# Package 'OmnipathR'

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<b>Description</b> Import data from https://www.omnipathdb.org webservice. It also includes functions to transform and print this data.				
License MIT + file LICENSE				
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 $.get\_annotation\_databases$ 

Get the different annotation databases integrated in Omnipath

### Description

get the names of the databases from <a href="http://omnipath.org/annotation">http://omnipath.org/annotation</a>

### Usage

```
.get_annotation_databases()
```

#### Value

character vector with the names of the annotation databases

### See Also

```
import_Omnipath_annotations
```

```
.get_annotation_databases()
```

```
.get\_complexes\_databases
```

```
.get_complexes_databases
```

Get the different complexes databases integrated in Omnipath

### Description

get the names of the databases from <a href="http://omnipath.org/complexes">http://omnipath.org/complexes</a>

### Usage

```
.get_complexes_databases()
```

#### Value

character vector with the names of the databases

#### See Also

```
import_Omnipath_complexes
```

### **Examples**

```
.get_complexes_databases()
```

.get\_interaction\_databases

Get the different interaction databases

### **Description**

get the names of the databases from <a href="http://omnipath.org/interactions">http://omnipath.org/interactions</a>

### Usage

```
.get_interaction_databases()
```

### Value

character vector with the names of the interaction databases

### See Also

```
import\_AllInteractions, import\_Omnipath\_Interactions, import\_PathwayExtra\_Interactions, import\_KinaseExtra\_Interactions, import\_National (and import\_Natio
```

```
.get_interaction_databases()
```

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```
.get_intercell_categories
```

Get the different intercell categories described in Omnipath

### **Description**

```
get the names of the categories from <a href="http://omnipath.org/intercell">http://omnipath.org/intercell</a>
```

### Usage

```
.get_intercell_categories()
```

#### Value

character vector with the different intercell categories

#### See Also

```
import_Omnipath_intercell
```

### **Examples**

```
.get_intercell_categories()
```

.get\_ptms\_databases

Get Post-translational modification (PTMs) databases

### Description

get the names of the different databases available for ptms databases <a href="http://omnipath.org/ptms">http://omnipath.org/ptms</a>

### Usage

```
.get_ptms_databases()
```

#### Value

character vector with the names of the PTMs databases

#### See Also

```
import_Omnipath_PTMS
```

```
.get_ptms_databases()
```

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get_complex_genes Get all the molecular complexes for a given gene(s)
---

### Description

This function returns all the molecular complexes where an input set of genes participate. User can choose to retrieve every complex where any of the input genes participate or just retrieve these complexes where all the genes in input set participate together.

### Usage

```
get_complex_genes(complexes = import_Omnipath_complexes(),
   select_genes = query_genes, total_match = FALSE)
```

### Arguments

complexes	complexes data frame (obtained using import_Omnipath_complexes)
select_genes	$vector\ containing\ the\ genes\ for\ whom\ complexes\ will\ be\ retrieved\ (hgnc\ format).$
total_match	[default=FALSE] logical indicating if the user wants to get all the complexes where any of the input genes participate (FALSE) or to get only the complexes where all the input genes participate together (TRUE)

#### Value

data.frame of complexes

### See Also

```
import_Omnipath_complexes)
```

### **Examples**

```
complexes = import_Omnipath_complexes(filter_databases=c("CORUM", "hu.MAP"))
query_genes = c("LMNA", "BANF1")
complexes_query_genes = get_complex_genes(complexes, query_genes)
```

```
get_signed_ptms get signs for ptms interactions
```

### **Description**

ptms data does not contain sign (activation/inhibition), we generate this information based on the interaction network

#### Usage

```
get_signed_ptms(ptms = import_Omnipath_PTMS(),
  interactions = import_Omnipath_Interactions())
```

#### **Arguments**

```
ptms data frame generated by import_Omnipath_PTMS
interactions interaction data frame generated by import_Omnipath_Interactions
```

#### Value

data.frame of ptms with is\_inhibition and is\_stimulation columns

#### See Also

```
import_Omnipath_PTMS import_Omnipath_Interactions
```

#### **Examples**

```
ptms = import_Omnipath_PTMS(filter_databases=c("PhosphoSite", "Signor"))
interactions = import_Omnipath_Interactions()
ptms = get_signed_ptms(ptms,interactions)
```

import\_AllInteractions

Imports from Omnipath webservice all the available interactions from the different datasets

#### **Description**

Imports the dataset from: http://omnipathdb.org/interactions?datasets=omnipath, pathwayextra, kinaseextra, ligrecextra, tfregulons, mirnatarget&fields=sources, references&genesymbols= 1, which contains all the different interactions available in the webserver:

### Usage

```
import_AllInteractions(from_cache_file = NULL,
  filter_databases = .get_interaction_databases(),
  select_organism = 9606)
```

#### **Arguments**

interactions not reported in these databases are removed. See .get\_interaction\_databases for more information.

```
select_organism
```

Interactions are available for human, mouse and rat. Choose among: 9606 human (default), 10116 rat and 10090 Mouse

#### **Details**

omnipath: the OmniPath data as defined in the paper, an arbitrary optimum between coverage and quality pathwayextra: activity flow interactions without literature reference kinaseextra: enzyme-substrate interactions without literature reference ligrecextra: ligand-receptor interactions without literature reference tfregulons: transcription factor (TF)-target interactions from DoRothEA mirnatarget: miRNA-mRNA and TF-miRNA interactions

#### Value

A dataframe containing all the datasets in the interactions query

#### See Also

```
.get_interaction_databases
```

#### **Examples**

```
interactions <- import_AllInteractions(filter_databases=c("HPRD","BioGRID"),
    select_organism = 9606)</pre>
```

```
import_KinaseExtra_Interactions
```

Imports from Omnipath webservice the interactions from kinaseextra dataset

#### Description

Imports the dataset from: http://omnipathdb.org/interactions?datasets=kinaseextra, which contains enzyme-substrate interactions without literature reference

#### Usage

```
import_KinaseExtra_Interactions(from_cache_file = NULL,
  filter_databases = .get_interaction_databases(),
  select_organism = 9606)
```

#### **Arguments**

interactions not reported in these databases are removed. See .get\_interaction\_databases for more information.

```
select_organism
```

Interactions are available for human, mouse and rat. Choose among: 9606 human (default), 10116 rat and 10090 Mouse

#### Value

A dataframe containing enzyme-substrate interactions without literature reference

#### See Also

```
.get_interaction_databases,import_AllInteractions
```

#### **Examples**

```
import_LigrecExtra_Interactions
```

Imports from Omnipath webservice the interactions from ligrecextra dataset

#### **Description**

Imports the dataset from: http://omnipathdb.org/interactions?datasets=ligrecextra, which contains ligand-receptor interactions without literature reference

#### Usage

```
import_LigrecExtra_Interactions(from_cache_file = NULL,
  filter_databases = .get_interaction_databases(),
  select_organism = 9606)
```

#### **Arguments**

```
from_cache_file

path to an earlier data file

filter_databases
```

interactions not reported in these databases are removed. See .get\_interaction\_databases for more information.

```
select_organism
```

Interactions are available for human, mouse and rat. Choose among: 9606 human (default), 10116 rat and 10090 Mouse

#### Value

A dataframe containing ligand-receptor interaction without literature reference

#### See Also

```
.get_interaction_databases,import_AllInteractions
```

#### **Examples**

```
import\_miRNAtarget\_Interactions
```

Imports from Omnipath webservice the interactions from miRNAtarget dataset

### **Description**

Imports the dataset from: http://omnipathdb.org/interactions?datasets=mirnatarget, which contains miRNA-mRNA and TF-miRNA interactions

### Usage

```
import_miRNAtarget_Interactions(from_cache_file = NULL,
  filter_databases = .get_interaction_databases())
```

### Arguments

interactions not reported in these databases are removed. See .get\_interaction\_databases for more information.

#### Value

A dataframe containing miRNA-mRNA and TF-miRNA interactions

#### See Also

```
.get_interaction_databases,import_AllInteractions
```

```
interactions <-
  import_miRNAtarget_Interactions(filter_databases=c("miRTarBase",
  "miRecords"))</pre>
```

```
import_Omnipath_annotations
```

Import Omnipath Annotations

### Description

imports the annotations stored in Omnipath database from http://omnipathdb.org/annotations

### Usage

```
import_Omnipath_annotations(from_cache_file = NULL,
    select_genes = NULL, filter_databases = .get_annotation_databases())
```

### **Arguments**

from\_cache\_file

path to an earlier data file

select\_genes

vector containing the genes for whom annotations will be retrieved (hgnc format). It is also possible to donwload complexes annotations. To do so, write "COMPLEX:" right before the genesymbols of the genes integrating the complex. Check the vignette for examples.

filter\_databases

annotations not reported in these databases are removed. See  $.get\_annotation\_databases$  for more information.

### Value

A data.frame containing different gene/complex annotations

#### See Also

```
.get_annotation_databases
```

```
annotations = import_Omnipath_annotations(select_genes=c("TP53","LMNA"),
    filter_databases=c("HPA"))
```

```
import_Omnipath_complexes
```

Import Omnipath Complexes

### Description

imports the complexes stored in Omnipath database from http://omnipathdb.org/complexes

### Usage

```
import_Omnipath_complexes(from_cache_file = NULL,
  filter_databases = .get_complexes_databases())
```

### **Arguments**

complexes not reported in these databases are removed. See .get\_complexes\_databases for more information.

#### Value

A dataframe containing information about complexes

#### See Also

```
.get_complexes_databases
```

### **Examples**

```
complexes = import_Omnipath_complexes(filter_databases=c("CORUM", "hu.MAP"))
```

```
import_Omnipath_Interactions
```

Import Omnipath interaction database

### **Description**

imports the database from <a href="http://omnipathdb.org/interactions">http://omnipathdb.org/interactions</a>, which contains only interactions with references. These interactions are the original ones from the first Omnipath version.

### Usage

```
import_Omnipath_Interactions(from_cache_file = NULL,
  filter_databases = .get_interaction_databases(),
  select_organism = 9606)
```

#### **Arguments**

```
from_cache_file

path to an earlier data file

filter_databases

interactions not reported in these databases are removed. See . get_interaction_databases
for more information.

select_organism

Interactions are available for human, mouse and rat. Choose among: 9606 human (default), 10116 rat and 10090 Mouse
```

#### Value

A dataframe containing information about protein-protein interactions

#### See Also

```
.get_interaction_databases,import_AllInteractions
```

#### **Examples**

```
import_Omnipath_intercell
```

Import Omnipath Intercell Data

#### **Description**

imports the intercell data stored in Omnipath database from <a href="http://omnipathdb.org/intercell">http://omnipathdb.org/intercell</a>. Intercell provides information on the roles in inter-cellular signaling. E.g. if a protein is a ligand, a receptor, an extracellular matrix (ECM) component, etc.

### Usage

```
import_Omnipath_intercell(from_cache_file = NULL,
    select_categories = .get_intercell_categories())
```

#### **Arguments**

vector containing the categories to be retrieved. All the genes belonging to that category will be returned. For furter information about the categories see .get\_intercell\_categories

### Value

A dataframe cotaining information about roles in inter-cellular signaling.

#### See Also

```
.get_intercell_categories
```

#### **Examples**

```
intercell = import_Omnipath_intercell(select_categories=c("ecm"))
```

#### **Description**

```
imports the PTMs database from http://omnipathdb.org/ptms
```

### Usage

```
import_Omnipath_PTMS(from_cache_file = NULL,
  filter_databases = .get_ptms_databases(), select_organism = 9606)
```

#### **Arguments**

```
from_cache_file

path to an earlier data file

filter_databases

PTMs not reported in these databases are removed. See .get_ptms_databases
for more information

select_organism

PTMs are available for human, mouse and rat. Choose among: 9606 human
(default), 10116 rat and 10090 Mouse
```

#### Value

A data frame containing the information about ptms

#### See Also

```
.get_ptms_databases,import_Omnipath_Interactions
```

```
import\_PathwayExtra\_Interactions
```

Imports from Omnipath webservice the interactions from Pathwayextra dataset

#### **Description**

Imports the dataset from: http://omnipathdb.org/interactions?datasets=pathwayextra, which contains activity flow interactions without literature reference

#### Usage

```
import_PathwayExtra_Interactions(from_cache_file = NULL,
  filter_databases = .get_interaction_databases(),
  select_organism = 9606)
```

#### **Arguments**

```
from\_cache\_file
```

path to an earlier data file

filter\_databases

interactions not reported in these databases are removed. See . get\_interaction\_databases for more information.

select\_organism

Interactions are available for human, mouse and rat. Choose one of those: 9606 human (default), 10116 rat or 10090 Mouse

#### Value

A dataframe containing activity flow interactions between proteins without literature reference

#### See Also

```
.get_interaction_databases,import_AllInteractions
```

```
import_TFregulons_Interactions
```

Imports from Omnipath webservice the interactions from Dorothea dataset

### Description

Imports the dataset from: http://omnipathdb.org/interactions?datasets=tfregulons which contains transcription factor (TF)-target interactions from DoRothEA https://github.com/saezlab/DoRothEA

### Usage

```
import_TFregulons_Interactions(from_cache_file = NULL,
  filter_databases = .get_interaction_databases(),
  select_organism = 9606)
```

### **Arguments**

```
from_cache_file

path to an earlier data file

filter_databases

interactions not reported in these databases are removed. See .get_interaction_databases

for more information.

select_organism
```

Interactions are available for human, mouse and rat. Choose among: 9606 human (default), 10116 rat and 10090 Mouse

#### Value

A dataframe containing TF-target interactions from DoRothEA

#### See Also

```
.get_interaction_databases,import_AllInteractions
```

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interaction\_graph Build Omnipath interaction graph

### **Description**

transforms the interactions data.frame to an igraph object

#### Usage

```
interaction_graph(interactions = interactions)
```

### **Arguments**

interactions

data.frame created by import\_Omnipath\_Interactions, import\_PathwayExtra\_Interactions,
import\_KinaseExtra\_Interactions, import\_LigrecExtra\_Interactions,
import\_TFregulons\_Interactions, import\_miRNAtarget\_Interactions or
import\_AllInteractions

#### Value

An igraph object

#### See Also

import\_Omnipath\_Interactions, import\_PathwayExtra\_Interactions, import\_KinaseExtra\_Interactions, import\_LigrecExtra\_Interactions, import\_TFregulons\_Interactions, import\_miRNAtarget\_Interactions or import\_AllInteractions

#### **Examples**

```
interactions = import_Omnipath_Interactions(filter_databases=c("SignaLink3"))
OPI_g = interaction_graph(interactions)
```

OmnipathR

The OmnipathR package

### **Description**

OmnipathR is an R package built to provide easy access to the data stored in the Omnipath webservice:

```
http://omnipathdb.org/
```

The webservice implements a very simple REST style API. This package make requests by the HTTP protocol to retreive the data. Hence, fast Internet access is required for a propser use of OmnipathR.

The package also provides some utility functions to filter, analyse and visualize the data.

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#### Author(s)

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### **Examples**

```
# Download post-translational modifications:
ptms = import_Omnipath_PTMS(filter_databases=c("PhosphoSite", "Signor"))
# Download protein-protein interactions
interactions = import_Omnipath_Interactions(filter_databases=c("SignaLink3"))
# Convert to igraph objects:
ptms_g = ptms_graph(ptms = ptms )
OPI_g = interaction_graph(interactions = interactions )
# Print some interactions:
print_interactions(head(ptms))
# interactions with references:
print_interactions(tail(ptms),writeRefs=TRUE)
# find interactions between kinase and substrate:
print_interactions(dplyr::filter(ptms,enzyme_genesymbol=="MAP2K1",
  substrate_genesymbol=="MAPK3"))
# find shortest paths on the directed network between proteins
printPath_es(shortest_paths(OPI_g,from = "TYRO3",to = "STAT3",
  output = 'epath')$epath[[1]],OPI_g)
# find all shortest paths between proteins
printPath_vs(all_shortest_paths(ptms_g,from = "SRC",to = "STAT1")$res,ptms_g)
```

printPath\_es

print network paths given by edge sequence

#### **Description**

prints the interactions in the path in a nice format

### Usage

```
printPath_es(edgeSeq, G)
```

### Arguments

```
edgeSeq edge sequence
G igraph object (from ptms or any interaction dataset)
```

printPath\_vs

#### See Also

```
printPath_vs
```

### **Examples**

```
interactions = import_Omnipath_Interactions(filter_databases=c("SignaLink3"))
OPI_g = interaction_graph(interactions = interactions)
printPath_es(shortest_paths(OPI_g,from = "TYRO3",to = "STAT3", output = 'epath')$epath[[1]],OPI_g)
```

printPath\_vs

print networks paths given by node sequence

### Description

prints the interactions in the path in a nice format

#### Usage

```
printPath_vs(nodeSeq, G)
```

### Arguments

nodeSeq node sequence
G igraph object (from ptms or interactions)

### See Also

```
printPath_es
```

```
interactions = import_Omnipath_Interactions(filter_databases=c("SignaLink3"))
OPI_g = interaction_graph(interactions = interactions )
printPath_vs(shortest_paths(OPI_g,from = "TYRO3",to = "STAT3")$vpath,OPI_g)

ptms = import_Omnipath_PTMS(filter_databases=c("PhosphoSite", "Signor"))
ptms_g = ptms_graph(ptms)
printPath_vs(all_shortest_paths(ptms_g,from = "SRC",to = "STAT1")$res,ptms_g)
```

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print\_interactions

print interactions

#### **Description**

prints the interactions/ptms in a nice format

### Usage

```
print_interactions(interDF, writeRefs = FALSE)
```

### **Arguments**

interDF data.frame with the interactions generated by any of the following functions:

import\_Omnipath\_PTMS, import\_Omnipath\_Interactions, import\_PathwayExtra\_Interactions,
import\_VineseExtra\_Interactions

 $import\_Kinase Extra\_Interactions, import\_Ligrec Extra\_Interactions, import\_TFregulons\_Interactions, import\_miRNA target\_Interactions or$ 

import\_AllInteractions

writeRefs [FALSE] writes also the PubMed IDs if available

#### **Examples**

```
ptms = import_Omnipath_PTMS()
print_interactions(head(ptms))
print_interactions(tail(ptms),writeRefs=TRUE)
print_interactions(dplyr::filter(ptms,enzyme_genesymbol=="MAP2K1",
    substrate_genesymbol=="MAPK3"))
```

ptms\_graph

Post-translational modifications (PTMs) graph

### **Description**

transforms the ptms interactions data.frame to igraph object

### Usage

```
ptms_graph(ptms)
```

### **Arguments**

ptms

data.frame created by import\_Omnipath\_PTMS

### Value

An igraph object

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### See Also

```
import\_Omnipath\_PTMS
```

```
ptms = import_Omnipath_PTMS(filter_databases=c("PhosphoSite", "Signor"))
ptms_g = ptms_graph(ptms = ptms )
```

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