

# Package ‘artMS’

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**Type** Package

**Title** Analytical R tools for Mass Spectrometry

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**Description** artMS provides a set of tools for the analysis of proteomics label-free datasets. It takes as input the MaxQuant search result output (evidence.txt file) and performs quality control, relative quantification using MSstats, downstream analysis and integration. artMS also provides a set of functions to re-format and make it compatible with other analytical tools, including, SAINTq, SAINTexpress, Phosfate, and PHOTON. Check [<http://artms.org>](<http://artms.org>) for details.

**License** GPL (>= 3) + file LICENSE

**URL** <http://artms.org>

**BugReports** <https://github.com/biodavidjm/artMS/issues>

**Depends** R (>= 3.6.0)

**Imports** AnnotationDbi, biomaRt, bit64, circlize, cluster, ComplexHeatmap, corrplot, data.table, dplyr, factoextra, FactoMineR, getopt, ggdendro, ggplot2, gplots, ggrepel, gProfileR, graphics, grDevices, grid, limma, MSstats, openxlsx, org.Hs.eg.db, org.Mm.eg.db, PerformanceAnalytics, pheatmap, plotly, plyr, RColorBrewer, scales, seqinr, stats, stringr, tidyr, UpSetR, utils, VennDiagram, yaml

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## R topics documented:

artmsAnalysisQuantifications . . . . .	3
artmsAnnotateSpecie . . . . .	5
artmsAnnotationUniprot . . . . .	6
artmsAvgIntensityRT . . . . .	7
artmsChangeColumnName . . . . .	8
artmsConvertMetabolomics . . . . .	8
artmsDataPlots . . . . .	9
artmsEnrichLog2fc . . . . .	10
artmsEnrichProfiler . . . . .	11
artmsEvidenceToSaintExpress . . . . .	13
artmsEvidenceToSAINTq . . . . .	14
artmsFilterEvidenceContaminants . . . . .	15
artmsGeneratePhSiteExtended . . . . .	16
artmsIsEvidenceNewVersion . . . . .	17
artmsIsSpeciesSupported . . . . .	17
artmsLeaveOnlyUniprotEntryID . . . . .	18
artmsMapUniprot2Entrez . . . . .	19
artmsMergeEvidenceAndKeys . . . . .	20
artmsMsstatsSummary . . . . .	20
artmsPhosphateOutput . . . . .	21
artmsPhotonOutput . . . . .	22
artmsPlotHeatmapQuant . . . . .	23
artmsProtein2SiteConversion . . . . .	24
artmsQualityControlEvidenceBasic . . . . .	25
artmsQualityControlEvidenceExtended . . . . .	27
artmsQualityControlMetabolomics . . . . .	30
artmsQualityControlSummaryExtended . . . . .	31
artmsQuantification . . . . .	32
artmsResultsWide . . . . .	33
artmsSILACtoLong . . . . .	34
artmsSpectralCounts . . . . .	35
artmsVolcanoPlot . . . . .	35
artmsWriteConfigYamlFile . . . . .	36

artms_config . . . . .	37
artms_data_corum_mito_database . . . . .	39
artms_data_pathogen_LPN . . . . .	39
artms_data_pathogen_TB . . . . .	40
artms_data_ph_config . . . . .	40
artms_data_ph_contrast . . . . .	41
artms_data_ph_evidence . . . . .	42
artms_data_ph_keys . . . . .	42
artms_data_ph_msstats_modelqc . . . . .	43
artms_data_ph_msstats_results . . . . .	43
artms_data_randomDF . . . . .	43

## Index 44

artmsAnalysisQuantifications

*Analysis of the Relative Quantifications*

### Description

Analysis of relative quantifications, including:

- Annotations
- Summary files in different format (xls, txt) and shapes (long, wide)
- Numerous summary plots
- Enrichment analysis using Gprofiler
- PCA of quantifications
- Clustering analysis
- Basic imputation of missing values

### Usage

```
artmsAnalysisQuantifications(
  log2fc_file,
  modelqc_file,
  species,
  output_dir = "analysis_quant",
  outliers = c("keep", "iqr", "std"),
  enrich = TRUE,
  l2fc_thres = 1,
  choosePvalue = c("adjpvalue", "pvalue"),
  isBackground = "nobackground",
  isPtm = "global",
  mnbr = 2,
  isFluomics = FALSE,
  pathogen = "nopathogen",
  plotPvaluesLog2fcDist = TRUE,
  plotAbundanceStats = TRUE,
  plotReproAbundance = TRUE,
  plotCorrConditions = TRUE,
```

```

plotCorrQuant = TRUE,
plotPCAabundance = TRUE,
plotFinalDistributions = TRUE,
plotPropImputation = TRUE,
plotHeatmapsChanges = TRUE,
plotTotalQuant = TRUE,
plotClusteringAnalysis = TRUE,
data_object = FALSE,
verbose = TRUE
)

```

### Arguments

log2fc_file	(char) MSstats results file location
modelqc_file	(char) MSstats modelqc file location
species	(char) Select one species. Species currently supported for a full analysis (including enrichment analysis): <ul style="list-style-type: none"> <li>• HUMAN</li> <li>• MOUSE</li> </ul> <p>To find out species supported only for annotation check ?artmsIsSpeciesSupported()</p>
output_dir	(char) Name for the folder to output the results from the function. Default is current directory (recommended to provide a new folder name).
outliers	(char) It allows to keep or remove outliers. Options: <ul style="list-style-type: none"> <li>• keep (default): it keeps outliers 'keep', 'iqr', 'std'</li> <li>• iqr (recommended): remove outliers +/- 6 x Interquartile Range (IQR)</li> <li>• std : 6 x standard deviation</li> </ul>
enrich	(logical) Performed enrichment analysis using GprofileR? Only available for species HUMAN and MOUSE. TRUE (default if "human" or "mouse" are the species) or FALSE
l2fc_thres	(int) log2fc cutoff for enrichment analysis (default, l2fc_thres = 1.5)
choosePvalue	(char) specify whether pvalue or adjpvalue should use for the analysis. The default option is adjpvalue (multiple testing correction). But if the number of biological replicates for a given experiment is too low (for example n = 2), then choosePvalue = pvalue is recommended.
isBackground	(char) background of gene names for enrichment analysis. nobackground (default) will use the total number of genes detected. Alternatively provided the file path name to the background gene list.
isPtm	(char) Is a ptm-site quantification? <ul style="list-style-type: none"> <li>• global (default),</li> <li>• ptmsites (for site specific analysis),</li> <li>• ptmph (Jeff Johnson script output evidence file)</li> </ul>
mnbr	(int) minimal number of biological replicates for imputation and filtering. Default: mnbr = 2 (Proteins must be found in one of the conditions in at least 2 of the biological replicates)
isFluomics	(logical) Does this data belong to the FluOMICs project? TRUE or FALSE (default)
pathogen	(char) Is there a pathogen in the dataset as well? if it does not, then use pathogen = nopathogen (default). Pathogens available: tb (Tuberculosis), lpn (Legionella)

plotPvaluesLog2fcDist	(logical) If TRUE (default) plots pvalues and log2fc distributions
plotAbundanceStats	(logical) If TRUE (default) plots stats graphs about abundance values
plotReproAbundance	(logical) If TRUE plots reproducibility based on normalized abundance values
plotCorrConditions	(logical) If TRUE plots correlation between the different conditions
plotCorrQuant	(logical) if TRUE plots correlation between the available quantifications (comparisons)
plotPCAabundance	(logical) if TRUE performs PCA analysis of conditions using normalized abundance values
plotFinalDistributions	(logical) if TRUE plots distribution of both log2fc and pvalues
plotPropImputation	(logical) if TRUE plots proportion of overall imputation
plotHeatmapsChanges	(logical) if TRUE plots heatmaps of quantified changes (both all and significant only)
plotTotalQuant	(logical) if TRUE plots barplot of total number of quantifications per comparison
plotClusteringAnalysis	(logical) if TRUE performs clustering analysis between quantified comparisons (more than 1 comparison required)
data_object	(logical) flag to indicate whether the required files are data objects. Default is FALSE
verbose	(logical) TRUE (default) shows function messages

**Value**

(data.frame) summary of quantifications, including annotations, enrichments, etc

**Examples**

```
# Testing that the files cannot be empty
artmsAnalysisQuantifications(log2fc_file = NULL,
                             modelqc_file = NULL,
                             species = NULL,
                             output_dir = NULL)
```

---

artmsAnnotateSpecie     *Adding a column with the species name*

---

**Description**

Adding the species name to every protein. This makes more sense if there are more than one species in the dataset, which must be specified in the pathogen option. Influenza is a special case that it does not need to be specified, as far as the proteins were originally annotated as INFLUENZAGENE\_STRAIN (strains covered H1N1, H3N2, H5N1), as for example, NS1\_H1N1

**Usage**

```
artmsAnnotateSpecie(df, pathogen = "nopathogen", species)
```

**Arguments**

df	(data.frame) with a Protein column (of uniprot ids)
pathogen	(char) Is there a pathogen in the dataset as well? if it does not, then use pathogen = nopathogen (default). Supported tb (Tuberculosis), lpn (Legionella)
species	(char) Host organism (supported for now: human or mouse)

**Value**

(data.frame) The same data.frame but with an extra column specifying the species

**Examples**

```
# Adding a new column with the main species of the data. Easy.
# But the main functionality is to add both the host-species and a pathogen,
# which is not illustrated in this example
data_with_specie <- artmsAnnotateSpecie(df = artms_data_ph_msstats_results,
                                         species = "human")
```

---

```
artmsAnnotationUniprot
```

*Annotate table with Gene Symbol and Name based on Uniprot ID(s)*

---

**Description**

Annotate gene name and symbol based on uniprot ids. It will take the column from your data.frame specified by the columnid argument, search for the gene symbol, name, and entrez based on the species (species argument) and merge the information back to the input data.frame

**Usage**

```
artmsAnnotationUniprot(x, columnid, species, verbose = TRUE)
```

**Arguments**

x	(data.frame) to be annotated (or file path and name)
columnid	(char) The column with the uniprotkb ids
species	(char) The species name. Check ?artmsMapUniprot2Entrez to find out more about supported species.
verbose	(logical) TRUE (default) shows function messages

**Value**

(data.frame) with two new columns: Gene and Protein.name

## Examples

```
# This example adds annotations to the evidence file available in
# artMS, based on the column 'Proteins'.

evidence_anno <- artmsAnnotationUniprot(x = artms_data_ph_evidence,
                                       columnid = 'Proteins',
                                       species = 'human')
```

---

artmsAvgIntensityRT     *Summarize average intensity and retention time per protein*

---

## Description

Input an evidence file from MaxQuant and a file containing a list of proteins of interest (optional). The function will summarize from the evidence file and report back the average intensity, average retention time, and the average calibrated retention time. If a list of proteins is provided, then only those proteins will be summarized and returned.

## Usage

```
artmsAvgIntensityRT(
  evidence_file,
  protein_file = NULL,
  output_file = FALSE,
  species,
  verbose = TRUE
)
```

## Arguments

evidence_file	(char) The filepath to the MaxQuant searched data (evidence) file (txt tab delimited file).
protein_file	(char) The filepath to a file or vector containing a list of proteins of interest.
output_file	(char) The file name for the results (must have the extension .txt). If empty, then the results will be returned as an R object.
species	(char) The species name. Check ?artmsMapUniprot2Entrez for supported species
verbose	(logical) TRUE (default) shows function messages

## Value

An R object with the results and a file with the results (if the output\_file argument is provided). It contains averages of Intensity, Retention Time, Calibrated Retention Time

## Examples

```
ave_int <- artmsAvgIntensityRT(evidence_file = artms_data_ph_evidence,
                              species = "human")
```

---

artmsChangeColumnName *Change a specific column name in a given data.frame*

---

### Description

Making easier to change a column name in any data.frame

### Usage

```
artmsChangeColumnName(dataset, oldname, newname)
```

### Arguments

dataset	(data.frame) with the column name you want to change
oldname	(char) the old column name
newname	(char) the new name for that column

### Value

(data.frame) with the new specified column name

### Examples

```
artms_data_ph_evidence <- artmsChangeColumnName(  
  dataset = artms_data_ph_evidence,  
  oldname = "Phospho..STY.",  
  newname = "PH_STY")
```

---

artmsConvertMetabolomics

*Convert Markview Metabolomics file (alignment table) into a artMS compatible format*

---

### Description

artMS enables the relative quantification of untargeted polar metabolites using the alignment table generated by Markview. MarkerView is an ABSciex software that supports the files generated by Analyst software (.wiff) used to run our specific mass spectrometer (ABSciex Triple TOF 5600+). It also supports .t2d files generated by the Applied Biosystems 4700/4800 MALDI-TOF. MarkerView software is used to align mass spectrometry data from several samples for comparison. Using the import feature in the software, .wiff files (also .t2d MALDI-TOF files and tab-delimited .txt mass spectra data in mass-intensity format) are loaded for retention time alignment. Once the data files are selected, a series of windows will appear wherein peak finding, alignment, and filtering options can be entered and selected. These options include minimum spectral peak width, minimum retention time peak width, retention time and mass tolerance, and the ability to filter out peaks that do not appear in more than a user selected number of samples.

‘artmsConvertMetabolomics’ processes the markview file to enable QC analysis and relative quantification using the artMS functions



**Usage**

```
artmsConvertMetabolomics(input_file, out_file, id_file = NULL, verbose = TRUE)
```

**Arguments**

```
input_file      (char) Markview input file
out_file        (char) Output file name
id_file         (char) KEGG database
verbose         (logical) TRUE (default) shows function messages
```

**Value**

(text file) Outputs the converted output name

**Examples**

```
# Testing that the arguments cannot be null
artmsConvertMetabolomics(input_file = NULL,
                          out_file = NULL)
```

---

artmsDataPlots

*Individual Normalized abundance dot plots for every protein*


---

**Description**

Protein abundance dot plots for each unique uniprot id. It can take a long time

**Usage**

```
artmsDataPlots(input_file, output_file, verbose = TRUE)
```

**Arguments**

```
input_file      (char) File path and name to the -normalized.txt output file from MSstats
output_file     (char) Output file (path) name (add the .pdf extension)
verbose         (logical) TRUE (default) shows function messages
```

**Value**

(pdf) file with each individual protein abundance plot for each conditions

**Examples**

```
artmsDataPlots(input_file = "results/ab-results-mss-normalized.txt",
                output_file = "results/ab-results-mss-normalized.pdf")
```

---

artmsEnrichLog2fc      *Enrichment of changes in protein abundance or PTMs*

---

## Description

Enrichment analysis of the selected proteins

## Usage

```
artmsEnrichLog2fc(  
  dataset,  
  species,  
  background,  
  heatmaps = FALSE,  
  output_name = "enrichment.txt",  
  verbose = TRUE  
)
```

## Arguments

dataset	(data.frame) with a Gene and Comparison or Label (with the name of the comparisons specified in the contrast file) columns
species	(char) Specie, only supported "human" or "mouse"
background	(vector) Background genes for the enrichment analysis.
heatmaps	(logical) if TRUE generates heatmaps (pdf), FALSE (default) otherwise.
output_name	(char) Name of the annotation files, which will be used as well for the heatmaps (if heatmaps is selected) Default output_name = "enrichment.txt"
verbose	(logical) TRUE (default) shows function messages

## Value

(data.frame) Results from the enrichment analysis using Gprofiler and heatmaps (if selected)

## Examples

```
# The data must be annotated (Protein and Gene columns)  
data_annotated <- artmsAnnotationUniprot(  
  x = artms_data_ph_msstats_results,  
  columnid = "Protein",  
  species = "human")  
  
# And then the enrichment  
enrich_set <- artmsEnrichLog2fc(  
  dataset = data_annotated,  
  species = "human",  
  background = unique(data_annotated$Gene))
```

---

 artmsEnrichProfiler     *Enrichment analysis using GprofileR*


---

## Description

This function simplifies the enrichment analysis performed by the excellent tool GprofileR.

## Usage

```
artmsEnrichProfiler(
  x,
  categorySource = c("GO"),
  species,
  background = NA,
  verbose = TRUE
)
```

## Arguments

- x** (list, data.frame) List of protein ids. It can be anything: either a list of ids, or you could also send a data.frame and it will find the columns with the IDs. Is not cool? Multiple list can be also sent simultaneously, as for example running: `tmp <- split(enrichment$Gene, enrichment$cl_number, drop= TRUE)`
- categorySource** (vector) Resources providing the terms on which the enrichment will be performed. The supported resources by gprofiler are:
- GO (GO:BP, GO:MF, GO:CC): Gene Ontology (see more below)
  - KEGG: Biological pathways
  - REAC: Biological pathways (Reactome)
  - TF: Regulatory motifs in DNA (TRANSFAC TFBS)
  - MI: Regulatory motifs in DNA (miRBase microRNAs)
  - CORUM: protein complexes database
  - HP: Human Phenotype Ontology
  - HPA: Protein databases (Human Protein Atlas)
  - OMIM: Online Mendelian Inheritance in Man annotations:
  - BIOGRID: BioGRID protein-protein interactions The type of annotations for Gene Ontology:
  - Inferred from experiment (IDA, IPI, IMP, IGI, IEP)
  - Direct assay (IDA) / Mutant phenotype (IMP)
  - Genetic interaction (IGI) / Physical interaction (IPI)
  - Traceable author (TAS) / Non-traceable author (NAS) / Inferred by curator (IC)
  - Expression pattern (IEP) / Sequence or structural similarity (ISS) / Genomic context (IGC)
  - Biological aspect of ancestor (IBA) / Rapid divergence (IRD)
  - Reviewed computational analysis (RCA) / Electronic annotation (IEA)
  - No biological data (ND) / Not annotated or not in background (NA)

species	(char) Specie code: Organism names are constructed by concatenating the first letter of the name and the family name. Example: human - 'hsapiens', mouse - 'mmusculus'. Check gProfileR to find out more about supported species.
background	(vector) gene list to use as background for the enrichment analysis. Default: NA
verbose	(logical) TRUE (default) shows function messages

## Details

This function uses the following gprofiler arguments as default:

- ordered\_query = FALSE
- significant = TRUE
- exclude\_iea = TRUE
- underrep = FALSE
- evcodes = FALSE
- region\_query = FALSE
- max\_p\_value = 0.05
- min\_set\_size = 0
- max\_set\_size = 0
- min\_isect\_size = 0
- correction\_method = "analytical" #Options: "gSCS", "fdr", "bonferroni"
- hier\_filtering = "none"
- domain\_size = "known" # annotated or known
- numeric\_ns = ""
- png\_fn = NULL
- include\_graph = TRUE

## Value

The enrichment results as provided by gprofiler

## Examples

```
# annotate the MSstats results to get the Gene name
data_annotated <- artmsAnnotationUniprot(
  x = artms_data_ph_msstats_results,
  columnid = "Protein",
  species = "human")

# Filter the list of genes with a log2fc > 2
filtered_data <-
unique(data_annotated$Gene[which(data_annotated$log2FC > 2)])

# And perform enrichment analysis
data_annotated_enrich <- artmsEnrichProfiler(
  x = filtered_data,
  categorySource = c('KEGG'),
  species = "hsapiens",
  background = unique(data_annotated$Gene))
```

---

`artmsEvidenceToSaintExpress`*MaxQuant evidence file to SAINTexpress format*

---

### Description

Converts the MaxQuant evidence file to the 3 required files by SAINTexpress. One can choose to either use the spectral counts (use `msspc`) or the intensities (use `msint`) for the analysis.

### Usage

```
artmsEvidenceToSaintExpress(  
  evidence_file,  
  keys_file,  
  ref_proteome_file,  
  quant_variable = c("msspc", "msint"),  
  output_file,  
  verbose = TRUE  
)
```

### Arguments

<code>evidence_file</code>	(char) The evidence file path and name
<code>keys_file</code>	(char) Keys file with a SAINT column specifying test (T) and control (C) conditions
<code>ref_proteome_file</code>	(char) Reference proteome path file name in fasta format
<code>quant_variable</code>	(char) choose either <ul style="list-style-type: none"><li>• <code>msspc</code> (spectral counts, default) or</li><li>• <code>msint</code> (MS Intensity)</li></ul>
<code>output_file</code>	(char) Output file name (must have extension <code>.txt</code> )
<code>verbose</code>	(logical) TRUE (default) shows function messages

### Value

The 3 required files by SAINTexpress:

- `interactions.txt`
- `preys.txt`
- `baits.txt`

### Examples

```
# Testing that the files cannot be empty  
artmsEvidenceToSaintExpress(evidence_file = NULL,  
  keys_file = NULL, ref_proteome_file = NULL)
```

---

artmsEvidenceToSAINTq *MaxQuant evidence file to SAINTq format*

---

### Description

Converts the MaxQuant evidence file to the required files by SAINTq. The user can choose to use either peptides with spectral counts (use `msspc`) or the all the peptides (use `all`) for the analysis. The quantitative can be also chosen (either MS Intensity or Spectral Counts)

### Usage

```
artmsEvidenceToSAINTq(
  evidence_file,
  keys_file,
  output_dir = "artms_saintq",
  sc_option = c("all", "msspc"),
  fractions = FALSE,
  quant_variable = c("msint", "msspc"),
  verbose = TRUE
)
```

### Arguments

<code>evidence_file</code>	(char or data.frame) The evidence file path and name, or data.frame
<code>keys_file</code>	(char) Keys file with a SAINT column specifying test (T) and control (C) conditions
<code>output_dir</code>	(char) New directory to create and save files. Default is current directory (recommended to provide a new folder name).
<code>sc_option</code>	(char). Filter peptides with spectral counts only. Two options: <ul style="list-style-type: none"> <li>• <code>msspc</code>: use only peptides with <code>spectral_counts</code></li> <li>• <code>all</code> (default): all peptides detected (including the one resulting from the MaxQuant 'Match between run' algorithm)</li> </ul>
<code>fractions</code>	(logical) TRUE for 2D proteomics (fractions). Default: FALSE
<code>quant_variable</code>	(char) Select the quantitative variable. Two options available: <ul style="list-style-type: none"> <li>• <code>msint</code>: MS Intensity (default)</li> <li>• <code>msspc</code>: MS.MS.count (Spectral Counts)</li> </ul>
<code>verbose</code>	(logical) TRUE (default) shows function messages

### Details

After running the script, the new specified folder should contain the following files:

- `saintq-config-peptides`
- `saintq-config-proteins`
- `saintq_input_peptides.txt`
- `saintq_input_proteins.txt`

Then cd into the new folder and run either of the following two options (assuming that saintq is installed in your linux/unix/mac os x system):

```
> saintq config-saintq-peptides
```

or

```
> saintq config-saintq-proteins
```

### Value

The input files requires to run SAINTq

### Examples

```
# Testing that the files cannot be empty
artmsEvidenceToSAINTq (evidence_file = NULL,
                      keys_file = NULL,
                      output_dir = NULL)
```

---

artmsFilterEvidenceContaminants

*Remove contaminants and empty proteins from the MaxQuant evidence file*

---

### Description

Remove contaminants and erroneously identified 'reverse' sequences by MaxQuant, in addition to empty protein ids

### Usage

```
artmsFilterEvidenceContaminants(x, verbose = TRUE)
```

### Arguments

x	(data.frame) of the Evidence file
verbose	(logical) TRUE (default) shows function messages

### Value

(data.frame) without REV\_\_ and CON\_\_ Protein ids

### Examples

```
ef <- artmsFilterEvidenceContaminants(x = artms_data_ph_evidence)
```





---

artmsIsEvidenceNewVersion

*Check if a given evidence file was generated by a new version of MaxQuant (v>1)*

---

### Description

MaxQuant introduced changes in the column names and number of columns for the evidence file in version 1 (we think). This function check whether the evidence comes from the latest version of MaxQuant.

### Usage

```
artmsIsEvidenceNewVersion(evidence_file)
```

### Arguments

evidence\_file the evidence file name

### Value

(logical) TRUE if it is a newer version of MaxQuant, FALSE otherwise

### Examples

```
artmsIsEvidenceNewVersion(evidence_file = artms_data_ph_evidence)
```

---

artmsIsSpeciesSupported

*Check if a species is supported and available*

---

### Description

Given a species name, it checks whether is supported, and if supported, check whether the annotation package is installed.

### Usage

```
artmsIsSpeciesSupported(species, verbose = TRUE)
```

### Arguments

species (char) The species name. Species currently supported as part of artMS:

- HUMAN
- MOUSE

And the following species can be used as well, but the user needs to install the corresponding org.db package:

- ANOPHELES (install.packages(org.Ag.eg.db))
- BOVINE (install.packages(org.Bt.eg.db))

- WORM (install.packages(org.Ce.eg.db))
- CANINE (install.packages(org.Cf.eg.db))
- FLY (install.packages(org.Dm.eg.db))
- ZEBRAFISH (install.packages(org.Dr.eg.db))
- CHICKEN (install.packages(org.Gg.eg.db))
- RHESUS (install.packages(org.Mmu.eg.db))
- CHIMP (install.packages(org.Pt.eg.db))
- RAT (install.packages(org.Rn.eg.db))
- YEAST (install.packages(org.Sc.sgd.db))
- PIG (install.packages(org.Ss.eg.db))
- XENOPUS (install.packages(org.Xl.eg.db))

verbose (logical) TRUE (default) shows function messages

**Value**

(string) Name of the package for the given species

**Examples**

```
# Should return TRUE
artmsIsSpeciesSupported(species = "HUMAN")
artmsIsSpeciesSupported(species = "CHIMP")
```

---

artmsLeaveOnlyUniprotEntryID

*Leave only the Entry ID from a typical full Uniprot IDs in a given column*

---

**Description**

Downloading a Reference Uniprot fasta database includes several Uniprot IDs for every protein. If the regular expression available in Maxquant is not activated, the full id will be used in the Proteins, Lead Protein, and Leading Razor Protein columns. This script leaves only the Entry ID.

For example, values in a Protein column like this:

```
sp|P12345|Entry_name;sp|P54321|Entry_name2
```

will be replace by

```
'P12345;P54321'
```

**Usage**

```
artmsLeaveOnlyUniprotEntryID(x, columnid)
```

**Arguments**

x (data.frame) that contains the columnid  
 columnid (char) Column name with the full uniprot ids

**Value**

(data.frame) with only Entry IDs.

**Examples**

```
# Example of data frame with full uniprot ids and sequences
p <- c("sp|A6NIE6|RN3P2_HUMAN;sp|Q9NYV6|RRN3_HUMAN",
      "sp|A7E2V4|ZSWM8_HUMAN",
      "sp|A5A6H4|ROA1_PANTR;sp|P09651|ROA1_HUMAN;sp|Q32P51|RA1L2_HUMAN",
      "sp|A0FGR8|ESYT2_HUMAN")
s <- c("ALENDFNSPPRK", "GWGSPGRPK", "SSGPYGGGQYFAK", "VLVALASEELAK")
evidence <- data.frame(Proteins = p, Sequences = s, stringsAsFactors = FALSE)

# Replace the Proteins column with only Entry ids
evidence <- artmsLeaveOnlyUniprotEntryID(x = evidence, columnid = "Proteins")
```

---

```
artmsMapUniprot2Entrez
```

*Map GENE SYMBOL, NAME, AND ENTREZID to a vector of Uniprot IDS*

---

**Description**

Map GENE SYMBOL, NAME, AND ENTREZID to a vector of Uniprot IDS

**Usage**

```
artmsMapUniprot2Entrez(uniprotkb, species)
```

**Arguments**

uniprotkb	(vector) Vector of UniprotKB IDs
species	(char) The species name. Species currently supported as part of artMS: check ?artmsIsSpeciesSupported() to find out the list of supported species'

**Value**

(data.frame) with ENTREZID and GENENAMES mapped on UniprotKB ids

**Examples**

```
# Load an example with human proteins
exampleID <- c("Q6P996", "B1N8M6")
artmsMapUniprot2Entrez(uniprotkb = exampleID,
                      species = "HUMAN")
```

---

```
artmsMergeEvidenceAndKeys
```

*Merge evidence.txt (or summary.txt) with keys.txt files*

---

### Description

Merge the evidence and keys files on the given columns

### Usage

```
artmsMergeEvidenceAndKeys(
  x,
  keys,
  by = c("RawFile"),
  isSummary = FALSE,
  verbose = TRUE
)
```

### Arguments

x	(data.frame or char) The evidence data, either as data.frame or the file name (and path). It also works for the summary.txt file
keys	The keys data, either as a data.frame or file name (and path)
by	(vector) specifying the columns use to merge the evidence and keys. Default: by=c('RawFile')
isSummary	(logical) TRUE or FALSE (default)
verbose	(logical) TRUE (default) shows function messages

### Value

(data.frame) with the evidence and keys merged

### Examples

```
evidenceKeys <- artmsMergeEvidenceAndKeys(x = artms_data_ph_evidence,
                                          keys = artms_data_ph_keys)
```

---

```
artmsMsstatsSummary
```

*Summarize the MSStats results and data quantification*

---

### Description

Converts the MSStats results file to wide format (unique Protein ID and columns are the comparisons), as well as adds BioReplicate information about

- the Number of Unique Peptides,
- Spectral Counts
- Intensities for each protein. In cases where there are multiple values for a Protein-BioReplicate pair due to minute changes in sequence, the maximum value is taken for the pair. Any pairs without a value are assigned a value of NA.

**Usage**

```
artmsMsstatsSummary(
  evidence_file,
  prot_group_file,
  keys_file,
  results_file,
  return_df = FALSE,
  verbose = TRUE
)
```

**Arguments**

`evidence_file` (char or data.frame) The filepath to the MaxQuant searched data (evidence) file (txt tab delimited file). Only works for the newer versions of the evidence file.

`prot_group_file` (char) The filepath to the MaxQuant proteinGroups.txt file (txt tab delimited file) or data.frame

`keys_file` (char) The filepath to the keys file used with MSSstats (txt tab delimited file).

`results_file` (char) The filepath to the MSSstats results file in the default long format (txt tab delimited file or data.frame).

`return_df` (data.frame) Whether or not to return the results to the R environment upon completion. This is useful if this is being used in an R pipeline and you want to feed the results directly into the next stage of analysis via an R environment/terminal. Regardless, the results will be written to file. Default = FALSE

`verbose` (logical) TRUE (default) shows function messages

**Value**

(data.frame or txt file) with the summary

**Examples**

```
# Testing warning if files are not submitted
test <- artmsMsstatsSummary(evidence_file = NULL,
  prot_group_file = NULL,
  keys_file = NULL,
  results_file = NULL)
```

---

artmsPhosphateOutput     *Generate Phosphate Input file*

---

**Description**

It takes as input the imputedL2fcExtended.txt results generated by the artmsAnalysisQuantifications() function and generates the **Phosphate** input file (or data.frame) Please, notice that the only species supported by Phosphate is humans.

**Usage**

```
artmsPhosphateOutput(inputFile, output_dir = ".", verbose = TRUE)
```

**Arguments**

inputFile	(char) the imputedL2fcExtended.txt file name and location
output_dir	(char) Name of the folder to output results (Default: current directory. Recommended: phosfate_input)
verbose	(logical) TRUE (default) to show function messages

**Value**

Multiple output files (inputs of phosfate)

**Examples**

```
artmsPhosfateOutput(inputFile)
```

---

artmsPhotonOutput	<i>Generate PHOTON Input file</i>
-------------------	-----------------------------------

---

**Description**

It takes as input the imputedL2fcExtended.txt results generated by the artmsAnalysisQuantifications() function and generates the **PHOTON** input file. Please, notice that the only species supported by PHOTON is humans.

**Usage**

```
artmsPhotonOutput(inputFile, output_dir = ".", verbose = TRUE)
```

**Arguments**

inputFile	(char) the imputedL2fcExtended.txt file name and location
output_dir	(char) Name of the folder to output results (Default: current. Recommended: "photon_input_files" or similar)
verbose	(logical) TRUE (default) to show function messages

**Value**

Multiple output files (inputs of phosfate)

**Examples**

```
artmsPhotonOutput(inputFile)
```

---

artmsPlotHeatmapQuant *Outputs a heatmap of the MSStats results created using the log2fold changes*

---

### Description

Heatmap of the Relative Quantifications (MSStats results)

### Usage

```
artmsPlotHeatmapQuant(
  input_file,
  output_file = "quantifications_heatmap.pdf",
  species,
  labels = "*",
  cluster_cols = FALSE,
  display = "log2FC",
  lfc_lower = -2,
  lfc_upper = 2,
  whatPvalue = "adj.pvalue",
  FDR = 0.05,
  verbose = TRUE
)
```

### Arguments

input_file	(char) MSstats results.txt file and location (or data.frame of results)
output_file	(char) Output file name (pdf format) and location. Default: "quantifications_heatmap.pdf"
species	(char). Specie name to be able to add the Gene name. To find out more about the supported species check ?artmsMapUniprot2Entrez
labels	(vector) of uniprot ids if only specific labes would like to be plotted. Default: all labels
cluster_cols	(boolean) True or False to cluster columns. Default: FALSE
display	Metric to be displayed. Options: <ul style="list-style-type: none"> <li>• log2fc (default)</li> <li>• adj.pvalue</li> <li>• pvalue</li> </ul>
lfc_lower	(int) Lower limit for the log2fc. Default: -2
lfc_upper	(int) Upper limit for the log2fc. Default: +2
whatPvalue	(char) pvalue or adj.pvalue (default)
FDR	(int) Upper limit false discovery rate (or pvalue). Default: 0.05
verbose	(logical) TRUE (default) shows function messages

### Value

(pdf or ggplot2 object) heatmap of the MSStats results using the selected metric

**Examples**

```
artmsPlotHeatmapQuant(input_file = artms_data_ph_msstats_results,
                      species = "human",
                      output_file = NULL,
                      whatPvalue = "pvalue",
                      lfc_lower = -1,
                      lfc_upper = 1)
```

---

```
artmsProtein2SiteConversion
```

*Converts the Protein ID column of the evidence file selected by the user to mod-site-specific notation: ProteinID to ProteinID\_AAnumber notation*

---

**Description**

It enables the modified-peptide specific quantification by converting the Protein column of the evidence file selected by the user to an ProteinID\_AAnumber notation. In this way, each of the modified peptides can be quantified independently across conditions.

!!

WARNING: we have detected a version of MaxQuant (>1.6.3.0) outputs a "Modified sequence" column of the evidence file that has two important changes for the annotation of phosphorylation:

- Uses p instead of (ph)
- The modified residue (i.e. STY) is the residue on the right of the p, instead of the residue to the left of (ph), as usual. We have introduced a modification to detect and address this issue, but we advice the user to double check both the new evidence file with the introduce new notation and the -mapping.txt file and check that there are no NA values for the notation of phophopeptides.

!!

**Usage**

```
artmsProtein2SiteConversion(
  evidence_file,
  ref_proteome_file,
  column_name = c("Leading razor protein", "Leading proteins", "Proteins"),
  output_file,
  mod_type,
  overwrite_evidence = FALSE,
  verbose = TRUE
)
```

**Arguments**

evidence\_file (char) The evidence file name and location

ref\_proteome\_file

(char) The reference proteome used as database to search the evidence.txt file with MaxQuant. It will be used to map the modified peptide to the protein sequence and find the site location. Therefore, it does not use the MaxQuant's Phospho (STY)Sites.txt



column_name	(char) The Protein Column Name to map. Options: <ul style="list-style-type: none"> <li>• Leadind razor protein (default)</li> <li>• Leading protein</li> <li>• Proteins It only supports Uniprot Entry IDs and RefSeq, but it might work for other database IDs</li> </ul>
output_file	(char) Output file name (ptmsites-evidence.txt recommended)
mod_type	(char) The posttranslational modification. Options: <ul style="list-style-type: none"> <li>• UB: Protein Ubiquitination</li> <li>• PH: Protein Phosphorylation</li> <li>• AC: Protein Acetylation</li> </ul>
overwrite_evidence	(logical) if <output_file> is the same as <evidence_file>, overwrite_evidence = FALSE (default) doesn't allow to overwrite the evidence file. Otherwise, overwrite_evidence = TRUE allows to overwrite the evidence_file (this option might be activated if the user allows to use the same ptm-sites-evidence.txt file to re-annotate all the Protein IDs columns)
verbose	(logical) TRUE (default) shows function messages

**Value**

(file) Return a new evidence file with the specified Protein id column modified by adding the sequence site location(s) + postranslational modification(s) to the uniprot entry / refseq id.

Output ID examples: A34890\_ph3; Q64890\_ph24\_ph456; Q64890\_ub34\_ub129\_ub234; Q64890\_ac35.

**Examples**

```
# Testing warning if files are not submitted.
artmsProtein2SiteConversion(evidence_file = NULL, ref_proteome_file = NULL,
output_file = NULL)
```

---

```
artmsQualityControlEvidenceBasic
```

*Quality Control analysis of the MaxQuant evidence file*

---

**Description**

Quality Control analysis of the MaxQuant evidence file

**Usage**

```
artmsQualityControlEvidenceBasic(
  evidence_file,
  keys_file,
  prot_exp = c("AB", "PH", "UB", "AC", "APMS"),
  fractions = 0,
  output_name = "qcBasic_evidence",
  isSILAC = FALSE,
  plotINTDIST = FALSE,
  plotREPRO = FALSE,
```

```

plotCORMAT = TRUE,
plotINTMISC = TRUE,
plotPTMSTATS = TRUE,
printPDF = TRUE,
verbose = TRUE
)

```

### Arguments

evidence_file	(char or data.frame) The evidence file path and name, or data.frame
keys_file	(char or data.frame) The keys file path and name or data.frame
prot_exp	(char) Proteomics experiment. 4 options available: <ul style="list-style-type: none"> <li>• APMS: affinity purification mass spectrometry</li> <li>• AB: protein abundance</li> <li>• PH: protein phosphorylation</li> <li>• UB: protein ubiquitination (aka ubiquitylation)</li> <li>• AC: protein acetylation</li> </ul>
fractions	(binary) Is a fractionated experiment? <ul style="list-style-type: none"> <li>• 1 yes</li> <li>• 0 no (default)</li> </ul>
output_name	(char) prefix output name (no extension). Default: "qcBasic_evidence"
isSILAC	if TRUE processes SILAC input files. Default is FALSE
plotINTDIST	if TRUE plots both <i>Box-dot plot</i> and <i>Jitter plot</i> of biological replicates based on MS (raw) intensity values, otherwise FALSE (default)
plotREPRO	if TRUE plots a correlation dotplot for all the combinations of biological replicates of conditions, based on MS Intensity values using features (peptide+charge). Otherwise FALSE (default)
plotCORMAT	if TRUE (default) plots a <ul style="list-style-type: none"> <li>• <i>Correlation matrix</i> for all the biological replicates using MS Intensity values,</li> <li>• <i>Clustering matrix</i> of the MS Intensities</li> </ul>
plotINTMISC	if TRUE (default) plots several pages, including bar plots of <i>Total Sum of Intensities in BioReplicates</i> , <i>Total Sum of Intensities in Conditions</i> , <i>Total Peptide Counts in BioReplicates</i> , <i>Total Peptide Counts in conditions</i> separated by categories: CON: contaminants, PROT peptides, REV reversed sequences used by MaxQuant to estimate the FDR; <i>Box plots</i> of MS Intensity values per biological replicates and conditions; <i>bar plots</i> of total intensity (excluding contaminants) by bioreplicates and conditions; Barplots of <i>total feature counts</i> by bioreplicates and conditions.
plotPTMSTATS	IF TRUE (default) plots stats related to the selected modification, including: <i>bar plot of peptide counts and intensities</i> , broken by PTM/other categories; bar plots of <i>total sum-up of MS intensity values</i> by other/PTM categories.
printPDF	If TRUE (default) prints out the pdfs. Warning: plot objects are not returned due to the large number of them.
verbose	(logical) TRUE (default) shows function messages

### Value

Quality control files and plots

**Examples**

```

artmsQualityControlEvidenceBasic(evidence_file = artms_data_ph_evidence,
                                keys_file = artms_data_ph_keys,
                                prot_exp = "PH",
                                isSILAC = FALSE,
                                plotINTDIST = FALSE,
                                plotREPRO = TRUE,
                                plotCORMAT = FALSE,
                                plotINTMISC = FALSE,
                                plotPTMSTATS = FALSE,
                                printPDF = FALSE,
                                verbose = FALSE)

# But we recommend the following test:
# 1. Go to a working directory:
# setwd("/path/to/your/working/directory/")
# 2. Run the following command to print out all the pdf files
# artmsQualityControlEvidenceBasic(evidence_file = artms_data_ph_evidence,
#                                   keys_file = artms_data_ph_keys,
#                                   prot_exp = "PH")
# 3. Check your working directory and you should find pdf files with
# all the QC plots

```

---

artmsQualityControlEvidenceExtended

*Extended Quality Control of the MaxQuant evidence.txt file*

---

**Description**

Performs quality control based on the information available in the MaxQuant evidence.txt file.

**Usage**

```

artmsQualityControlEvidenceExtended(
  evidence_file,
  keys_file,
  output_name = "qcExtended_evidence",
  isSILAC = FALSE,
  plotPSM = TRUE,
  plotIONS = TRUE,
  plotTYPE = TRUE,
  plotPEPTIDES = TRUE,
  plotPEPTOVERLAP = TRUE,
  plotPROTEINS = TRUE,
  plotPROTOVERLAP = TRUE,
  plotPIO = TRUE,
  plotCS = TRUE,
  plotME = TRUE,
  plotMOCD = TRUE,
  plotPEPICV = TRUE,
  plotPEPDETECT = TRUE,
  plotPROTICV = TRUE,

```

```

plotPROTDETECT = TRUE,
plotIDoverlap = TRUE,
plotPCA = TRUE,
plotSP = TRUE,
printPDF = TRUE,
verbose = TRUE
)

```

## Arguments

evidence_file	(char or data.frame) The evidence file path and name, or data.frame
keys_file	(char or data.frame) The keys file path and name or data.frame
output_name	(char) prefix output name (no extension). Default: "qcExtended_evidence"
isSILAC	if TRUE processes SILAC input files. Default is FALSE
plotPSM	(logical) TRUE generates peptide-spectrum-matches (PSMs) statistics plot: Page 1 shows the number of PSMs confidently identified in each BioReplicate. If replicates are present, Page 2 shows the mean number of PSMs per condition with error bar showing the standard error of the mean. Note that potential contaminant proteins are plotted separately.
plotIONS	(logical) TRUE generates peptide ion statistics plot: A peptide ion is defined in the context of m/z, in other words, an unique peptide sequence may give rise to multiple ions with different charge state and/or amino acid modification. Page 1 shows the number of ions confidently identified in each BioReplicate . If replicates are present, Page 2 shows the mean number of peptide ions per condition with error bar showing the standard error of the mean. Note that potential contaminant proteins are plotted separately.
plotTYPE	(logical) TRUE generates identification type statistics plot: MaxQuant classifies each peptide identification into different categories (e.g., MSMS, MULTI-MSMS, MULTI-SECPEP). Page 1 shows the distribution of identification type in each BioReplicate
plotPEPTIDES	(logical) TRUE generates peptide statistics plot: Page 1 shows the number of unique peptide sequences (disregard the charge state or amino acid modifications) confidently identified in each BioReplicate. If replicates are present, Page 2 shows the mean number of peptides per condition with error bar showing the standard error of the mean. Note that potential contaminant proteins are plotted separately. Pages 3 and 4 show peptide identification intersection between BioReplicates (the bars are ordered by degree or frequency, respectively), and Page 4 shows the intersections across conditions instead of BioReplicates.
plotPEPTOVERLAP	(logical) TRUE Show peptide identification intersection between BioReplicates and Conditions
plotPROTEINS	(logical) TRUE generates protein statistics plot: Page 1 shows the number of protein groups confidently identified in each BioReplicate. If replicates are present, Page 2 shows the mean number of protein groups per condition with error bar showing the standard error of the mean. Note that potential contaminant proteins are plotted separately. Pages 3 and 4 show peptide identification intersection between BioReplicates (the bars are ordered by degree or frequency, respectively), and Page 4 shows the intersections across conditions instead of BioReplicates.
plotPROTOVERLAP	(logical) TRUE Show protein identification intersection between BioReplicates and Conditions

plotPIO	(logical) TRUE generates oversampling statistics plot: Page 1 shows the proportion of all peptide ions (including peptides matched across runs) fragmented once, twice and thrice or more. Page 2 shows the proportion of peptide ions (with intensity detected) fragmented once, twice and thrice or more. Page 3 shows the proportion of peptide ions (with intensity detected and MS/MS identification) fragmented once, twice and thrice or more
plotCS	(logical) TRUE generates charge state plot: Page 1 shows the charge state distribution of PSMs confidently identified in each BioReplicate.
plotME	(logical) TRUE generates precursor mass error plot: Page 1 shows the distribution of precursor error for all PSMs confidently identified in each BioReplicate.
plotMOCD	(logical) TRUE generates precursor mass-over-charge plot: Page 1 shows the distribution of precursor mass-over-charge for all PSMs confidently identified in each BioReplicate.
plotPEPICV	(logical) TRUE generates peptide intensity coefficient of variance (CV) plot: The CV is calculated for each feature (peptide ion) identified in more than one replicate. Page 1 shows the distribution of CV's for each condition, while Page 2 shows the distribution of CV's within 4 bins of intensity (i.e., 4 quantiles of average intensity).
plotPEPDETECT	(logical) TRUE generates peptide detection frequency plot: Page 1 summarizes the frequency that each peptide is detected across BioReplicates of each condition, showing the percentage of peptides detected once, twice, thrice, and so on (for whatever number of replicates each condition has).
plotPROTICV	(logical) TRUE generates protein intensity coefficient of variance (CV) plot: The CV is calculated for each protein (after summing the peptide intensities) identified in more than one replicate. Page 1 shows the distribution of CV's for each condition, while Page 2 shows the distribution of CV's within 4 bins of intensity (i.e., 4 quantiles of average intensity).
plotPROTDETECT	(logical) TRUE generates protein detection frequency plot: Page 1 summarizes the frequency that each protein group is detected across BioReplicates of each condition, showing the percentage of proteins detected once, twice, thrice, and so on (for whatever number of replicates each condition has). Page 2 shows the feature (peptide ion) intensity distribution within each BioReplicate (potential contaminant proteins are plot separately). Page 3 shows the density of feature intensity for different feature types (i.e., MULTI-MSMS, MULTI-SECPEP).
plotIDoverlap	(logical) TRUE generates pairwise identification heatmap overlap: Pages 1 and 2 show pairwise peptide and protein overlap between any 2 BioReplicates, respectively.
plotPCA	(logical) TRUE generates PCA and pairwise intensity correlation: Page 1 and 3 show pairwise peptide and protein intensity correlation and scatter plot between any 2 BioReplicates, respectively. Page 2 and 4 show Principal Component Analysis at the intensity level for both peptide and proteins, respectively.
plotSP	(logical) TRUE generates sample quality metrics: Page 1 shows missing cleavage distribution of all peptides confidently identified in each BioReplicate. Page 2 shows the fraction of peptides with at least one methionine oxidized in each BioReplicate.
printPDF	If TRUE (default) prints out the pdfs. Warning: plot objects are not returned due to the large number of them.
verbose	(logical) TRUE (default) shows function messages

**Details**

all the plots are generated by default

**Value**

A number of QC plots based on the evidence file

**Examples**

```
# Testing warning if files are not submitted
test <- artmsQualityControlEvidenceExtended(evidence_file = NULL,
keys_file = NULL)
```

---

```
artmsQualityControlMetabolomics
```

*Quality Control analysis of the evidence-like metabolomics dataset*

---

**Description**

Quality Control analysis of the evidence-like metabolomics dataset

**Usage**

```
artmsQualityControlMetabolomics(
  evidence_file,
  keys_file,
  met_exp = c("MV"),
  output_name = "qcPlots_metab",
  plotINTDIST = FALSE,
  plotCORMAT = TRUE,
  plotINTMISC = TRUE,
  printPDF = TRUE,
  verbose = TRUE
)
```

**Arguments**

evidence_file	(char or data.frame) The evidence file path and name, or data.frame
keys_file	(char or data.frame) The keys file path and name or data.frame
met_exp	(char) Metabolomics experiment. Only one option available (so far): <ul style="list-style-type: none"> <li>• MV: Markview output</li> </ul>
output_name	(char) prefix output name (no extension). Default: "qcPlots_metab"
plotINTDIST	if TRUE (default) plots both <i>Box-dot plot</i> and <i>Jitter plot</i> of biological replicates based on MS (raw) intensity values.
plotCORMAT	if TRUE (default) generates up to 3 pdf files for technical replicates, biological replicates, and conditions. Each pdf file contains: <ul style="list-style-type: none"> <li>• <i>Correlation matrix</i> for all the biological replicates using MS Intensity values,</li> <li>• <i>Clustering matrix</i> of the MS Intensities and correlation distribution</li> </ul>

- *histogram* of the distribution of correlations

plotINTMISC	if TRUE (default) plots several pages, including bar plots of <i>Total Sum of Intensities in BioReplicates</i> , <i>Total Sum of Intensities in Conditions</i> , <i>Total Feature Counts in BioReplicates</i> , <i>Total Feature Counts in conditions</i> separated by categories (INT: has a intensity value NOINT: no intensity value ) <i>Box plots</i> of MS Intensity values per biological replicates and conditions; <i>bar plots</i> of total intensity by bioreplicates and conditions; Barplots of <i>total feature counts</i> by bioreplicates and conditions.
printPDF	If TRUE (default) prints out the pdfs. Warning: plot objects are not returned due to the large number of them.
verbose	(logical) TRUE (default) shows function messages

**Value**

Quality control files and plots for metabolomics

**Examples**

```
# Testing that input arguments cannot be null
artmsQualityControlMetabolomics(evidence_file = NULL,
                                keys_file = NULL,
                                met_exp = "MV")
```

---

artmsQualityControlSummaryExtended  
*Quality Control of the MaxQuant summary.txt file*

---

**Description**

Performs quality control based on the information available in the MaxQuant summary.txt file.

**Usage**

```
artmsQualityControlSummaryExtended(
  summary_file,
  keys_file,
  output_name = "qcExtended_summary",
  isFractions = FALSE,
  plotMS1SCANS = TRUE,
  plotMS2 = TRUE,
  plotMSMS = TRUE,
  plotISOTOPE = TRUE,
  printPDF = TRUE,
  verbose = TRUE
)
```

**Arguments**

summary_file	(char or data.frame) The evidence file path and name, or data.frame
keys_file	(char or data.frame) The keys file path and name or data.frame
output_name	(char) prefix output name (no extension). Default: "qcExtended_summary"
isFractions	(logical) TRUE if it is a 2D experiment (fractions). Default: FALSE
plotMS1SCANS	(logical) TRUE generates MS1 scan counts plot: Page 1 shows the number of MS1 scans in each BioReplicate. If replicates are present, Page 2 shows the mean number of MS1 scans per condition with error bar showing the standard error of the mean. If isFractions TRUE, each fraction is a stack on the individual bar graphs.
plotMS2	(logical) TRUE generates MS2 scan counts plot: Page 1 shows the number of MSs scans in each BioReplicate. If replicates are present, Page 2 shows the mean number of MS1 scans per condition with error bar showing the standard error of the mean. If isFractions TRUE, each fraction is a stack on the individual bar graphs.
plotMSMS	(logical) TRUE generates MS2 identification rate (%) plot: Page 1 shows the fraction of MS2 scans confidently identified in each BioReplicate. If replicates are present, Page 2 shows the mean rate of MS2 scans confidently identified per condition with error bar showing the standard error of the mean. If isFractions TRUE, each fraction is a stack on the individual bar graphs.
plotISOTOPE	(logical) TRUE generates Isotope Pattern counts plot: Page 1 shows the number of Isotope Patterns with charge greater than 1 in each BioReplicate. If replicates are present, Page 2 shows the mean number of Isotope Patterns with charge greater than 1 per condition with error bar showing the standard error of the mean. If isFractions TRUE, each fraction is a stack on the individual bar graphs.
printPDF	If TRUE (default) prints out the pdfs. Warning: plot objects are not returned due to the large number of them.
verbose	(logical) TRUE (default) shows function messages

**Value**

A number of plots from the summary file

**Examples**

```
# Testing warning if files are not submitted
test <- artmsQualityControlSummaryExtended(summary_file = NULL,
keys_file = NULL)
```

---

artmsQuantification     *Relative quantification using MSstats*

---

**Description**

Relative quantification using MSstats including:

- plots
- quantifications (log2fc, pvalues, etc)
- normalized abundance values



**Usage**

```
artmsQuantification(yaml_config_file, data_object = FALSE, verbose = TRUE)
```

**Arguments**

```
yaml_config_file      (char) The yaml file name and location
data_object           (logical) flag to indicate whether the configuration file is a string to a file that
                        should be opened or
verbose               (logical) TRUE (default) shows function messages
```

**Value**

The relative quantification of the conditions and comparisons specified in the keys/contrast file resulting from running MSstats, in addition to quality control plots (if selected)

**Examples**

```
artmsQuantification("artms-ab-config.yaml")
```

---

artmsResultsWide	<i>Reshape the MSstats results file from long to wide format</i>
------------------	--

---

**Description**

Converts the normal MSstats results.txt file into "wide" format where each row represents a unique protein's results, and each column represents the comparison made by MSstats. The fold change and p-value of each comparison will be its own column.

**Usage**

```
artmsResultsWide(
  results_msstats,
  output_file = NULL,
  select_pvalues = c("adjpvalue", "pvalue"),
  species,
  verbose = TRUE
)
```

**Arguments**

```
results_msstats      (char) Input file name and location (MSstats results.txt file)
output_file          (char) Output file name and location (e.g. results-wide.txt). If NULL (de-
                        fault) returns an R object (data.frame)
select_pvalues       (char) Either
                        • pvalue or
                        • adjpvalue (default)
```

species (char) Specie name for annotation purposes. Check ?artmsMapUniprot2Entrez to find out more about the supported species (e.g species = "human")

verbose (logical) TRUE (default) shows function messages

**Value**

(output file tab delimited) reshaped file with unique protein ids and as many columns log2fc and adj.pvalues as comparisons available

**Examples**

```
ph_results_wide <- artmsResultsWide(
  results_msstats = artms_data_ph_msstats_results,
  output_file = NULL,
  species = "human")
```

---

artmsSILACtoLong      *Convert the SILAC evidence file to MSstats format*

---

**Description**

Converting the evidence file from a SILAC search to a format compatible with MSstats. It basically modifies the Raw.files adding the Heavy and Light label

**Usage**

```
artmsSILACtoLong(evidence_file, output = NULL, verbose = TRUE)
```

**Arguments**

evidence\_file (char) Text filepath to the evidence file

output (char) Text filepath of the output name. If NULL it does not write the output

verbose (logical) TRUE (default) shows function messages

**Value**

(data.frame) with SILAC data processed for MSstats (and output file)

**Examples**

```
evidence2silac <- artmsSILACtoLong(evidence_file = "silac.evidence.txt",
  output = "silac-evidence.txt")
```

---

artmsSpectralCounts     *Outputs the spectral counts from the MaxQuant evidence file.*

---

### Description

Outputs the spectral counts from the MaxQuant evidence file.

### Usage

```
artmsSpectralCounts(  
  evidence_file,  
  keys_file,  
  output_file = NULL,  
  verbose = TRUE  
)
```

### Arguments

evidence\_file     (char) Maxquant evidence file or data object  
keys\_file         (char) Keys file with the experimental design or data object  
output\_file       (char) Output file name (add .txt extension). If NULL (default) it returns a data.frame object  
verbose           (logical) TRUE (default) shows function messages

### Value

A txt file with biological replicates, protein id, and spectral count columns

### Examples

```
summary_spectral_counts <- artmsSpectralCounts(  
  evidence_file = artms_data_ph_evidence,  
  keys_file = artms_data_ph_keys)
```

---

artmsVolcanoPlot     *Volcano plot (log2fc / pvalues)*

---

### Description

It generates a scatter-plot used to quickly identify changes

**Usage**

```
artmsVolcanoPlot(
  mss_results,
  output_name = "volcano_plot.pdf",
  lfc_upper = 1,
  lfc_lower = -1,
  whatPvalue = "adj.pvalue",
  FDR = 0.05,
  PDF = TRUE,
  decimal_threshold = 16,
  verbose = TRUE
)
```

**Arguments**

mss_results	(data.frame or file) Selected MSstats results
output_name	(char) Name for the output file (don't forget the .pdf extension)
lfc_upper	(numeric) log2fc upper threshold (positive value)
lfc_lower	(numeric) log2fc lower threshold (negative value)
whatPvalue	(char) pvalue or adj.pvalue (default)
FDR	(numeric) False Discovery Rate threshold
PDF	(logical) Option to generate pdf format. Default: T
decimal_threshold	(numeric) Decimal threshold for the pvalue. Default: 16 (10 <sup>-16</sup> )
verbose	(logical) TRUE (default) shows function messages

**Value**

(pdf) of a volcano plot

**Examples**

```
artmsVolcanoPlot(mss_results = artms_data_ph_msstats_results,
  whatPvalue = "pvalue",
  PDF = FALSE)
```

---

artmsWriteConfigYamlFile

*Write out a template file of the artMS configuration file (yaml)*

---

**Description**

Creates a template file of the artMS configuration file, which is required to run `artmsQuantification`. Check `?artms_config` and the vignettes to find out more about the details of the structure of the file and how to fill it up

**Usage**

```
artmsWriteConfigYamlFile(
  config_file_name = "artms_config_file.yaml",
  verbose = TRUE
)
```

**Arguments**

`config_file_name` (char) The name for the configuration file. It must have a .yaml extension. If NULL, it returns the config as a yaml object

`verbose` (logical) TRUE (default) shows function messages

**Value**

A file (or yaml data object) of the artMS configuration file

**Examples**

```
config_empty <- artmsWriteConfigYamlFile(config_file_name = NULL)
```

---

artms_config	<i>artMS configuration template</i>
--------------	-------------------------------------

---

**Description**

The configuration file in yaml format contains the configuration details required to run `artmsQuantification()`, which includes quality control functions

**Usage**

```
artms_config
```

**Format**

The configuration (yaml) file contains the following sections:

- files**
  - `evidence` : /path/to/the/evidence.txt
  - `keys` : /path/to/the/keys.txt
  - `contrasts` : /path/to/the/contrast.txt
  - `summary` : /path/to/the/summary.txt
  - `output` : /path/to/the/output/results/results.txt
- qc**
  - `basic`: 1 # 1 = yes; 0 = no
  - `extended`: 1 # 1 = yes; 0 = no
  - `extendedSummary`: 0 # 1 = yes; 0 = no
- data**
  - `enabled` : 1 # 1 = yes; 0 = no
  - `fractions`:
    - `enabled` : 0 # 1 for protein fractionation
  - `silac`:

- enabled : 0 # 1 for SILAC experiments
- filters:
  - enabled : 1
- contaminants : 1
- protein\_groups : remove #remove, keep
- modifications : ab # PH, UB, AB, APMS
- sample\_plots : 1 # correlation plots
- msstats**
  - enabled : 1
  - msstats\_input : # blank if not previous msstats input file is available
  - profilePlots : none # before, after, before-after, none
  - normalization\_method : equalizeMedians # globalStandards (include a reference protein(s) ), equalizeMedians, quantile, 0
  - normalization\_reference : #should be a value in the Protein column
  - summaryMethod : TMP # "TMP"(default) means Tukey's median polish, which is robust estimation method. "linear" uses linear mixed model. "logOfSum" conducts log2 (sum of intensities) per run.
  - censoredInt : NA # Missing values are censored or at random. 'NA' (default) assumes that all 'NA's in 'Intensity' column are censored. '0' uses zero intensities as censored intensity. In this case, NA intensities are missing at random. The output from Skyline should use '0'. Null assumes that all NA intensities are randomly missing.
  - cutoffCensored : minFeature # Cutoff value for censoring. only with censoredInt='NA' or '0'. Default is 'minFeature', which uses minimum value for each feature.'minFeatureNRun' uses the smallest between minimum value of corresponding feature and minimum value of corresponding run. 'minRun' uses minimum value for each run.
  - MBimpute : 1 # only for summaryMethod="TMP" and censoredInt='NA' or '0'. TRUE (default) imputes 'NA' or '0' (depending on censoredInt option) by Accelerated failure model. FALSE uses the values assigned by cutoffCensored.
  - feature\_subset: all # allhighQuality : highQuality seems to be buggy right now
- output\_extras**
  - output\_extras :
    - enabled : 1 # if 0, it wont do anything in this section
  - annotate :
    - enabled: 1 # 1|0 whether to annotate the proteins in the results or not
  - species : HUMAN # Supported species: HUMAN, MOUSE, ANOPHELES, ARABIDOPSIS, BOVINE, WORM, CANINE, FLY, ZEBRAFISH, ECOLI\_STRAIN\_K12, ECOLI\_STRAIN\_SAKAI, CHICKEN, RHESUS, MALARIA, CHIMP, RAT, YEAST, PIG, XENOPUS
  - plots:
    - volcano: 1
    - heatmap: 1
    - LFC : -1.5 1.5 # Range of minimal log2fc
    - FDR : 0.05
    - heatmap\_cluster\_cols : 0
    - heatmap\_display : log2FC # log2FC or pvalue

---

artms\_data\_corum\_mito\_database

*CORUM Protein Complexes database use for complex enrichment analysis*

---

### Description

The list of protein complexes has been enriched with mitochondria proteins from mouse, as described in this paper:

2018 - Ruchi Masand, Esther Paulo, Dongmei Wu , Yangmeng Wang, Danielle L. Swaney, David Jimenez-Morales, Nevan J. Krogan, and Biao Wang Proteome Imbalance of Mitochondrial Electron Transport Chain in Brown Adipocytes Leads to Metabolic Benefits. *Cell Metab.* 2018 Mar 06; 27(3):616-629.e4

### Usage

artms\_data\_corum\_mito\_database

### Format

Tab delimited file.

To find out more about the format and columns available at CORUM, please visit this [link](#)

### Details

LAST CORUM DOWNLOAD DATE: 2017-08-01

---

artms\_data\_pathogen\_LPN

*LPN PATHOGEN: Legionella pneumophila subsp. pneumophila (strain Philadelphia 1 / ATCC 33152 / DSM 7513) UNIPROT IDS*

---

### Description

LPN PATHOGEN: Legionella pneumophila subsp. pneumophila (strain Philadelphia 1 / ATCC 33152 / DSM 7513) UNIPROT IDS

### Usage

artms\_data\_pathogen\_LPN

### Format

A data.frame of Entry IDs

---

artms\_data\_pathogen\_TB

*TB PATHOGEN: Mycobacterium tuberculosis (strain ATCC 35801 / TMC 107 / Erdman) UNIPROTS IDS*

---

### Description

TB PATHOGEN: Mycobacterium tuberculosis (strain ATCC 35801 / TMC 107 / Erdman) UNIPROTS IDS

### Usage

artms\_data\_pathogen\_TB

### Format

A data.frame of Entry IDs

---

artms\_data\_ph\_config *artMS configuration for the available PH dataset*

---

### Description

The configuration file with default options to run the available PH dataset with ‘artmsQuantification()’

### Usage

artms\_data\_ph\_config

### Format

The configuration (yaml) file contains the following sections:

- files**
  - evidence : artms\_data\_ph\_evidence
  - keys : artms\_data\_ph\_keys
  - contrasts : artms\_data\_ph\_contrast
  - summary :
  - output : "results.txt"
- qc**
  - basic: 0
  - extended: 0
  - extendedSummary: 0 =
- data**
  - enabled : 1
  - fractions:
    - enabled : 0
  - silac:
    - enabled : 0
  - filters:



- enabled : 1
- contaminants : 1
- protein\_groups : remove
- modifications : PH
- sample\_plots : 1
- msstats**
  - enabled : 1
  - msstats\_input : # blank if not previous msstats input file is available
  - profilePlots : none # before, after, before-after, none
  - normalization\_method : equalizeMedians
  - normalization\_reference : #should be a value in the Protein column
  - summaryMethod : TMP
  - censoredInt : NA
  - cutoffCensored : minFeature
  - MBimpute : 1
  - feature\_subset: all
- output\_extras**
  - output\_extras :
    - enabled : 1
  - annotate :
    - enabled: 1
  - species : HUMAN
  - plots:
    - volcano: 1
    - heatmap: 1
    - LFC : -1 1
    - FDR : 0.05
    - heatmap\_cluster\_cols : 0
    - heatmap\_display : log2FC

---

artms\_data\_ph\_contrast

*Contrast example for the PH dataset*


---

## Description

Contrast file with the relative quantification to be performed for the two conditions available in the example dataset: "Cal33-HSC6". See vignette for more details on how to prepare the contrast file.

## Usage

```
artms_data_ph_contrast
```

## Format

list with one comparison: "Cal33-HSC6"

---

artms\_data\_ph\_evidence

*Evidence file example*

---

### Description

Evidence file from a PH experiment consisting of two head and neck cancer cell lines ("Conditions" "Ca133" and "HSC6").

Unfortunately, the number of lines was reduced to 1/8 due to bioconductor limitations on data size, which means that this data is not very representative of a real evidence file. However, both the full evidence.txt and keys.txt file are available at: <http://kroganlab.ucsf.edu/artms/ph/evidence.txt>  
<http://kroganlab.ucsf.edu/artms/ph/keys.txt>

### Usage

artms\_data\_ph\_evidence

### Format

A data frame with all the columns available in an evidence file generated with MaxQuant version 1.6.2.3

---

artms\_data\_ph\_keys

*Keys File Example*

---

### Description

the artMS keys file provides the details of the experimental design for any given proteomics experiment.

This particular example belongs to a PH experiment consisting of two head and neck cancer cell lines ("Conditions" "Ca133" and "HSC6"), with 2 biological replicates each (in this reduced version)

### Usage

artms\_data\_ph\_keys

### Format

Tab delimited file with the following columns:

**Raw.file** Raw file processed. Each one should be a unique biological (or technical) replicate

**IsotopeLabelType** Type of labeling. L is used for label free experiments

**Condition** Label for conditions. VERY IMPORTANT: Only alpha-numeric characters and underscore (\_) are allowed

**BioReplicate** Label for the Biological replicates. VERY IMPORTANT: Use the same labeling for bioreplicate as the Condition, but adding a dash (-) corresponding to the number of biological replicate. For example, for Condition "Ca1", use Ca1-1, Ca1-2, Ca1-3, etc for the bioreplicates

**Run** The MS run number

---

artms\_data\_ph\_msstats\_modelqc  
*MSstats modelQC example*

---

**Description**

Normalized data obtained from the `artmsQuantification()` step of the PH dataset (global analysis)

**Usage**

```
artms_data_ph_msstats_modelqc
```

**Format**

A data frame resulting from running the latest version of `MSstats::groupComparison` function required as input for `artmsAnalysisQuantifications()`

---

artms\_data\_ph\_msstats\_results  
*MSstats results example*

---

**Description**

Relative quantification results obtained running `MSstats` on the available PH datasets (global analysis). Changes in protein phosphorylation were quantified between two conditions (check `artms_data_ph_contrast`)

**Usage**

```
artms_data_ph_msstats_results
```

**Format**

A data frame resulting from running the latest version of `MSstats`

---

artms\_data\_randomDF *Random data set*

---

**Description**

Dataset randomly generated for testing purposes

**Usage**

```
artms_data_randomDF
```

**Format**

A data frame with 100 rows and 10 variables:

Dataset generated using this code

```
data.frame(replicate(10, sample(0:1, 100, rep=TRUE)))
```

# Index

- \* **APMS**
  - artmsEvidenceToSaintExpress, [13](#)
  - artmsEvidenceToSAINTq, [14](#)
- \* **MSSStats**,
  - artmsMsstatsSummary, [20](#)
- \* **MaxQuant**,
  - artmsAvgIntensityRT, [7](#)
  - artmsMsstatsSummary, [20](#)
- \* **QC**,
  - artmsQualityControlEvidenceBasic, [25](#)
  - artmsQualityControlMetabolomics, [30](#)
- \* **SAINT**,
  - artmsEvidenceToSaintExpress, [13](#)
  - artmsEvidenceToSAINTq, [14](#)
- \* **SAINTExpress**,
  - artmsEvidenceToSaintExpress, [13](#)
- \* **SAINTq**,
  - artmsEvidenceToSAINTq, [14](#)
- \* **abundance**,
  - artmsDataPlots, [9](#)
- \* **ac**
  - artmsProtein2SiteConversion, [24](#)
- \* **analysis**,
  - artmsAnalysisQuantifications, [3](#)
- \* **annotation**,
  - artmsAnnotateSpecie, [5](#)
  - artmsAnnotationUniprot, [6](#)
  - artmsIsSpeciesSupported, [17](#)
  - artmsLeaveOnlyUniprotEntryID, [18](#)
  - artmsMapUniprot2Entrez, [19](#)
- \* **caliberated**
  - artmsAvgIntensityRT, [7](#)
- \* **check**
  - artmsIsEvidenceNewVersion, [17](#)
- \* **cleanup**,
  - artmsFilterEvidenceContaminants, [15](#)
- \* **columns**
  - artmsChangeColumnName, [8](#)
- \* **config**,
  - artmsWriteConfigYamlFile, [36](#)
- \* **contaminants**
  - artmsFilterEvidenceContaminants, [15](#)
- \* **control**,
  - artmsQualityControlEvidenceBasic, [25](#)
  - artmsQualityControlMetabolomics, [30](#)
- \* **convert**,
  - artmsProtein2SiteConversion, [24](#)
  - artmsSILACtoLong, [34](#)
- \* **convert**
  - artmsConvertMetabolomics, [8](#)
- \* **data.frame**,
  - artmsChangeColumnName, [8](#)
- \* **datasets**
  - artms\_config, [37](#)
  - artms\_data\_corum\_mito\_database, [39](#)
  - artms\_data\_pathogen\_LPN, [39](#)
  - artms\_data\_pathogen\_TB, [40](#)
  - artms\_data\_ph\_config, [40](#)
  - artms\_data\_ph\_contrast, [41](#)
  - artms\_data\_ph\_evidence, [42](#)
  - artms\_data\_ph\_keys, [42](#)
  - artms\_data\_ph\_msstats\_modelqc, [43](#)
  - artms\_data\_ph\_msstats\_results, [43](#)
  - artms\_data\_randomDF, [43](#)
- \* **dotplots**,
  - artmsDataPlots, [9](#)
- \* **driver**,
  - artmsQuantification, [32](#)
- \* **enrichment**
  - artmsEnrichLog2fc, [10](#)
  - artmsEnrichProfiler, [11](#)
- \* **evidence**,
  - artmsAvgIntensityRT, [7](#)
  - artmsIsEvidenceNewVersion, [17](#)
  - artmsMergeEvidenceAndKeys, [20](#)
  - artmsMsstatsSummary, [20](#)
  - artmsProtein2SiteConversion, [24](#)
  - artmsQualityControlEvidenceExtended, [27](#)
  - artmsQualityControlMetabolomics, [30](#)

- 30
- \* **evidence**
  - artmsQualityControlEvidenceBasic, 25
  - artmsSILACtoLong, 34
  - artmsSpectralCounts, 35
- \* **external**,
  - artmsGeneratePhSiteExtended, 16
- \* **file**,
  - artmsIsEvidenceNewVersion, 17
- \* **files**
  - artmsPhosphateOutput, 21
  - artmsPhotonOutput, 22
- \* **function**
  - artmsQuantification, 32
- \* **generate**,
  - artmsPhosphateOutput, 21
  - artmsPhotonOutput, 22
- \* **heatmap**,
  - artmsPlotHeatmapQuant, 23
- \* **ids**
  - artmsLeaveOnlyUniprotEntryID, 18
  - artmsMapUniprot2Entrez, 19
- \* **input**,
  - artmsIsEvidenceNewVersion, 17
- \* **intensity**,
  - artmsAvgIntensityRT, 7
- \* **keys**
  - artmsMergeEvidenceAndKeys, 20
  - artmsQualityControlEvidenceExtended, 27
  - artmsQualityControlSummaryExtended, 31
- \* **log2fc**
  - artmsPlotHeatmapQuant, 23
- \* **main**,
  - artmsQuantification, 32
- \* **merge**,
  - artmsMergeEvidenceAndKeys, 20
- \* **metabolomics**,
  - artmsConvertMetabolomics, 8
- \* **metabolomics**
  - artmsQualityControlMetabolomics, 30
- \* **msstats**,
  - artmsResultsWide, 33
- \* **outputs**,
  - artmsPhosphateOutput, 21
  - artmsPhotonOutput, 22
- \* **ph**,
  - artmsProtein2SiteConversion, 24
- \* **phosphate**
  - artmsGeneratePhSiteExtended, 16
- \* **plot**,
  - artmsVolcanoPlot, 35
- \* **plot**
  - artmsDataPlots, 9
- \* **ptm**,
  - artmsProtein2SiteConversion, 24
- \* **qc**,
  - artmsQualityControlEvidenceExtended, 27
  - artmsQualityControlSummaryExtended, 31
- \* **quality**,
  - artmsQualityControlEvidenceBasic, 25
  - artmsQualityControlMetabolomics, 30
- \* **quantifications**
  - artmsAnalysisQuantifications, 3
- \* **rename**,
  - artmsChangeColumnName, 8
- \* **reshape**
  - artmsResultsWide, 33
- \* **results**,
  - artmsResultsWide, 33
- \* **retention**
  - artmsAvgIntensityRT, 7
- \* **silac**,
  - artmsSILACtoLong, 34
- \* **species**
  - artmsAnnotateSpecie, 5
  - artmsIsSpeciesSupported, 17
- \* **spectral\_counts**,
  - artmsSpectralCounts, 35
- \* **summary**,
  - artmsAvgIntensityRT, 7
  - artmsMergeEvidenceAndKeys, 20
  - artmsQualityControlSummaryExtended, 31
- \* **summary**
  - artmsMsstatsSummary, 20
- \* **time**,
  - artmsAvgIntensityRT, 7
- \* **tools**,
  - artmsGeneratePhSiteExtended, 16
- \* **ub**,
  - artmsProtein2SiteConversion, 24
- \* **uniprot**
  - artmsAnnotationUniprot, 6
- \* **version**
  - artmsIsEvidenceNewVersion, 17
- \* **volcano**

- artmsVolcanoPlot, 35
- \* **wide**,
  - artmsResultsWide, 33
- \* **yaml**
  - artmsWriteConfigYamlFile, 36

artms\_config, 37

artms\_data\_corum\_mito\_database, 39

artms\_data\_pathogen\_LPN, 39

artms\_data\_pathogen\_TB, 40

artms\_data\_ph\_config, 40

artms\_data\_ph\_contrast, 41

artms\_data\_ph\_evidence, 42

artms\_data\_ph\_keys, 42

artms\_data\_ph\_msstats\_modelqc, 43

artms\_data\_ph\_msstats\_results, 43

artms\_data\_randomDF, 43

artmsAnalysisQuantifications, 3

artmsAnnotateSpecie, 5

artmsAnnotationUniprot, 6

artmsAvgIntensityRT, 7

artmsChangeColumnName, 8

artmsConvertMetabolomics, 8

artmsDataPlots, 9

artmsEnrichLog2fc, 10

artmsEnrichProfiler, 11

artmsEvidenceToSaintExpress, 13

artmsEvidenceToSAINTq, 14

artmsFilterEvidenceContaminants, 15

artmsGeneratePhSiteExtended, 16

artmsIsEvidenceNewVersion, 17

artmsIsSpeciesSupported, 17

artmsLeaveOnlyUniprotEntryID, 18

artmsMapUniprot2Entrez, 19

artmsMergeEvidenceAndKeys, 20

artmsMsstatsSummary, 20

artmsPhosphateOutput, 21

artmsPhotonOutput, 22

artmsPlotHeatmapQuant, 23

artmsProtein2SiteConversion, 24

artmsQualityControlEvidenceBasic, 25

artmsQualityControlEvidenceExtended,  
27

artmsQualityControlMetabolomics, 30

artmsQualityControlSummaryExtended, 31

artmsQuantification, 32

artmsResultsWide, 33

artmsSILACToLong, 34

artmsSpectralCounts, 35

artmsVolcanoPlot, 35

artmsWriteConfigYamlFile, 36