```

---

compare\_genome\_stats *Compare user-defined assembly statistics with statistics of NCBI genomes*

---

### Description

This function helps users analyze their genome assembly stats in a context by comparing metrics obtained by users with "reference" metrics in closely-related organisms.

### Usage

```
compare_genome_stats(ncbi_stats = NULL, user_stats = NULL)
```

### Arguments

|            |  |
|------------|--|
| ncbi_stats | A data frame of summary statistics for a particular taxon obtained from the NCBI, as obtained with the function <code>get_genome_stats</code> .  |
| user_stats | A data frame with assembly statistics obtained by the user. A column named <b>accession</b> is mandatory, and it must contain unique identifiers for the genome(s) analyzed by the user. Dummy variables can be used as identifiers (e.g., "my_genome_001"), as long as they are unique. All other column containing assembly stats must have the same names as their corresponding columns in the data frame specified in <b>ncbi_stats</b> . For instance, stats on total number of genes and sequence length must be in columns named "gene_count_total" and "sequence_length", as in the <b>ncbi_stats</b> data frame. |



## Details

For each genome assembly statistic (e.g., "gene\_count\_total"), values in **user\_stats** are compared to a distribution of values from **ncbi\_stats**, and their percentile and rank in the distributions are reported.

## Value

A data frame with the following variables:

**accession** character, unique identifier as in user\_stats\$accession.

**variable** character, name of the genome assembly metric (e.g., "CC\_ratio").

**percentile** numeric, percentile in the distribution.

**rank** numeric, rank in the distribution (highest to lowest). For the variable "CC\_ratio", ranks go from lowest to highest.

## Examples

```
# Use case: user assembled a maize (Zea mays) genome

## Obtain stats for maize genomes on the NCBI
ncbi_stats <- get_genome_stats(taxon = "Zea mays")

## Create a data frame of stats for fictional maize genome
user_stats <- data.frame(
  accession = "my_lovely_maize",
  sequence_length = 2.4 * 1e9,
  gene_count_total = 50000,
  CC_ratio = 1
)

# Compare stats
compare_genome_stats(ncbi_stats, user_stats)
```

---

compare\_orthogroups     *Compare inferred orthogroups to a reference set*

---

## Description

Compare inferred orthogroups to a reference set

## Usage

```
compare_orthogroups(ref_orthogroups = NULL, test_orthogroups = NULL)
```

**Arguments**

ref\_orthogroups

Reference orthogroups in a 3-column data frame with columns **Orthogroup**, **Species**, and **Gene**. This data frame can be created from the 'Orthogroups.tsv' file generated by OrthoFinder with the function read\_orthogroups().

test\_orthogroups

Test orthogroups that will be compared to *ref\_orthogroups* in the same 3-column data frame format.

**Details**

This function compares a test set of orthogroups to a reference set and returns which orthogroups in the reference set are fully preserved in the test set (i.e., identical gene repertoire) and which are not. Species names (column 2) must be the same between reference and test set. If some species are not shared between reference and test sets, they will not be considered for the comparison.

**Value**

A 2-column data frame with the following variables:

**Orthogroup** Character of orthogroup IDs.

**Preserved** A logical vector of preservation status. It is TRUE if the orthogroup in the reference set is fully preserved in the test set, and FALSE otherwise.

**Examples**

```
set.seed(123)
data(og)
og <- og[1:5000, ]
ref <- og
# Shuffle genes to simulate a different set
test <- data.frame(
  Orthogroup = sample(og$Orthogroup, nrow(og), replace = FALSE),
  Species = og$Species,
  Gene = og$Gene
)
comparison <- compare_orthogroups(ref, test)

# Calculating percentage of preservation
sum(comparison$Preserved) / length(comparison$Preserved)
```

---

fit\_sft

*Goodness of fit test for the scale-free topology model*


---

**Description**

Goodness of fit test for the scale-free topology model

**Usage**

```
fit_sft(edges)
```

**Arguments**

**edges** A 2-column data frame with network edges represented in each. Columns 1 and 2 represent nodes 1 and 2 of each edge.

**Value**

A numeric scalar with the R squared for the scale-free topology fit.

**Examples**

```
data(synnet)
edges <- synnet
fit_sft(edges)
```

---

|                  |   |
|------------------|---|
| get_genome_stats | <i>Get summary statistics for genomes on NCBI using the NCBI Datasets API</i> |
|------------------|---|

---

**Description**

Get summary statistics for genomes on NCBI using the NCBI Datasets API

**Usage**

```
get_genome_stats(taxon = NULL, filters = NULL)
```

**Arguments**

**taxon** Taxon for which summary statistics will be retrieved, either as a character scalar (e.g., "brassicaceae") or as a numeric scalar representing NCBI Taxonomy ID (e.g., 3700).

**filters** (optional) A list of filters to use when querying the API in the form of key-value pairs, with keys in list names and values in list elements (e.g., `list(filters.reference_only = "true")`), see examples for details).

**Details**

Possible filters for the **filters** parameter can be accessed at [https://www.ncbi.nlm.nih.gov/datasets/docs/v2/reference-docs/rest-api/#get-/genome/taxon/-taxons-/dataset\\_report](https://www.ncbi.nlm.nih.gov/datasets/docs/v2/reference-docs/rest-api/#get-/genome/taxon/-taxons-/dataset_report).

**Value**

A data frame with the following variables:

**accession** character, accession number.  
**source** character, data source.  
**species\_taxid** numeric, NCBI Taxonomy ID.  
**species\_name** character, species' scientific name.  
**species\_common\_name** character, species' common name.  
**species\_ecotype** character, species' ecotype.  
**species\_strain** character, species' strain.  
**species\_isolate** character, species' isolate.  
**species\_cultivar** character, species' cultivar.  
**assembly\_level** factor, assembly level ("Complete", "Chromosome", "Scaffold", or "Contig").  
**assembly\_status** character, assembly status.  
**assembly\_name** character, assembly name.  
**assembly\_type** character, assembly type.  
**submission\_date** character, submission date (YYYY-MM-DD).  
**submitter** character, submitter name.  
**sequencing\_technology** character, sequencing technology.  
**atypical** logical, indicator of wheter the genome is atypical.  
**refseq\_category** character, RefSeq category.  
**chromosome\_count** numeric, number of chromosomes.  
**sequence\_length** numeric, total sequence length.  
**ungapped\_length** numeric, ungapped sequence length.  
**contig\_count** numeric, number of contigs.  
**contig\_N50** numeric, contig N50.  
**contig\_L50** numeric, contig L50.  
**scaffold\_N50** numeric, contig N50.  
**scaffold\_L50** numeric, contig L50.  
**GC\_percent** numeric, GC percentage (0-100).  
**annotation\_provider** character, name of annotation provider.  
**annotation\_release\_date** character, annotation release date (YYYY-MM-DD).  
**gene\_count\_total** numeric, total number of genes.  
**gene\_count\_coding** numeric, number of protein-coding genes.  
**gene\_count\_noncoding** numeric, number of non-coding genes.  
**gene\_count\_pseudogene** numeric, number of pseudogenes.  
**gene\_count\_other** numeric, number of other genes.  
**CC\_ratio** numeric, ratio of the number of contigs to the number of chromosomes.

## Examples

```
# Example 1: Search for A. thaliana genomes by tax ID
ex1 <- get_genome_stats(taxon = 3702)

# Example 2: Search for A. thaliana genomes by name
ex2 <- get_genome_stats(taxon = "Arabidopsis thaliana")

# Example 3: Search for chromosome-level Brassicaceae genomes
ex3 <- get_genome_stats(
  taxon = "brassicaceae",
  filters = list(filters.assembly_level = "chromosome")
)
```

---

interpro\_ath

*Intepro annotation for Arabidopsis thaliana's genes*

---

## Description

The annotation data were retrieved from PLAZA Dicots 5.0.

## Usage

```
data(interpro_ath)
```

## Format

A 2-column data frame:

**Gene** Character of gene IDs.

**Annotation** Character of Interpro domains.

## References

Van Bel, M., Silvestri, F., Weitz, E. M., Kreft, L., Botzki, A., Coppens, F., & Vandepoele, K. (2021). PLAZA 5.0: extending the scope and power of comparative and functional genomics in plants. *Nucleic acids research*.

## Examples

```
data(interpro_ath)
```

---

|              |   |
|--------------|---|
| interpro_bol | <i>Interpro annotation for Brassica oleraceae's genes</i> |
|--------------|---|

---

**Description**

The annotation data were retrieved from PLAZA Dicots 5.0.

**Usage**

```
data(interpro_bol)
```

**Format**

A 2-column data frame:

**Gene** Character of gene IDs.

**Annotation** Character of Interpro domains.

**References**

Van Bel, M., Silvestri, F., Weitz, E. M., Kreft, L., Botzki, A., Coppens, F., & Vandepoele, K. (2021). PLAZA 5.0: extending the scope and power of comparative and functional genomics in plants. *Nucleic acids research*.

**Examples**

```
data(interpro_bol)
```

---

|                     |                             |
|---------------------|-----------------------------|
| list_busco_datasets | <i>List BUSCO data sets</i> |
|---------------------|-----------------------------|

---

**Description**

List BUSCO data sets

**Usage**

```
list_busco_datasets()
```

**Value**

A hierarchically organized list of available data sets as returned by `busco --list-datasets`.

**Examples**

```
if(busco_is_installed()) {  
  list_busco_datasets()  
}
```

---

og *Orthogroups between Arabidopsis thaliana and Brassica oleraceae*

---

**Description**

Data obtained from PLAZA Dicots 5.0.

**Usage**

```
data(og)
```

**Format**

A 3-column data frame with the following variables:

**Orthogroup** Orthogroup ID.

**Species** Abbreviation for species' name.

**Gene** Gene ID

**References**

Van Bel, M., Silvestri, F., Weitz, E. M., Kreft, L., Botzki, A., Coppens, F., & Vandepoele, K. (2021). PLAZA 5.0: extending the scope and power of comparative and functional genomics in plants. *Nucleic acids research*.

**Examples**

```
data(og)
```

---

plot\_busco *Plot BUSCO summary output*

---

**Description**

Plot BUSCO summary output

**Usage**

```
plot_busco(summary_df = NULL)
```

**Arguments**

summary\_df Data frame with BUSCO summary output as returned by read\_busco().

**Value**

A ggplot object with a barplot of BUSCOs in each class.

### Examples

```
# Single file
result_dir <- system.file("extdata", package = "cogeqc")
summary_df <- read_busco(result_dir)
# Batch mode
data(batch_summary)
plot_busco(summary_df)
plot_busco(batch_summary)
```

---

plot\_duplications      *Plot species-specific duplications*

---

### Description

Plot species-specific duplications

### Usage

```
plot_duplications(stats_list = NULL)
```

### Arguments

`stats_list`      A list of data frames with Orthofinder summary stats as returned by the function `read_orthofinder_stats`.

### Value

A ggplot object with a barplot of number of species-specific duplications.

### Examples

```
dir <- system.file("extdata", package = "cogeqc")
stats_list <- read_orthofinder_stats(dir)
plot_duplications(stats_list)
```

---

plot\_genes\_in\_ogs      *Plot percentage of genes in orthogroups for each species*

---

### Description

Plot percentage of genes in orthogroups for each species

### Usage

```
plot_genes_in_ogs(stats_list = NULL)
```



**Arguments**

`stats_list` A list of data frames with Orthofinder summary stats as returned by the function `read_orthofinder_stats`.

**Value**

A ggplot object with a barplot of percentages of genes in orthogroups for each species.

**Examples**

```
dir <- system.file("extdata", package = "cogeqc")
stats_list <- read_orthofinder_stats(dir)
plot_genes_in_ogs(stats_list)
```

---

`plot_genome_stats` *Plot statistics on genome assemblies on the NCBI*

---

**Description**

Plot statistics on genome assemblies on the NCBI

**Usage**

```
plot_genome_stats(ncbi_stats = NULL, user_stats = NULL)
```

**Arguments**

`ncbi_stats` A data frame of summary statistics for a particular taxon obtained from the NCBI, as obtained with the function `get_genome_stats`.

`user_stats` (Optional) A data frame with assembly statistics obtained by the user. Statistics in this data frame are highlighted in red if this data frame is passed. A column named **accession** is mandatory, and it must contain unique identifiers for the genome(s) analyzed by the user. Dummy variables can be used as identifiers (e.g., "my\_genome\_001"), as long as they are unique. All other column containing assembly stats must have the same names as their corresponding columns in the data frame specified in `ncbi_stats`. For instance, stats on total number of genes and sequence length must be in columns named "gene\_count\_total" and "sequence\_length", as in the `ncbi_stats` data frame.

**Value**

A composition of ggplot objects made with patchwork.

## Examples

```
# Example 1: plot stats on maize genomes on the NCBI
## Obtain stats for maize genomes on the NCBI
ncbi_stats <- get_genome_stats(taxon = "Zea mays")

plot_genome_stats(ncbi_stats)

## Plot stats
# Example 2: highlight user-defined stats in the distribution
## Create a data frame of stats for fictional maize genome
user_stats <- data.frame(
  accession = "my_lovely_maize",
  sequence_length = 2.4 * 1e9,
  gene_count_total = 50000,
  CC_ratio = 1
)

plot_genome_stats(ncbi_stats, user_stats)
```

---

plot\_og\_overlap

*Plot pairwise orthogroup overlap between species*

---

## Description

Plot pairwise orthogroup overlap between species

## Usage

```
plot_og_overlap(stats_list = NULL, clust = TRUE)
```

## Arguments

|            |  |
|------------|--|
| stats_list | A list of data frames with Orthofinder summary stats as returned by the function <code>read_orthofinder_stats</code> . |
| clust      | Logical indicating whether to clust data based on overlap. Default: TRUE   |

## Value

A ggplot object with a heatmap.

## Examples

```
dir <- system.file("extdata", package = "cogeqc")
stats_list <- read_orthofinder_stats(dir)
plot_og_overlap(stats_list)
```

---

plot\_og\_sizes                      *Plot orthogroup sizes per species*

---

### Description

Plot orthogroup sizes per species

### Usage

```
plot_og_sizes(orthogroups = NULL, log = FALSE, max_size = NULL)
```

### Arguments

|             |   |
|-------------|---|
| orthogroups | A 3-column data frame with columns <b>Orthogroup</b> , <b>Species</b> , and <b>Gene</b> . This data frame can be created from the 'Orthogroups.tsv' file generated by OrthoFinder with the function read_orthogroups(). |
| log         | Logical indicating whether to transform orthogroups sizes with natural logarithms. Default: FALSE.  |
| max_size    | Numeric indicating the maximum orthogroup size to consider. If this parameter is not NULL, orthogroups larger than max_size (e.g., 100) will not be considered. Default: NULL.  |

### Value

A ggplot object with a violin plot.

### Examples

```
data(og)
plot_og_sizes(og, log = TRUE)
plot_og_sizes(og, max_size = 100)
plot_og_sizes(og, log = TRUE, max_size = 100)
```

---

plot\_orthofinder\_stats                      *Plot a panel with a summary of Orthofinder stats*

---

### Description

This function is a wrapper for plot\_species\_tree, plot\_duplications, plot\_genes\_in\_ogs, plot\_species\_specific\_ogs.

### Usage

```
plot_orthofinder_stats(tree = NULL, stats_list = NULL, xlim = c(0, 1))
```

**Arguments**

|            |   |
|------------|---|
| tree       | Tree object as returned by <code>treeio::read.*</code> , a family of functions in the <b>treeio</b> package to import tree files in multiple formats, such as Newick, Phylip, NEXUS, and others. If your species tree was inferred with Orthofinder (using STAG), the tree file is located in <i>Species_Tree/SpeciesTree_rooted_node_labels.txt</i> . Then, it can be imported with <code>treeio::read_tree(path_to_file)</code> . |
| stats_list | (optional) A list of data frames with Orthofinder summary stats as returned by the function <code>read_orthofinder_stats</code> . If this list is given as input, nodes will be labeled with the number of duplications.  |
| xlim       | Numeric vector of x-axis limits. This is useful if your node tip labels are not visible due to margin issues. Default: <code>c(0, 1)</code> .   |

**Value**

A panel of ggplot objects.

**Examples**

```
data(tree)
dir <- system.file("extdata", package = "cogeqc")
stats_list <- read_orthofinder_stats(dir)
plot_orthofinder_stats(tree, xlim = c(0, 1.5), stats_list = stats_list)
```

---

plot\_species\_specific\_ogs

*Plot number of species-specific orthogroups*

---

**Description**

Plot number of species-specific orthogroups

**Usage**

```
plot_species_specific_ogs(stats_list = NULL)
```

**Arguments**

|            |  |
|------------|--|
| stats_list | A list of data frames with Orthofinder summary stats as returned by the function <code>read_orthofinder_stats</code> . |
|------------|--|

**Value**

A ggplot object with a barplot of number of species-specific orthogroups for each species.

**Examples**

```
dir <- system.file("extdata", package = "cogeqc")
stats_list <- read_orthofinder_stats(dir)
plot_species_specific_ogs(stats_list)
```

---

plot\_species\_tree      *Plot species tree*

---

**Description**

Plot species tree

**Usage**

```
plot_species_tree(tree = NULL, xlim = c(0, 1), stats_list = NULL)
```

**Arguments**

|            |   |
|------------|---|
| tree       | Tree object as returned by <code>treeio::read.*</code> , a family of functions in the <b>treeio</b> package to import tree files in multiple formats, such as Newick, Phylip, NEXUS, and others. If your species tree was inferred with Orthofinder (using STAG), the tree file is located in <i>Species_Tree/SpeciesTree_rooted_node_labels.txt</i> . Then, it can be imported with <code>treeio::read_tree(path_to_file)</code> . |
| xlim       | Numeric vector of x-axis limits. This is useful if your node tip labels are not visible due to margin issues. Default: <code>c(0, 1)</code> .   |
| stats_list | (optional) A list of data frames with Orthofinder summary stats as returned by the function <code>read_orthofinder_stats</code> . If this list is given as input, nodes will be labeled with the number of duplications.  |

**Value**

A `ggtree/ggplot` object with the species tree.

**Examples**

```
data(tree)
plot_species_tree(tree)
```

---

read\_busco      *Read and parse BUSCO's summary report*

---

**Description**

Read and parse BUSCO's summary report

**Usage**

```
read_busco(result_dir = NULL)
```

**Arguments**

**result\_dir** Path to the directory where BUSCO results are stored. This function will look for the short\_summary\* file (single run) or short\_summary\* file (batch mode).

**Value**

A data frame with the following variables:

**Class** BUSCO class. One of **Complete\_SC**, **Complete\_duplicate**, **Fragmented**, or **Missing**

**Frequency** Frequency of BUSCOs in each class. If BUSCO was run in batch mode, this variable will contain relative frequencies. If BUSCO was run for a single file, it will contain absolute frequencies.

**Lineage** Name of the lineage dataset used.

**File (batch mode only)** Name of the input FASTA file.

**Examples**

```
result_dir <- system.file("extdata", package = "cogeqc")
df <- read_busco(result_dir)
```

---

read\_orthofinder\_stats

*Read and parse Orthofinder summary statistics*

---

**Description**

Read and parse Orthofinder summary statistics

**Usage**

```
read_orthofinder_stats(stats_dir = NULL)
```

**Arguments**

**stats\_dir** Path to directory containing Orthofinder's comparative genomics statistics. In your Orthofinder results directory, this directory is named **Comparative\_Genomics\_Statistics**.

**Value**

A list of data frames with the following elements:

1. **stats** A data frame of summary stats per species with the following variables:
  - Species** Factor of species names.
  - N\_genes** Numeric of number of genes.
  - N\_genes\_in\_OGs** Numeric of number of genes in orthogroups.
  - Perc\_genes\_in\_OGs** Numeric of percentage of genes in orthogroups.

**N\_ssOGs** Numeric of number of species-specific orthogroups.

**N\_genes\_in\_ssOGs** Numeric of number of genes in species-specific orthogroups.

**Perc\_genes\_in\_ssOGs** Numeric of percentage of genes in species-specific orthogroups.

**Dups** Integer with number of duplications per species.

2. **og\_overlap** A symmetric data frame of pairwise orthogroup overlap between species.
3. **duplications** A 2-column data frame with node IDs in the first column and number of gene duplications (50% support) in the second column.

### Examples

```
stats_dir <- system.file("extdata", package = "cogeqc")
ortho_stats <- read_orthofinder_stats(stats_dir)
```

---

|                  |   |
|------------------|---|
| read_orthogroups | <i>Read and parse orthogroups file created by OrthoFinder</i> |
|------------------|---|

---

### Description

This function converts the orthogroups file named **Orthogroups.tsv** to a parsed data frame.

### Usage

```
read_orthogroups(orthogroups_path = NULL)
```

### Arguments

**orthogroups\_path**  
Path to Orthogroups/Orthogroups.tsv file generated by OrthoFinder.

### Value

A 3-column data frame with orthogroups, species IDs and gene IDs, respectively.

### Author(s)

Fabricio Almeida-Silva

### Examples

```
path <- system.file("extdata", "Orthogroups.tsv.gz", package = "cogeqc")
og <- read_orthogroups(path)
```

run\_busco

*Run BUSCO assessment of assembly and annotation quality***Description**

Run BUSCO assessment of assembly and annotation quality

**Usage**

```
run_busco(
  sequence = NULL,
  outlabel = NULL,
  mode = c("genome", "transcriptome", "proteins"),
  lineage = NULL,
  auto_lineage = NULL,
  force = FALSE,
  threads = 1,
  outpath = NULL,
  download_path = tempdir()
)
```

**Arguments**

|               |   |
|---------------|---|
| sequence      | An object of class DNAStringSet/AAStringSet/RNAStringSet or path to FASTA file with the genome, transcriptome, or protein sequences to be analyzed. If there are many FASTA files in a directory, you can input the path to this directory, so BUSCO will be run in all FASTA files inside it.  |
| outlabel      | Character with a recognizable short label for analysis directory and files.   |
| mode          | Character with BUSCO mode. One of 'genome', 'transcriptome', or 'proteins'.   |
| lineage       | Character with name of lineage to be used.  |
| auto_lineage  | Character indicating whether BUSCO should determine optimum lineage path automatically. One of 'euk', 'prok', 'all', or NULL. If 'euk', it will determine optimum lineage path on eukaryote tree. If 'prok', it will determine optimum lineage path on non-eukaryote trees. If 'all', it will determine optimum lineage path for all trees. If NULL, it will not automatically determine lineage, and <i>lineage</i> must be manually specified. Default: NULL. |
| force         | Logical indicating whether existing runs with the same file names should be overwritten. Default: FALSE.  |
| threads       | Numeric with the number of threads/cores to use. Default: 1.  |
| outpath       | Path to results directory. If NULL, results will be stored in the current working directory. Default: NULL.   |
| download_path | Path to directory where BUSCO datasets will be stored after downloading. Default: tempdir().  |



**Value**

A character vector with the names of subdirectories and files in the results directory.

**Examples**

```
sequence <- system.file("extdata", "Hse_subset.fa", package = "cogeqc")
download_path <- paste0(tempdir(), "/datasets")
if(busco_is_installed()) {
  run_busco(sequence, outlabel = "Hse", mode = "genome",
            lineage = "burkholderiales_odb10",
            outpath = tempdir(), download_path = download_path)
}
```

---

synnet

*Synteny network for Brassica oleraceae, B. napus, and B. rapa*

---

**Description**

Synteny network for Brassica oleraceae, B. napus, and B. rapa

**Usage**

```
data(synnet)
```

**Format**

A 2-column data frame with the variables **anchor1** and **anchor2**, containing names of loci in anchor 1 and anchor 2, respectively.

**References**

Zhao, T., & Schranz, M. E. (2019). Network-based microsynteny analysis identifies major differences and genomic outliers in mammalian and angiosperm genomes. *Proceedings of the National Academy of Sciences*, 116(6), 2165-2174.

**Examples**

```
data(synnet)
```

---

tree

*Species tree for model species*

---

**Description**

The data used to create this object was retrieved from Orthofinder's example output for model species, available in [https://bioinformatics.plants.ox.ac.uk/davidemms/public\\_data/](https://bioinformatics.plants.ox.ac.uk/davidemms/public_data/).

**Usage**

```
data(tree)
```

**Format**

An object of class "phylo" as returned by `treeio::read.tree()`.

**References**

Emms, D. M., & Kelly, S. (2019). OrthoFinder: phylogenetic orthology inference for comparative genomics. *Genome biology*, 20(1), 1-14.

**Examples**

```
data(tree)
```

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