

# Package: cageminer (via r-universe)

June 18, 2026

**Title** Candidate Gene Miner

**Version** 1.18.0

**Description** This package aims to integrate GWAS-derived SNPs and coexpression networks to mine candidate genes associated with a particular phenotype. For that, users must define a set of guide genes, which are known genes involved in the studied phenotype. Additionally, the mined candidates can be given a score that favor candidates that are hubs and/or transcription factors. The scores can then be used to rank and select the top n most promising genes for downstream experiments.

**License** GPL-3

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

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

**biocViews** Software, SNP, FunctionalPrediction, GenomeWideAssociation, GeneExpression, NetworkEnrichment, VariantAnnotation, FunctionalGenomics, Network

**Encoding** UTF-8

**LazyData** false

**Roxygen** list(markdown = TRUE)

**RoxygenNote** 7.3.2

**Imports** ggplot2, rlang, ggbio, ggtext, GenomeInfoDb, GenomicRanges, IRanges, reshape2, methods, BioNERO

**Depends** R (>= 4.1)

**Suggests** testthat (>= 3.0.0), SummarizedExperiment, knitr, BiocStyle, rmarkdown, covr, sessioninfo

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**VignetteBuilder** knitr

**Config/pak/sysreqs** cmake libglpk-dev make libbz2-dev libicu-dev libjpeg-dev liblzma-dev libpng-dev libuv1-dev libxml2-dev libssl-dev perl xz-utils zlib1g-dev

**Repository** <https://bioc-release.r-universe.dev>

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chr_length	<i>Pepper chromosome lengths</i>
------------	----------------------------------

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### Description

Lengths of pepper chromosomes 1-12 in a GRanges object. The genome for which lengths were calculated (v1.55) was downloaded from <http://peppergenome.snu.ac.kr/download.php>

### Usage

```
data(chr_length)
```

### Format

A GRanges object

### Examples

```
data(chr_length)
```

---

gcn	<i>Simulation of the output list from BioNERO::exp2gcn() with pepper data</i>
-----	---

---

**Description**

This object is a list as returned by BioNERO::exp2gcn(), but only the element genes\_and\_modules is included. For running time issues, only genes in the cyan module were kept in the element genes\_and\_modules. All other list elements have been assigned NULL. The network was inferred using the code from the vignette.

**Usage**

```
data(gcn)
```

**Format**

A list with the elements returned by BioNERO::exp2gcn().

**Examples**

```
data(gcn)
```

---

gene_ranges	<i>Genomic coordinates of pepper genes</i>
-------------	--

---

**Description**

GRanges object with genomic coordinates of pepper genes downloaded from <http://peppergenome.snu.ac.kr/download.php>.

**Usage**

```
data(gene_ranges)
```

**Format**

A GRanges object

**Examples**

```
data(gene_ranges)
```

---

guides

*Guide genes associated with defense and resistance to oomycetes*

---

**Description**

The GO annotation was retrieved from PLAZA 4.0 Dicots.

**Usage**

```
data(guides)
```

**Format**

A data frame with genes in the first column and GO description in the second column.

**References**

Van Bel, M., Diels, T., Vancaester, E., Kreft, L., Botzki, A., Van de Peer, Y., ... & Vandepoele, K. (2018). PLAZA 4.0: an integrative resource for functional, evolutionary and comparative plant genomics. *Nucleic acids research*, 46(D1), D1190-D1196.

**Examples**

```
data(guides)
```

---

hubs

*Example hub genes for the network stored in the gcn object*

---

**Description**

The data frame was created using the code from the vignette.

**Usage**

```
data(hubs)
```

**Format**

Data frame with gene IDs, module and intramodular degree.

**Examples**

```
data(hubs)
```

---

mine_candidates	<i>Mine high-confidence candidate genes in a single step</i>
-----------------	--

---

## Description

Mine high-confidence candidate genes in a single step

## Usage

```
mine_candidates(
  gene_ranges = NULL,
  marker_ranges = NULL,
  window = 2,
  expand_intervals = TRUE,
  gene_col = "ID",
  exp = NULL,
  gcn = NULL,
  guides = NULL,
  metadata,
  metadata_cols = 1,
  sample_group,
  min_cor = 0.2,
  alpha = 0.05,
  ...
)
```

## Arguments

gene_ranges	A GRanges object with genomic coordinates of all genes in the genome.
marker_ranges	Genomic positions of SNPs. For a single trait, a GRanges object. For multiple traits, a GRangesList or CompressedGRangesList object, with each element of the list representing SNP positions for a particular trait.
window	Sliding window (in Mb) upstream and downstream relative to each SNP. Default: 2.
expand_intervals	Logical indicating whether or not to expand markers that are represented by intervals. This is particularly useful if users want to use a custom interval defined by linkage disequilibrium, for example. Default: TRUE.
gene_col	Column of the GRanges object containing gene ID. Default: "ID", the default for gff/gff3 files imported with rtracklayer::import.
exp	Expression data frame with genes in row names and samples in column names or a SummarizedExperiment object.
gcn	Gene coexpression network returned by BioNERO::exp2gcn().
guides	Guide genes as a character vector or as a data frame with genes in the first column and gene annotation class in the second column.

metadata	Sample metadata with samples in row names and sample information in the first column. Ignored if <code>exp</code> is a <code>SummarizedExperiment</code> object, as the <code>colData</code> will be extracted from the object.
metadata_cols	A vector (either numeric or character) indicating which columns should be extracted from column metadata if <code>exp</code> is a <code>SummarizedExperiment</code> object. The vector can contain column indices (numeric) or column names (character). By default, all columns are used.
sample_group	Level of sample metadata to be used for filtering in gene-trait correlation.
min_cor	Minimum correlation value for <code>BioNERO::gene_significance()</code> . Default: 0.2
alpha	Numeric indicating significance level. Default: 0.05
...	Additional arguments to <code>BioNERO::gene_significance</code> .

**Value**

A data frame with mined candidate genes and their correlation to the condition of interest.

**Examples**

```
data(pepper_se)
data(snp_pos)
data(gene_ranges)
data(guides)
data(gcn)
set.seed(1)
candidates <- mine_candidates(gene_ranges, snp_pos, exp = pepper_se,
                             gcn = gcn, guides = guides$Gene,
                             sample_group = "PRR_stress")
```

---

mine\_step1

*Step 1: Get all putative candidate genes for a given sliding window*


---

**Description**

For a user-defined sliding window relative to each SNP, this function will subset all genes whose genomic positions overlap with the sliding window.

**Usage**

```
mine_step1(gene_ranges, marker_ranges, window = 2, expand_intervals = TRUE)
```

**Arguments**

gene_ranges	A GRanges object with genomic coordinates of all genes in the genome.
marker_ranges	Genomic positions of SNPs. For a single trait, a GRanges object. For multiple traits, a GRangesList or CompressedGRangesList object, with each element of the list representing SNP positions for a particular trait.
window	Sliding window (in Mb) upstream and downstream relative to each SNP. Default: 2.
expand_intervals	Logical indicating whether or not to expand markers that are represented by intervals. This is particularly useful if users want to use a custom interval defined by linkage disequilibrium, for example. Default: TRUE.

**Value**

A GRanges or GRangesList object with the genomic positions of all putative candidate genes.

**See Also**

[findOverlaps-methods](#)

**Examples**

```
data(snp_pos)
data(gene_ranges)
genes <- mine_step1(gene_ranges, snp_pos, window = 2)
```

---

mine\_step2

*Step 2: Get candidates in modules enriched in guide genes*

---

**Description**

Step 2: Get candidates in modules enriched in guide genes

**Usage**

```
mine_step2(exp, gcn, guides, candidates, ...)
```

**Arguments**

exp	Expression data frame with genes in row names and samples in column names or a SummarizedExperiment object.
gcn	Gene coexpression network returned by <code>BioNERO::exp2gcn()</code> .
guides	Guide genes as a character vector or as a data frame with genes in the first column and gene annotation class in the second column.
candidates	Character vector of all candidate genes to be inspected.
...	Additional arguments to <code>BioNERO::module_enrichment</code>

**Value**

A list of 2 elements:

**candidates** Character vector of candidates after step 2

**enrichment** Data frame of results for enrichment analysis

**Examples**

```
data(pepper_se)
data(guides)
data(gcn)
set.seed(1)
mine2 <- mine_step2(
  exp = pepper_se,
  gcn = gcn,
  guides = guides$Gene,
  candidates = rownames(pepper_se)
)
```

---

mine\_step3

---

*Step 3: Select candidates based on gene significance*


---

**Description**

Step 3: Select candidates based on gene significance

**Usage**

```
mine_step3(
  exp,
  metadata,
  metadata_cols = 1,
  candidates,
  sample_group,
  min_cor = 0.2,
  alpha = 0.05,
  ...
)
```

**Arguments**

exp	Expression data frame with genes in row names and samples in column names or a SummarizedExperiment object.
metadata	Sample metadata with samples in row names and sample information in the first column. Ignored if exp is a SummarizedExperiment object, as the colData will be extracted from the object.

metadata_cols	A vector (either numeric or character) indicating which columns should be extracted from column metadata if <b>exp</b> is a SummarizedExperiment object. The vector can contain column indices (numeric) or column names (character). By default, all columns are used.
candidates	Character vector of candidate genes to be inspected.
sample_group	Level of sample metadata to be used for filtering in gene-trait correlation.
min_cor	Minimum correlation value for <code>BioNERO::gene_significance()</code> . Default: 0.2
alpha	Numeric indicating significance level. Default: 0.05
...	Additional arguments to <code>BioNERO::gene_significance</code> .

**Value**

A data frame with mined candidate genes and their correlation to the condition of interest.

**Examples**

```
data(pepper_se)
data(snp_pos)
data(gene_ranges)
data(guides)
data(gcn)
data(mine2)
set.seed(1)
mine3 <- mine_step3(
  exp = pepper_se,
  candidates = mine2$candidates,
  sample_group = "PRR_stress"
)
```

---

mine2

*Example output from mine\_step2()*

---

**Description**

The list was created using the example code from `mine_step()`.

**Usage**

```
data(mine2)
```

**Format**

List with elements 'candidates' (character vector) and 'enrichment' (data frame).

**Examples**

```
data(mine2)
```

---

mined_candidates	<i>Example output from mined_candidates()</i>
------------------	---

---

**Description**

The data frame was created using the code from the vignette.

**Usage**

```
data(mined_candidates)
```

**Format**

Data frame with an example of the output from mined\_candidates

**Examples**

```
data(mined_candidates)
```

---

pepper_se	<i>Gene expression data from Kim et al., 2018.</i>
-----------	--

---

**Description**

The data were filtered to keep only the top 4000 genes with highest RPKM values in PRR stress-related samples.

**Usage**

```
data(pepper_se)
```

**Format**

A SummarizedExperiment object.

**References**

Kim, MS., Kim, S., Jeon, J. et al. Global gene expression profiling for fruit organs and pathogen infections in the pepper, *Capsicum annuum* L.. *Sci Data* 5, 180103 (2018). <https://doi.org/10.1038/sdata.2018.103>

**Examples**

```
data(pepper_se)
```

---

plot_snp_circos	<i>Circos plot of SNP distribution across chromosomes</i>
-----------------	---

---

**Description**

Circos plot of SNP distribution across chromosomes

**Usage**

```
plot_snp_circos(genome_ranges, gene_ranges, marker_ranges)
```

**Arguments**

genome_ranges	A GRanges object with chromosome lengths.
gene_ranges	A GRanges object with genomic coordinates of all genes in the genome.
marker_ranges	Genomic positions of SNPs. For a single trait, a GRanges object. For multiple traits, a GRangesList or CompressedGRangesList object, with each element of the list representing SNP positions for a particular trait.

**Value**

A ggplot object with a circos plot of molecular marker distribution across chromosomes.

**Examples**

```
data(snp_pos)
data(gene_ranges)
data(chr_length)
p <- plot_snp_circos(chr_length, gene_ranges, snp_pos)
```

---

plot_snp_distribution	<i>Plot a barplot of SNP distribution across chromosomes</i>
-----------------------	--

---

**Description**

Plot a barplot of SNP distribution across chromosomes

**Usage**

```
plot_snp_distribution(marker_ranges)
```

**Arguments**

marker_ranges	Genomic positions of SNPs. For a single trait, a GRanges object. For multiple traits, a GRangesList or CompressedGRangesList object, with each element of the list representing SNP positions for a particular trait. List elements must have names for proper labelling.
---------------	---

**Value**

A ggplot object.

**Examples**

```
data(snp_pos)
p <- plot_snp_distribution(snp_pos)
```

---

score\_genes

*Score candidate genes and select the top n genes*

---

**Description**

Score candidate genes and select the top n genes

**Usage**

```
score_genes(
  mined_candidates,
  hubs = NULL,
  tfs = NULL,
  pick_top = 10,
  weight_tf = 2,
  weight_hub = 2,
  weight_both = 3
)
```

**Arguments**

mined_candidates	Data frame resulting from mine_candidates() or mine_step().
hubs	Character vector of hub genes.
tfs	Character vector of transcription factors.
pick_top	Number of top genes to select. Default: 10.
weight_tf	Numeric scalar with the weight to which correlation coefficients will be multiplied if the gene is a TF. Default: 2.
weight_hub	Numeric scalar with the weight to which correlation coefficients will be multiplied if the gene is a hub. Default: 2.
weight_both	Numeric scalar with the weight to which correlation coefficients will be multiplied if the gene is both a TF and a hub. Default: 3.

**Value**

Data frame with top n candidates and their scores.

## Examples

```
data(tfs)
data(hubs)
data(mined_candidates)
set.seed(1)
scored <- score_genes(mined_candidates, hubs$Gene, tfs$Gene_ID)
```

---

simulate_windows	<i>Simulate number of genes for each sliding window</i>
------------------	---

---

## Description

This function counts genes that are contained in sliding windows related to each SNP.

## Usage

```
simulate_windows(
  gene_ranges,
  marker_ranges,
  windows = seq(0.1, 2, by = 0.1),
  expand_intervals = TRUE
)
```

## Arguments

gene_ranges	A GRanges object with genomic coordinates of all genes in the genome.
marker_ranges	Genomic positions of SNPs. For a single trait, a GRanges object. For multiple traits, a GRangesList or CompressedGRangesList object, with each element of the list representing SNP positions for a particular trait.
windows	Sliding windows (in Mb) upstream and downstream relative to each SNP. Default: seq(0.1, 2, by = 0.1).
expand_intervals	Logical indicating whether or not to expand markers that are represented by intervals. This is particularly useful if users want to use a custom interval defined by linkage disequilibrium, for example. Default: TRUE.

## Details

By default, the function creates 20 sliding windows by expanding upstream and downstream boundaries for each SNP from 0.1 Mb (100 kb) to 2 Mb.

## Value

A ggplot object summarizing the results of the simulations.

## See Also

[findOverlaps-methods](#)

**Examples**

```
data(snp_pos)
data(gene_ranges)
simulate_windows(gene_ranges, snp_pos)
```

---

snp_pos	<i>Capsicum annuum</i> SNPs associated with resistance to <i>Phytophthora</i> root rot.
---------	---

---

**Description**

The SNPs in this data set were retrieved from Siddique et al., 2019, and they are associated to resistance to *Phytophthora* root rot.

**Usage**

```
data(snp_pos)
```

**Format**

A GRanges object.

**References**

Siddique, M.I., Lee, H.Y., Ro, N.Y. et al. Identifying candidate genes for *Phytophthora capsici* resistance in pepper (*Capsicum annuum*) via genotyping-by-sequencing-based QTL mapping and genome-wide association study. *Sci Rep* 9, 9962 (2019). <https://doi.org/10.1038/s41598-019-46342-1>

**Examples**

```
data(snp_pos)
```

---

tfs	<i>Pepper transcription factors</i>
-----	-------------------------------------

---

**Description**

Pepper transcription factors and their families retrieved from PlantTFDB 4.0.

**Usage**

```
data(tfs)
```

**Format**

A data frame with gene IDs in the first column and TF families in the second column.

**References**

Jin, J., Tian, F., Yang, D. C., Meng, Y. Q., Kong, L., Luo, J., & Gao, G. (2016). PlantTFDB 4.0: toward a central hub for transcription factors and regulatory interactions in plants. *Nucleic acids research*, gkw982.

**Examples**

```
data(tfs)
```

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